Evaluation of Nitrification Inhibition Using Bench-Scale Rate Measurements, Profile Sampling, and Process Simulation Modeling

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Abstract

The Hampton Roads Sanitation District (HRSD) operates thirteen treatment plants in the eastern Virginia area with a combined capacity of 231 million gallons per day (mgd). The Nansemond Treatment Plant (NTP) is one of the larger facilities, and is designed to treat 30 mgd using a 3-stage Virginia Initiative Process (VIP) biological nutrient removal (BNR) process. The majority of the influent is domestic, but there is also a large industrial contribution, particularly from a hog processing facility, landfill leachate, and significant loads from septage and grease deliveries (Bilyk et al, 2008). NTP is currently being upgraded to a 5-stage Bardenpho process to achieve improved total nitrogen (TN) removal. For several years starting in about 2001, NTP has experienced continuous and sporadic nitrification upsets that cannot be explained by plant operations events. Sporadic nitrification upsets are characterized by sharp increases in effluent ammonia and nitrite with decreases in nitrate concentrations due to reduced growth rates in bacteria. The result is reduced overall total nitrogen (TN) removal. Continuous inhibition is evidenced by a previous engineering report by Hazen and Sawyer, P.C. (2007), whereby it was suggested that the ammonia oxidizing bacteria (AOB) maximum specific growth rate (μ_{max}) be reduced from 0.9 to 0.57 days⁻¹. This has significant implications in terms of the required aeration volume for consistent nitrification at cold temperatures.

The objective of this project was to determine whether the NTP influent wastewater does in fact exhibit inhibition to ammonia (AOB) and nitrite oxidizing bacteria (NOB), evaluated independently, and to determine the impact on polyphosphate accumulating organism activity

(PAO). Because the historical operational experiences and data analysis suggested inhibited AOB and NOB activity, an investigation was initiated targeting the source of that inhibition. After conducting seventeen weeks of batch experiments the source of inhibition was not determined. Batch experiments however, did reveal other possible sources of inhibition including large amounts of chemical toilet waste received at NTP possibly containing quaternary ammonium compounds (QACs).

Due to available blower capacity during construction it was planned that nitrification would not be maintained during the fall of 2009. In an effort to stop nitrification, the solids retention time (SRT) was purposely reduced over a period of about one month (as wastewater temperature cooled) until additional blower capacity was available. This provided an opportunity to study baseline nitrification kinetics and determine the potential for continuous inhibition through profile sampling. Simulation modeling of the profile sampling and plant data was performed with *Biowin 3.1* (EnviroSim, Ltd.) as a means for comparison and to generate μ_{max} values for AOB to compare with the original design μ_{max} of 0.57⁻¹.

Profile sampling was conducted from the primary effluent to the secondary effluent with samples collected along the length of the BNR process. This was being done to address the following issues:

• Conduct baseline sampling prior to a more detailed nitrification inhibition study estimated to begin in May 2010, which will include influent sampling and the operation of bench-scale sequencing batch reactors. This will be used to establish "normal" COD, nutrient and DO profiles though the VIP process without (and possibly with) the impact of inhibitory conditions, specifically with respect to N conversions and P release and uptake along the process.

- Evaluate the potential for nitrite accumulation in the process and its potential effect on aerobic phosphate uptake by phosphorus accumulating organisms (PAOs).
- Evaluate the impact of sporadic ferric chloride addition to the biological process as a means of preventing effluent TP exceedances.
- Evaluate the design μ_{max} to the actual observed μ_{max} for AOB through simulation modeling.
- Compare modeling and observed profile data for signs of any continuous nitrification inhibition.

Experimental results from batch-rate testing confirmed the sporadically inhibitory nature of NTP primary effluent when combined with other stable nitrifying biomasses. Investigation into quaternary ammonium compounds (QACs) which were contained in the chemical toilet waste suggested that QACs at higher concentrations caused some inhibition of NOB activity, but no significant impact on AOB activity. Profile sampling demonstrated no signs of sporadic or continuous nitrification inhibition or impact of nitrite accumulation and ferric chloride addition on biological treatment processes. Modeling of the profile data generated similar profiles; however, there were slight variations as the model predicted nitrification to stop earlier than what was actually observed. From the modeling it was also determined that the maximum specific growth rate (μ_{max}) of ammonia oxidizing bacteria (AOB) was in the range of 0.50 – 60 days⁻¹. This supported batch and profile work that showed NTP PE exhibited some degree of continuous inhibition. Diurnal loadings however, were not accounted for in the modeling which could slightly underestimate the actual AOB μ_{max} value. Several suspected inhibitors were eliminated as potential causes of inhibition, including waste from a hog processing facility, landfill leachate,

the addition of ferric chloride, plant internal recycle streams, branches of the collection system, and chemical toilet disinfectants containing QACs.

References

Bilyk, K., Cubbage, L., Stone, A., Pitt, P., Dano, J., and Balzer, B. 2008. Unlocking the Mystery of Biological Phosphorus Removal Upsets and Inhibited Nitrification at a 30 mgd BNR Facility. *Proceedings of the Water Environment Federation Technical Conference and Exposition*, 2008.

Hazen and Sawyer. 2007. Nansemond Treatment Plant Nutrient Reduction Improvement Technical Memorandum.

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Introduction

1.1 Project Background

The Nansemond Treatment Plant (NTP) is one of the larger facilities of the thirteen plants operated by the Hampton Roads Sanitation District (HRSD) with a maximum monthly design capacity of 30 million gallons per day (MGD). This facility was originally designed as a 3-stage Virginia Initiative Process (VIP) biological nutrient removal (BNR) system as shown in Figure 1.1 (Bilyk et al., 2008) and is currently being upgraded to a 5-stage Bardenpho process with external carbon addition. The majority of the influent is domestic, but there is also a large industrial contribution, particularly from a hog processing facility, landfill leachate, and significant loads from septage and grease deliveries (Bilyk et al., 2008).

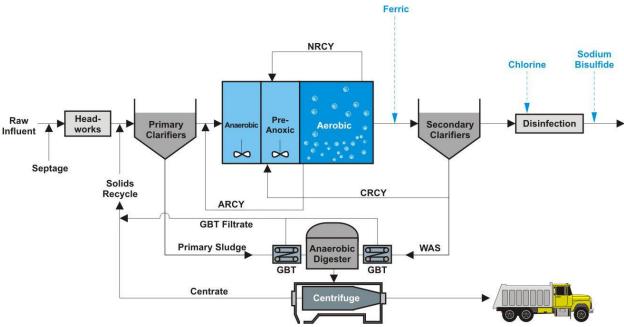


Figure 1.1 3-Stage VIP (2008) NTP Process Flow Diagram (Bilyk et al., 2008)

NTP began operations as a 10 mgd secondary treatment plant in 1983. Expansions and upgrades were completed in May 1998, which converted the facility into a 30 mgd BNR facility. Since the upgrade the facility has experienced mixed success in the BNR mode (Balzer et al.,

2005). This determination was based upon the plant's efficiency in removing nitrogen. The VIP plant in Norfolk, VA which also employs the VIP process began BNR operations approximately seven (7) years prior to similar Nansemond operations (Balzer et al., 2005). Nitrogen removal was similar at both facilities (Table 1.1) during the first two years of side-by-side operation (1999 & 2000) (Balzer et al., 2005). Nansemond experienced an unexpected decline in nitrogen removal efficiency starting in 2001, which has continued to the present (Balzer et al., 2005). This deficiency in performance has been variable and has not been consistent from 2001 to the present time.

Table 1.1 Percent TN Removal for HRSD's Nansemond and VIP Treatment Plants (1999-2009)

PLANT	NP	VIP	
YEAR	[% Removal]	[% Removal]	
1999	69.9	71.5	
2000	64.2	67.0	
2001	53.1	66.3	
2002	64.4	67.1	
2003	45.0	62.5	
2004	55.6	71.9	
2005	50.6	67.83	
2006	70.8	69.67	
2007	68	65.83	
2008	70.3	72.58	
2009	50	60	
MEAN	60.2	67.5	

NTP achieves continuous complete nitrification during certain periods consistently meeting desired effluent limits. In spite of this, NTP has also experienced unexplained sporadic nitrification upsets for a number of years and even in summer months when compared to the VIP plant (Figure 1.2 and 1.3) and some indication of continuous nitrification inhibition, as demonstrated by calibration of a process simulation model to historical plant performance data (Hazen & Sawyer, 2007). The previous target total nitrogen (TN) treatment objective was 12 mg/L on a seasonal basis. This has changed more recently to a permitted limit of 8 mg/L TN on an annual average basis (Bilyk et al., 2008).

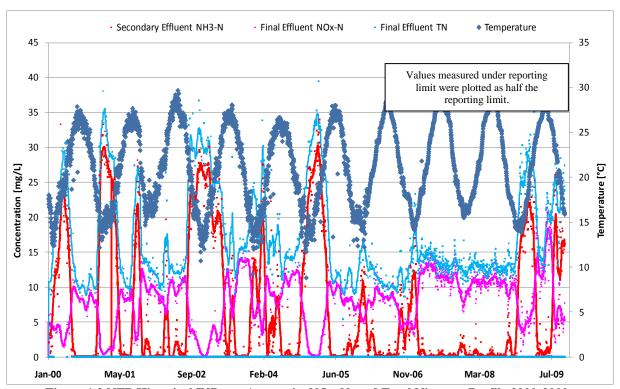


Figure 1.2 NTP Historical Effluent Ammonia, NOx-N, and Total Nitrogen Profile 2000-2009 (Lines represent 30-day rolling averages).

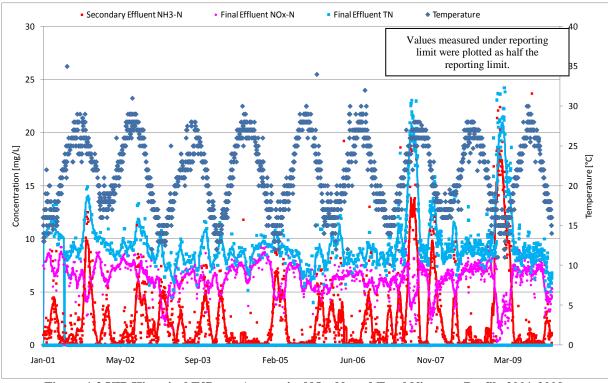


Figure 1.3 VIP Historical Effluent Ammonia, NOx-N, and Total Nitrogen Profile 2001-2009 (Lines represent 30-day rolling averages).

In addition the plant also experienced regular biological phosphorus removal upsets (Figure 1.4) which forced NTP to add large quantities of ferric chloride to avoid permit violations for effluent TP (Bilyk et al., 2008). It was the goal of NTP to meet a treatment objective of 1 mg/L of total phosphorus (TP) on an annual average basis (Bilyk et al., 2008).

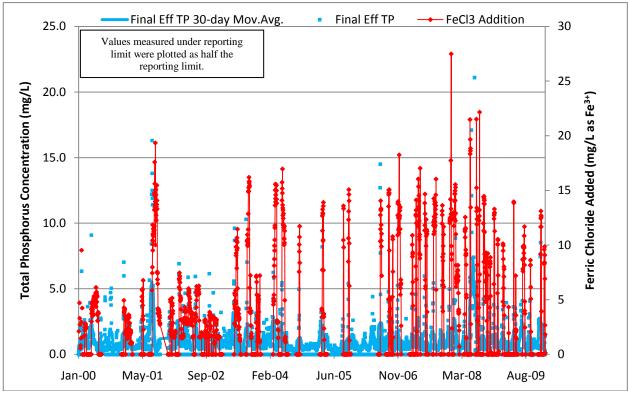


Figure 1.4 NTP Historical Effluent TP and Ferric Chloride Addition 2000-2009

1.1.1 Investigation of Inhibition

HRSD has expended considerable time and resources investigating possible sources of inhibition. It was originally presumed that contributions from industrial loads, either from a hog processing facility or landfill leachate, were the culprits for sporadic failure in nitrification and bio-P upsets. These sources were tested specifically as part of this work.

1.1.2 Facility Upgrades

In order to meet future permit limits NTP is currently being upgraded to a 5-stage Bardenpho process (Figure 1.5). This upgrade also will provide increased aeration volume/capacity to enhance both nitrification and Bio-P removal. The new upgrades will also incorporate a full-scale proprietary technology developed by Ostara that uses a fluidized bed reactor to recover phosphorus and ammonia through struvite precipitation from the centrate being generated at NTP. The harvested struvite can then be utilized as a slow release fertilizer (Ostara, 2007).

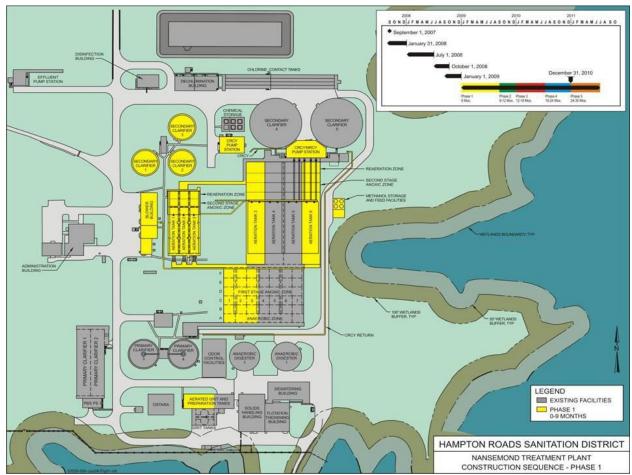


Figure 1.5 NTP with Upgrades to 5-stage Bardenpho Process.

A preliminary engineering report by Hazen and Sawyer, P.C. for the current nutrient removal upgrade, suggested that the ammonia oxidizing bacteria (AOB) maximum specific growth rate ($\mu_{max, AOB}$) be reduced from the default value of 0.90 days⁻¹ to 0.57 days⁻¹ to account

for high effluent ammonia data during the calibration period (Hazen & Sawyer, 2007). This has significant implications in terms of the required aeration volume for consistent nitrification at cold temperatures. NTP along with the other HRSD James River basin facilities are required to meet a combined annual discharge limit of 6 million pounds TN to the James River – bubble permit limit (Balzer et al., 2005). It was originally assumed that NTP would nitrify year-round with this background inhibition. This would only be made possible however, by reconfiguring the BNR process to allow the 2nd anoxic zone to operate aerobically (swing zone). Removing an anoxic zone would reduce overall TN removal because the process would essentially change from the 5-stage Bardenpho process to an A2O Process. If the inhibition can be eliminated then there would no longer be a need to configure the 2nd anoxic zone to run aerobically, providing improved annual average TN removal. This would ensure that NTP and the other James River HRSD facilities meet the TN bubble permit limit with more certainty.

1.2 Research Objectives

The objectives of this study were the following:

- Establish the inhibitory characteristics of the NTP influent wastewater through benchscale experimentation
- Study baseline nitrification kinetics and to evaluate the presence of a source of continuous inhibition
- Model the baseline nitrification kinetics and compare these values to previously calculated kinetic values.

1.2.1 Bench-scale Experimentation

The initial objectives of this research were to investigate the inhibition of nitrification at NTP through independent evaluation of ammonia oxidizing bacteria (AOB) and nitrite oxidizing

bacteria (NOB) rates, as well as, determine the impact of inhibition on polyphosphate accumulating organism activity (PAO). This was accomplished through the use of a wide variety of NTP, targeted industry, and control wastewater and biomass samples to attempt to identify possible sources of inhibition through batch rate measurements using bench-scale reactors.

1.2.2 Baseline Profile Sampling

In an effort to better understand the BNR process at the facility, profile sampling was also performed in conjunction with the bench-scale reactor experiments during the summer and fall of 2009. During the initial upgrades to the facility, nitrification was maintained; however, due to available blower capacity during construction it was planned that nitrification would not be maintained during the fall of 2009. In an effort to stop nitrification, the solids retention time (SRT) was purposely reduced over a period of about one month (as wastewater temperature cooled) until additional blower capacity was available. This coincided with the profile sampling creating an opportunity to study nitrification kinetics as the plant stopped nitrifying. Profile sampling was conducted until the plant completely ceased nitrifying in the fall of 2009. This work consisted of collecting grab samples through the BNR process and analyzing the samples for ammonia (NH₃-N), nitrite (NO₂-N), nitrate (NO₃-N), ortho-phosphate (PO₄-P), and soluble COD (sCOD), pH, and DO.

1.2.3 Biowin Simulation Modeling

The profile sampling data were combined with plant operating data and modeled using *Biowin 3.1* (EnviroSim, Ltd.). The profile sampling period (7-23-09 to 11-5-09) was divided into 5 separate periods to generate a calibrated model. The modeling for four of the periods was first performed in steady-state conditions. The period in which NTP stopped nitrifying had to be calibrated with a dynamic model. A calibrated simulation was then generated over the entire

period of profile samples. These simulations were compared to data collected during the profile sampling to better understand the level of continuous nitrification inhibition.

1.3 Thesis Organization

- Chapter two is dedicated to providing literature reviews of nitrification inhibition with known and suspected sources of inhibition, batch-rate measurements, and profile sampling.
- Chapter three provides methodologies for grab sampling, batch reactor construction,
 batch reactor operation, profile sampling, analysis of nutrients, and modeling.
- Two manuscripts were included which encompass the different aspects of this research:
 Bench-Scale Batch Testing, profile sampling, and simulation modeling of the process.

This research provides insight into measuring nitrification inhibition kinetics, evaluation of possible inhibitors, data exhibiting a fully-nitrifying plant falling out of nitrification, and modeling work which provides a comparison of measured plant performance data versus simulation modeling & profile sampling data.

1.4 References

Balzer, B., Cox, M., Deluna, J., Ghosn, S., Hogg, P., Kennedy, G., Pletl, J. 2005. The Status of Biological Nutrient Removal at HRSD's Nansemond Treatment Plant. NITRO Team Report.

Bilyk, K., Cubbage, L., Stone, A., Pitt, P., Dano, J., and Balzer, B. 2008. Unlocking the Mystery of Biological Phosphorus Removal Upsets and Inhibited Nitrification at a 30 mgd BNR Facility. *Proceedings of the Water Environment Federation Technical Conference and Exposition*, WEFTEC 08.

Hazen and Sawyer. 2007. Nansemond Treatment Plant Nutrient Reduction Improvement Technical Memorandum.

Ostara. 2007. Proposal: Full Scale Struvite Recovery Project at the Nansemond Wastewater Treatment Facility. *Ostara Nutrient Recovery Technologies*.

2. Literature Review

2.1 Nitrification Inhibition

Biological nutrient removal processes involving nitrification are widely incorporated in wastewater treatment plants. The need for nitrification in wastewater treatment arises from water quality concerns over (1) the effect of ammonia on receiving water with respect to DO concentrations (2) the toxicity of ammonia in receiving waters to aquatic and marine life, (3) the need to provide nitrogen removal to control eutrophication, and (4) the need to provide nitrogen control for water-reuse applications including groundwater infiltration (Metcalf & Eddy, 2003). Nitrification is well recognized as the most sensitive process in biological nutrient removal (BNR) systems and is susceptible to problems arising from pH, temperature, dissolved oxygen (DO) concentrations, substrate concentrations, and chemical inhibitors (Juliastuti et al., 2003a). Conventional aerobic nitrification involves the oxidation of ammonia to nitrate by two different types of bacteria, ammonia oxidizing bacteria (AOB) and nitrite oxidizing bacteria (NOB) (Metcalf & Eddy, 2003).

$$NH_4^+ + 1.5 O_2 \rightarrow 2 H^+ + H_2O + NO_2^-$$
 (1)

$$NO_2^- + 0.5 O_2 \rightarrow NO_3^- \tag{2}$$

The biochemical reactions above take place in nitrification processes in which ammonium is oxidized to nitrite by AOB and then nitrite is oxidized to nitrate through NOB. Based on the chemical reactions it can be noted that nitrification is an aerobic autotrophic process. Nitrification has been used for years to remove ammonia from wastewater prior to discharge.

Nitrifying bacteria have maximum specific growth rates that are much slower than heterotrophic bacteria in activated sludge processes. Generally, the minimum allowable solids

retention time (SRT) for operation of a nitrifying activated sludge process is controlled by nitrification kinetics per the commonly used design equations presented below:

$$\frac{1}{SRT_{\min}} = \mu_a \tag{3}$$

$$\mu_a = \left(\frac{\mu_{\text{max},a} N}{K_N + N}\right) \left(\frac{DO}{K_o + DO}\right) + b_a \tag{4}$$

where μ_a is the nitrifier specific growth rate, $\mu_{max,a}$ is the nitrifier maximum specific growth rate, N is the target effluent ammonia-N concentration, K_N is the nitrifier Monod half-saturation coefficient for ammonia, DO is the aeration basin dissolved oxygen concentration, K_o is the nitrifier Monod half-saturation coefficient for oxygen, and b_a is the autotrophic decay rate (Metcalf & Eddy, 2003).

The actual SRT used for process design is typically determined by multiplying the minimum SRT for nitrification by a safety factor ranging from 1.5 to a value as high as 5.0 (extended aeration). It is commonly found that chemical inhibitors tend to reduce the nitrifier maximum specific growth rate, $\mu_{max,a}$, but can also potentially increase the nitrifier half-saturation coefficient for ammonia, K_N . With the input of chemical inhibitor, the SRT_{min} would increase, possibly approaching the design SRT. If a system is operating at a given design SRT, this suggests that the presence of a chemical inhibitor may not affect process performance (effluent ammonia-N concentration) but could make the system much more susceptible to nitrification problems as a result of wastewater temperature fluctuations or peak ammonia loading events. As a result, it is critical to evaluate the effect of chemical inhibitors on nitrification kinetic parameters or on direct surrogates of those parameters (e.g. nitrate generation rate) (Kelly et al, 2004; Daigger and Sadick, 1998; Hockenbury and Grady, 1977).

Nitrification is not only used for ammonia removal, but is also commonly used in conjunction with denitrification processes for total nitrogen removal. The denitrification process is less sensitive to changes in pH, temperature, and other factors than the nitrification process; therefore it is practical to investigate the more sensitive nitrification process (Pagga et al, 2006). The specific inhibition of nitrification could have dramatic effects on a plants ability to treat wastewater properly and discharge within permitted limits. For many of the compounds, the concentrations that inhibit the nitrifying bacteria may be an order of magnitude lower than the concentrations which inhibit the heterotrophic bacteria that remove biodegradable organic matter (Daigger and Sadick, 1998). For this reason, it is important to know the inhibition potential of substances on nitrification to prevent disturbances (Pagga et al, 2006). It is generally advised to test chemicals and wastewaters which might be considered toxic or inhibitory by suitable laboratory tests in advance (Pagga et al, 2006). Many facilities have reported experiencing biotreatment process upset conditions based on a WERF Report survey conducted in 2000 (Love and Bott, 2000). Ineffective nitrification was reported as the second most inhibited process to COD/BOD removal out of survey conducted on 110 different treatment facilities (Love and Bott, 2000). This suggests that nitrification inhibition is a widespread and common problem found at wastewater treatment facilities.

There are several types of experiments and methods used to examine nitrification inhibition. Experiments can be carried out using continuously stirred tank reactors, batch reactors, or nitrifying bioreactors in bench, pilot, and full scale studies (Hu et al, 2004).

Nitrification inhibition can be measured through measuring oxygen uptake rates, also known as respirometry, and nitrate generation or ammonia uptake rate (NGR or AUR). In most cases, the experiments listed above utilize nitrifying biomass or activated sludge from a wastewater

treatment plant that is being studied and add different possible or known inhibitors to the reactors to examine oxygen, ammonia, nitrate, and nitrite uptake and generation rates. Although there is extensive literature on nitrification inhibition each situation must be independently evaluated to assess the cause of inhibition and solution to the problem. There are many potential sources of nitrification inhibition in wastewater which are discussed further in subsequent sections

2.1.1 Chemical Inhibition

Many transient upset events are known to be caused by shock loads of toxic chemicals. Furthermore, studies have shown that chemical toxins can detrimentally affect all the essential processes within an activated sludge treatment system (Henriques et al, 2007). Organic compounds at certain concentrations can promote inhibition of nitrification. According to literature it has been observed that chlorobenzene and trichloroethylene are capable of causing inhibition at much lower concentrations than phenol or ethylbenzene (Juliastuti et al, 2003a). Previous research by others has found nitrification to be highly susceptible to upset in wastewater treatment (Blum and Speece, 1991; Wood et al, 1981). Full-scale and laboratoryscale studies have shown that industrial contaminants are frequently the source of such upset events, and these sources can adversely affect the nitrification process for weeks (Hu et al, 2002; Nowak and Svardal, 1993). Literature has also shown that inhibitory chemicals can often be formed in the solids processing trains of wastewater treatment facilities (Daigger and Sadick, 1998). Numerous studies using pure cultures of nitrifying bacteria have reported that industrial toxins can be inhibitory (Anthonisen et al, 1976; Blum and Speece, 1991; Grunditz and Dalhammar, 2001). Furthermore, recovery of nitrification after an inhibitory event can take time (Stasinakis et al, 2003), leaving the treatment system vulnerable to permit violations and the downstream environment vulnerable to ecological damage (Kelly et al, 2004).

2.1.2 Impact of Heavy Metals on Biological Treatment

Metals have been found in significant concentrations in various wastewater streams. Contributions from industrial sources have been assumed as a cause of nitrification inhibition in more recent times (Hu et al, 2002). This is supported by the increasing trend of discharging industrial effluent to publicly owned treatment works (POTW) for treatment, which increases the possibility of contamination in the influent by metal ions (Stasinakis et al, 2003). Although a constant low-level exposure to metals does not typically affect microbial activity due to biomass acclimation, shock loads of metals can lead to complete failure of biological processes (Hu et al, 2004). The presence of heavy metals can also adversely affect the operation of biological treatment processes by accumulating to inhibitory concentrations. Many studies have investigated the effects of heavy metals in biological systems alone or in combination with others.

A high variation is seen in the reported inhibitory range for metals, since different experimental conditions exist in all studies (i.e. exposure time, type of buffer, pH, type and concentration of ligands) (Semerci et al, 2007). In addition to this, interpretation of results is based on different metal species such as total, labile, free or biosorbed metal. Under these circumstances, it is very difficult to compare the inhibitory concentration ranges (Semerci et al, 2007). Previous short-term studies have demonstrated that nitrification inhibition generally correlates well with the aqueous free metal cation concentration (Hu et al, 2004). There are many different types of metals which can be found in wastewaters of both municipal and industrial origin; however, nickel, cadmium, copper, and zinc are some of the more commonly found metals because of their widespread industrial use (Hu et al, 2004). Inhibition of nitrification is clear for heavy metals such as cadmium, zinc, and copper when the biomass is exposed to higher

shock load (24-hour period) concentrations of the metal (0.2 - 0.65 mg/L Cd²⁺, 0.5 - 3 mg/L Zn²⁺, 10 - 12.5 mg/L Cu²⁺) as supported by literature (Hu et al, 2002, Kelly and Love, 2004, Semerci et al, 2007, Madoni et al, 1999). Metals also have a dual effect on microbial growth and act either as trace elements or as inhibitors (Juliastuti et al, 2003).

2.1.3 Quaternary Ammonium Compounds (QAC)

Quaternary ammonium compounds (QACs) or quaternary ammonium salts (quaternary ammines) are salts of quaternary ammonium cations with a coordinating anion (e.g. chloride). They are organic compounds that contain four functional groups attached covalently to a positively charged central nitrogen atom (R_4N_+). These functional groups (R_4) include at least one long chain alkyl group, and the rest are either methyl or benzyl groups. QACs are extensively used in domestic and industrial applications as surfactants, emulsifiers, fabric softeners, disinfectants and corrosion inhibitors (Tezel et al, 2007).

QACs are of importance here because it has been determined that acute inhibition of both heterotrophic COD removal and nitrification, especially the nitrite oxidation process, can be inhibited at high concentrations (Kreuzinger et al, 2007). Boethling (Boethling, 1984) suggests that at higher concentrations, for which no acclimation has occurred, the presence of QACs causes significant inhibition. There has been some investigation into the impact of QACs on biological wastewater treatment, which have found that acclimation may occur, one reason for which appears to be complexation of QACs with anionic surfactants and/or adsorption to particulate matter (Yang et al, 2008). The focus of this review is the effect of common QACs used as disinfectants and deodorants for chemical toilet liquids and their impact on aerobic biological wastewater treatment.

The Hampton Roads Sanitation District (HRSD) Nansemond Treatment Plant (NTP) receives large quantities of chemical toilet waste that are discharged to the plant in slug doses via tank truck to the septage receiving station (see Tables 2.1 and 2.2). It was hypothesized that the QACs contained in this chemical toilet waste could be a possible source for the sporadic nitrification inhibition that is experienced at the NTP.

Table 2.1 Septage Waste Received for Several HRSD WWTPs

Wastewater	Residential Septage	Chemical Toilet Waste	Total Delivered Waste	Average Daily Flow	Chemical Toilet Waste	Total Septage
Treatment Plant	[gallons/year]	[gallons/year]	[gallons/year]	[MGD]	[gallons/day]	[gallons/day]
Atlantic	501,937	560,258	1,885,655	40	1,535	5,166
Boat Harbor	90,762	852,656	1,618,515	14.9	2,336	4,434
Chesapeake-Elizabeth	454,906	369,426	1,284,278	13.2	1,012	3,519
Nansemond	1,800,999	1,291,602	4,427,292	18.5	3,539	12,130
Williamsburg	1,549,416	272,326	4,000,171	15.7	746	10,959
York River	2,592,031	76,418	2,730,400	12.9	209	7,481

Table 2.2 Septage Waste Received as a Fraction of Average Daily Plant Flow

WWTP Loading	Chemical Toilet Waste	Chemical Toilet Waste
Highest to Lowest	[gallons/day]	[% Avg Day Flow]
Nansemond	3,539	0.019
Boat Harbor	2,336	0.016
Chesapeake-Elizabeth	1,012	0.008
Williamsburg	746	0.005
Atlantic	1,535	0.004
York River	209	0.002

2.1.3.1 Impact of QACs on Biological Treatment

Relevant QACs associated with disinfectants or chemical toilet additives are BAC, DDAC, dichlorobenzyl dimethyl dodecyl ammonium chloride, cetyltrlmethylammonium bromide, and laurylpyridlnlum methosulfate (Gerike et al, 1990). Previous research has been focused on BAC and DDAC, although one study was conducted which examined all the above QACs and the biodegradability and inhibitory threshold concentrations of these compounds (Geirke et al, 1990). It was found from this study that inhibitory effects were noted either in an oxygen consumption inhibition test (respirometry) or by comparing the degradation performance in a biological test culture with disinfectant added to a control (Gerike et al, 1990).

Study from the work of Boethling suggests that even with equivalent amounts of anionic surfactants and acclimation, slug loading which would result in temporarily high concentration in treatment plants could upset plant function and completely inhibit nitrification (Boethling, 1984). This implies that increases in QAC concentration would have a disruptive effect on sewage treatment even when anionic surfactants are also present (Boethling, 1984). This is especially true for the nitrification process, which appears to be somewhat more sensitive to inhibition by QACs than heterotrophic COD removal (Boethling, 1984).

In an another study, it was determined that the second step of nitrification for which nitrite oxidizing bacteria (NOB) convert nitrite (NO₂⁻) to nitrate (NO₃⁻) exhibited the greatest inhibition in response to QAC loading (Kreuzinger et al, 2007). The study used both short and long term tests to look at both acute and chronic inhibition of nitrification due to exposure to QAC compounds. For acute tests, oxygen consumption for carbon removal and nitrification were measured after addition of various concentrations (0.02, 0.2, 0.5, 1, 5, 10, 25, 50, 100 mg/L) of the single QACs, BAC-C12–16 and DDACC10–18, and these data were used to estimate acute maximum autotrophic growth rates (Kreuzinger et al, 2007). Chronic inhibition was evaluated in bench-scale biological treatment systems with continuous dosage of the test substances in synthetic wastewater (peptone, meat extract, urea, NaCl, CaCl₂, MgSO₄, K₂HPO₄) for a duration of 6 months, applying final concentrations of 0.1, 1 and 2 mg/L QAC mixture described above for at least 1 month. Chronic effects were assessed by comparing COD removal and nitrification efficiency with a control that did not receive QAC (Kreuzinger et al, 2007). It was found that the following QACs and concentrations were inhibitory to nitrification;

Benzyldimethyldodecylammonium chloride (BDMDAC):8-22 mg/L,

dichlorobenzyldimethyldodecylammonium chloride (DCBDMDAC):16 mg/L, and

cetyltrlmethylammonlum bromide (CTAB):15 mg/L (Kreuzinger et al, 2007). These values were measured using the oxygen consumption inhibition test method.

Results from recent research on alkyl benzel dimethyl ammonium chloride (AB), using mixed aerobic and nitrifying cultures, demonstrated that the mixed aerobic cultures are able to efficiently degrade up to 50 mg/L AB when fed with dextrin and peptone (Yang et al, 2008). Nitrification was complete at a concentration 20 mg/L AB only after an acclimation period, but was almost fully inhibited at 50 mg/L (Yang et al, 2008). This supports previous work which suggests that aerobic cultures can efficiently degrade QACs if properly acclimated at lower concentrations, but at higher concentrations or slug loads can experience complete inhibition. A mixed aerobic culture was also fed only AB as the external nitrogen and carbon source and was able to achieve high AB degradation at both 20 and 50 mg/L AB (Yang et al, 2008).

Nitrifying cultures were also examined with the addition of AB from 2-20 mg/L (Yang et al, 2008). Results from this assessment showed that at lower concentrations (2 and 5 mg/L AB) there was no significant inhibition compared to the control, although the rate of the first step of nitrification (NH₃-N to NO₂-N) was slower than the control (Yang et al, 2008). Inhibition starting at 10-20 mg/L AB was apparent in the nitrifying cultures as ammonia was not fully utilized at 10 mg/L and complete inhibition occurred above 15 mg/L (Yang et al, 2008). An interesting finding from this study was that no nitrite accumulation was observed which suggests that in this particular work AOB were more sensitive to AB than NOB (Yang et al., 2008).

2.1.4 Impact of Nitrification Inhibition on Biological Nutrient Removal (BNR)

The inhibition of nitrification, whatever the cause may be, has a tremendous impact on BNR process performance. Ineffective nitrification can lead to nitrite accumulation which could

also have an effect on biological phosphorus removal (Bio-P) (Meinhold et al, 1999). Nitrite accumulation at higher concentrations has been found to interfere with PAO metabolism causing PHA utilization and anoxic phosphate uptake to cease (Meinhold et al, 1999). Nitrate is preferred for use during PAO metabolism as it can be utilized in both nitrogen and phosphorus removal creating a "double use" which will result in reduced sludge production. Also nitrate is preferred over oxygen as it can reduce aeration demand (Meinhold et al, 1999).

Nitrification inhibition can also create issues for denitrification. Nitrate is provided from the return activated sludge (RAS), which is recycled from the secondary clarifiers. If NOB activity is inhibited than less nitrate would be produced, thus impacting denitrifiers which utilize nitrate as a primary electron acceptor.

A typical uninhibited AOB maximum specific growth rate is 0.80-1.0 d⁻¹; however, conditions that lead to continuous nitrifier inhibition could have growth rates significantly less than this range. For process design, AOB maximum specific growth rate generally controls the minimum SRT; therefore inhibited rates can create issues with the activated sludge process in a BNR system. Increase in the minimum SRT due to slower nitrification kinetics would require larger aeration tank volumes and associated aeration in addition to potentially larger secondary clarifiers.

2.2 Bench-Rate Measurements

2.2.1 AOB & NOB Rate Measurement

AOB and NOB activity is important when reviewing a plant's nitrification performance. The kinetic parameters associated with these rates, such as the maximum specific growth rate, are important when examining plant performance and are also a good indication of whether the plant is nitrifying properly. Traditional single-step modeling of nitrification is adequate under

sole rate limitation by NH₄⁺–N to NO₂⁻–N oxidation (Chandran et al, 2000b). However, such modeling yields meaningless kinetic parameter estimates when NO₂⁻–N to NO₃⁻–N oxidation or both oxidation steps limit overall nitrification during periods of the test assay (Chandran et al, 2000b). Respirometry - based two step nitrification models can permit biokinetic estimation of both NH₄⁺–N to NO₂⁻–N oxidation and NO₂⁻–N to NO₃⁻–N oxidation from a single NH₄⁺–N to NO₃⁻–N oxidation respirogram (Chandran et al, 2000). However, for any nitrification design or control efforts based on batch respirometry derived biokinetic estimates, it is beneficial to identify the rate-limiting step in overall nitrification by estimating and comparing the kinetics of each step (Chandran et al, 2005).

Both full-scale and laboratory scale studies using the activated sludge process have suggested that a wide variety of industrial chemicals can inhibit the nitrification process for extended periods (Stasinakis et al, 2003; Kelly et al, 2004; Nowak et al, 1993), and that recovery from this inhibition can take a significant amount of time (Kelly et al, 2004; Stasinakis et al, 2003). Respirometry, which measures oxygen uptake rate and nitrate generation rate (NGR), which determines the nitrate production rate, are most commonly used to measure nitrification inhibition (Kelly and Love, 2004). Another common method for determining nitrification rates and inhibition is through the use of titrimetric techniques which examine nitrification rates based on the rate of base addition for pH stabilization (Kelly and Love, 2004).

The two main methods used to evaluate nitrification kinetics related to this work and, specifically the effect of chemical toxins on nitrification:

1. <u>Respirometry</u>: involves the measurement of oxygen uptake rate (OUR) for microbes associated with biological treatment. A sample of mixed liquor is removed from a full, pilot, or bench-scale system, placed in a sealed reactor, possibly amended with substrate

or nitrification inhibitor, and the rate of oxygen consumption is monitored over time. To evaluate nitrification kinetics, a sample of mixed liquor is added to a temperaturecontrolled respirometer reactor with and without (control) a chemical stressor. The mixed liquor can be supplemented with primary effluent (PE), ammonia-spiked PE, secondary effluent (SE), ammonia, or nitrite. When conducting experiments to measure nitrification kinetics it is desired to start the experiment with relatively high levels of ammonia to allow for a longer experimental run and to ensure that the maximum nitrification rate is maintained. If this is the case, careful control of pH must be maintained to ensure the pH does not drop below about 6.8-7.0 (alkalinity is typically added in the form of sodium carbonate). If organic substrate is added (e.g. PE), these experiments can be run with and without nitrification inhibitor to distinguish between heterotrophic and autotrophic oxygen uptake. Since endogenous heterotrophic oxygen uptake can occur without organic substrate addition, experiments are often run with and without nitrification inhibitor even when ammonia is the only substrate added to the mixed liquor. It is possible to calculate nitrification kinetic parameters based on specific oxygen uptake rate (SOUR) profiles (note that the term "specific" indicates that the OUR has been normalized to the biomass concentration).

2. <u>Nitrate/Nitrite Generation Rate</u>: In order to evaluate the nitrification process fully independent of heterotrophic activity, kinetic rates directly related to the consumption or production of reactants and products of the nitrification process itself can be measured, specifically nitrate/nitrite generation rate (NGR) or ammonia uptake rate (AUR). The specific nitrification rate (SNR) can be obtained by normalizing to biomass concentration, and this data can be used to determine the autotrophic kinetic parameters

described above. For these experiments, a sample of mixed liquor is added to small temperature controlled reactors (approximately 3.0 L). The reactor is mixed and aerated, and the inhibitor of interest added to the stressed reactor. Once all residual organic substrate associated with the mixed liquor sample itself is consumed, ammonia or nitrite is spiked into the reactor, and caution must be taken to ensure that pH and alkalinity remain within acceptable limits – preferable pH 7 to 8 at all times. The nitrate, nitrite, and ammonia concentrations are monitored over time using typical analytical methods (APHA, 1998). It is critical to rapidly separate the mixed liquor from the soluble supernatant as quickly as possible after removing a sample from the reaction reactor. Typically, samples are removed from the reaction reactors at predetermined time intervals and rapidly centrifuged. The supernatant is poured off and immediately filtered through a 0.45 µm membrane filter. The filtrate can then be preserved for subsequent analysis.

Respirometry is a good method of measuring nitrification inhibition, but also has proven useful for indicating activated sludge process stability both in lab and full-scale experimentation (Kelly and Love, 2004). Respirometry has an advantage over NGR in that it is rapid and doesn't require extensive sample analyses. Unfortunately respirometry measures the total oxygen uptake rate of a biomass. This requires several iterations of tests to determine the respiration rate of only the nitrifying bacteria in a mixed community like mixed liquor because the nitrifying bacteria must be specifically inhibited from other species. To do this, a total respirometry must be performed as well as a respirometry where the biomass has been inhibited for nitrification using compounds such as heavy metals, organic compounds, QACs or industrial wastes (Kelly and Love, 2004).

NGR provides a direct measure of the rate of nitrification, as it measures the generation of nitrate

and nitrite, the product of two-stage nitrification (Kelly and Love, 2004). This can be used to measure AOB rates based on NOx-N generation or NOB based on NO₃-N generation. Although it provides a direct measure, it also requires more time to complete than respirometry.

Kelly et al (2004b) compared the use of respirometry to NGR measurement for two different chemical compounds and suggested that NGR is the preferred method for determining nitrification inhibition because the NGR test yields a direct measure of the nitrification rate through measurement of the final product while respirometry only provided indirect measurement of nitrifier activity (Kelly and Love, 2004). It was also noted that during one of the experiments in comparing respirometry and NGR measurement due to cadmium inhibition, respirometry seemed to over predict inhibition at lower concentrations and under predict inhibition at higher concentrations (Kelly and Love, 2004). NGR test results also exhibited better reproducibility within duplicate reactors (Kelly and Love, 2004).

2.2.2 Biological Phosphorus Rate Measurement

The use of enhanced biological phosphorus removal (EBPR) has been shown to be an economical and environmentally acceptable method for reducing phosphorus from wastewaters (Erdal, et al, 2006). The most common and widely used test method for measuring EBPR is the uptake and release test (URT). URTs provide a qualitative assessment of phosphorus accumulating organisms (PAO) activity in activated sludge that can be used to develop kinetic parameters for process simulation model calibration, such as the ratio of phosphate released to acetate consumed, and providing information on phosphate uptake kinetics that can be used to estimate the time required to remove quantities of phosphate in the aerobic zone (Neethling et al., 2005). By adding excess volatile fatty acids (VFA) such as sodium acetate to mixed liquor, PAO activity and the amount of stored phosphorus limit the measured phosphate release

(Neethling et al, 2005). The ratio of uptake HRT to release HRT can be used as a performance indicator as it correlates well to the observed performance in full-scale facilities according to Neethling et al (2005). Ratio between uptake and release rate can also be used as a performance indicator if data is consistent and accurate in determining aerobic contact time in full-scale treatment (Neethling et al, 2005).

There are several parameters known to influence phosphorus removal efficiency. Availability of readily biodegradable substrate in the influent of the system plays an important role in performance (Mota et al, 2001). VFAs used by PAOs are generated by fermentation under anaerobic conditions and some may be present in more septic wastewaters (Mota et al, 2001). The presence of nitrate in the anoxic stages of EBPR processes is widely recognized to have a repressive effect on phosphorus release and on net phosphorus removal (Mota et al, 2001). This is because the presence of nitrate creates a competition between denitrifying bacteria and PAOs for the VFAs which are the typical carbon source for the formation of polyhydroxyalkanoates (PHAs) (Mota et al, 2001). Studies have shown that better system performance was related to reduced competition for substrate in the non-oxic zones, which results in larger populations of PAOs, and thus, greater EBPR efficiency (Erdal et al, 2006). Solids retention time and temperature have also shown to play an important role in EBPR efficiency which has been supported by several different studies (Whang et al, 2001; Erdal et al, 2006). Other factors which can contribute to reliable EBPR performance include the role of fermenters or other processes used to enhance EBPR, the management of return flows from anaerobic solids processing steps, and chemical addition for phosphorus polishing and EBPR backup (Neethling et al, 2005).

2.3 Profile Sampling

Profile sampling of wastewater treatment plants provides an effective method for evaluating the performance of a BNR processes by sampling for species such as ammonia, nitrate, nitrite, orthophosphate, soluble COD, etc. through the treatment train. Profiles are most commonly incorporated with DO and nutrient profiles (NH₃-N, NO₃-N, etc.), but can be applied to different applications; examining different bacterial species or the solids content of a reactor. Although there is not much literature on this specific sampling method, it is very much often used in conjunction with modeling work and evaluating existing and future nutrient removal processes (Kochany et al, 2007; Scott et al, 2008; Kim et al, 2009). Profile sampling has also been used in determining the fate and transport of chemicals in wastewater treatment. There has been work conducted on the fate and transport of mercury, where samples were collected from various process locations and analyzed for mercury and methylmercury (Downing et al, 2008). When conducting profile sampling it is useful to determine what parameters need to be profiled to generate useful quantitative data. Another use for profile sampling is to determine various rates such as ammonia uptake rates, nitrate production rates, and phosphorus uptake and release rates. Grab sampling or composite sampling are both viable methods for profile sampling; however, composite sampling would require an automatic filtration system or manual composite sampling, since samples would require immediate filtration. The method for sampling is primarily based on the application, but generally involves the collection of various samples at different periods of time or various locations along the length of a treatment process.

One study used DO profiling during a feed cycle as an indication of upset conditions and monitoring SBR biomass recovery. It was found that based on DO profiling, upset conditions could be early detected and recovery measures quickly applied (Kochany et al, 2007). The same study using SBRs found that there was a correlation between the ammonia removal during a feed

cycle and the oxygen concentration in the reactors providing the same air supply (Kochany et al, 2007). This suggests that monitoring the oxygen profile (DO) can be used instead of ammonia analyses. Since DO measurements are faster as compared to ammonia analyses, determining DO profile during a feed cycle can provide useful information about efficiency of the biological system, without using expensive ammonia on-line analyzers (Kochany et al, 2007).

2.4 References

Al-Ahmad, A., Wiedmann-Al-Ahmad, M., Schön, G., Daschner, F.D., Kümmerer, K. 2000. Role of *Acinetobacter* for Biodegradability of Quaternary Ammonium Compounds. *Bulletin of Environmental Contamination & Toxicology*, 64 (6), 764-770.

American Public Health Association, American Water Works Association, Water Environment Federation. (1998). *Standard Methods for the Examination of Water and Wastewater*; 20th ed.: Washington D.C.

Anthonisen A.C., Loehr R.C., Prakasam, T.B.S. Srinath, E.G. 1976. Inhibition of Nitrification by Ammonia and Nitrous-Acid. *Journal Water Pollution Control Federation*, 48 (5), 835-852.

Bilyk, K., Cubbage, L., Stone, A., Pitt, P., Dano, J., Balzer, B., 2007. Unlocking the Mystery of Biological Phosphorus Removal Upsets and Inhibited Nitrification at a 30 mgd BNR Facility. *Water Environmental Federation*, 5133-5151.

Blum, D.J.W., Speece, R.E. 1991. A Database of Chemical Toxicity to Environmental Bacteria and its use in Interspecies Comparisons and Correlations. *Research Journal of the Water Pollution Control Federation*, 63 (3), 198-207.

Boethling, R.S. 1984 Environmental fate and toxicity in wastewater treatment of quaternary ammonium surfactants, *Water Research*, 18 (9), 1061–1076.

Bohm, B. 1994. A Test Method to Determine Inhibition of Nitrification by Industrial Wastewaters. *Water Science and Technology*, 30 (6), 169-172.

Bott, C.B., Love, N.G. 2002. Investigating a Mechanistic Cause for Activated-Sludge Deflocculation in Response to Shock Loads of Toxic Electrophilic Chemicals. *Water Environment Research*, 74 (3), 306-315.

Bull, J.P., Serreqi, A.N., Chen, T., Breuil, C. 1998. Development of an Immunoassay for a Quaternary Ammonium Compound, Benzyldimethyldodecylammonium Chloride. *Water Research*, 32 (12), 3621-3630.

Chandran, K., Smets, B.F., 2000. Applicability of two-step models in estimating nitrification kinetics from batch respirograms under different relative dynamics of ammonia and nitrite oxidation. *Biotechnol. Bioeng.* 70, 54–64.

Chandran, K., Smets, B.F., 2000b. Single-step nitrification models erroneously describe batch ammonia oxidation profiles when nitrite oxidation becomes rate limiting. *Biotechnol. Bioeng.* 68, 396–406.

Chandran, K., Smets, B.F. 2005. Optimizing experimental design to estimate ammonia and nitrite oxidation biokinetic parameters from batch respirograms. *Water Research*, 39 (20), 4969-4978.

Daigger, G., Sadick, T.E. 1998. Evaluation of methods to detect and control nitrification inhibition with specific application to incinerator flue-gas scrubber water. *Water Environment Research*, 70 (7), 1248-1256.

Downing, J., Dunlavey, E., Enoki, N., Schafer, P., Tucker, D. 2008. Fate and Transport of Mercury in a Large Advanced Wastewater Treatment Plant. *WEFETC 08*, 339-358.

Erdal, U.G., Erdal, Z.K., Randall, C.W. 2006. The Mechanism of Enhanced Biological Phosphorus Removal Washout and Temperature Relationships. *Water Environment Research*, 78 (7), 710-715.

Garcia, M.T., Campos, E., Sanchez-Leal, J., Bibosa, I. 1999. Effect of the Alkyl Chain Length on the Anaerobic Biodegradability and Toxicity of Quaternary Ammonium Based Surfactants. *Chemosphere*, 38 (15), 3473-3483.

Gendig, C., Domogala, G., Agnoli, F., Pagga, U., Strotmann, U.J. 2003. Evaluation and Further Development of the Activated Sludge Respiration Inhibition Test. *Chemosphere*, 52, 143-149.

Gerike, P., Fischer, W. K., Jasiak, W. 1978. Surfactant Quaternary Ammonium Salts in Aerobic Sewage Digestion. *Water Research*, 12, 1117-1122.

Gerike, P., Gode, P. 1990. The Biodegradability and Inhibitory Threshold Concentration of Some Disinfectants. *Chemosphere*, 21 (6), 799-812.

Grey, G., Schillinger, G., Anderson, K. 2008. Investigation of the Impact of a High Salt Wastewater on Biological Nitrification. *Water Environment Federation: WEFTEC 08*. 239-252.

Grillitsch, B., Gans, O., Kreuzinger, N., Scharf, S., Uhl. M., Fuerhacker, M. 2006. Environmental risk assessment for quaternary ammonium compounds: a case study from Austria. *Water Science & Technology*, 54 (11-12), 111-118.

Grunditz, C., Dalhammar, G. 2001. Development of Nitrification Inhibition Assays using Pure Cultures of *Nitrosomonas* and *Nitrobacter*. *Water Research*, 35 (2), 433-440.

Gu, A.Z., Majed, N., Benisch, M., Neethling, J.B. 2009. Fractionation and Treatability Assessment of Phosphorus in Wastewater Effluents – Implications on Meeting Stringent Limits. *WEFTEC 09*, 480-500.

Gue´rin-Me´chin, L., Dubois-Brissonnet, F., Heyd, B., Leveau, J.Y. 1999. Specific variations of fatty acid composition of *Pseudomonas aeruginosa* ATCC 15442 induced by Quaternary Ammonium Compounds and relation with resistance to bactericidal activity. *Journal of Applied Microbiology*, 87, 735-742.

Hazen and Sawyer. 2007. Nansemond Treatment Plant Nutrient Reduction Improvement Technical Memorandum.

Henriques, I.D.S., Holbrook, R.D., Kelly, R.T., Love, N.G. 2005. The impact of floc size on respiration inhibition by soluble toxicants—a comparative investigation. *Water Research*, 39, 2559-2568.

Henriques, I.D.S., Kelly, R.T., Dauphinais, J.L., Love, N.G. 2007. Activated Sludge Inhibition by Chemical Stressors—A Comprehensive Study. *Water Environment Research*, 79 (9), 940-951.

Higgins, M.J., Novak, J.T. 1997. The Effect of Cations on the Settling and Dewatering of Activated Sludges: Laboratory Results. *Water Environment Research*, 69 (2), 215-224.

Hockenbury, M.R., Grady, C.P.L. 1977. Inhibition of Nitrification – Effects of Selected Organic Compounds. *Research Journal WPCF*, 49 (5), 768-777.

Hu, Z., Chandran, K., Grasso, D., Smets, B.F. 2002. Effect of Nickel and Cadmium Speciation on Nitrification Inhibition. *Environmental Science and Technology*, 36, 3074-3078.

Hu, Z., Chandran, K., Grasso, D., Smets, B.F. 2003. Impact of Metal Sorption and Internalization on Nitrification Inhibition. *Environmental Science and Technology*, 37, 728-734.

Hu, Z., Chandran, K., Grasso, D., Smets, B.F., 2004. Comparison of nitrification inhibition by metals in batch and continuous flow reactors. *Water Research*, 38 (18), 3949-3959. Huang, J.Y.C. Metal Inhibition of Nitrification. *Industrial Waste Conference*, 37, 85-93.

Jonsson, K., Grunditz, C., Dalhammer, G., Jansen J.L.C. 2000. Occurrence of Nitrification Inhibition in Swedish Municipal Wastewaters. *Water Research*, 34 (9), 2455-2462.

Juliastuti, S.R., Baeyens, J., Creemers, C., 2003a. Inhibition of Nitrification by Heavy Metals and Organic Compounds: The ISO 9509 Test. *Environmental Engineering Science*, 20 (2), 79-90.

- Juliastuti, S.R., Baeyens, J., Creemers, C., 2003b. The Inhibitory Effects of Heavy Metals and Organic Compounds on the net maximum specific growth rate of the autotrophic biomass in activated sludge. *Journal of Hazardous Materials*, B100, 271–283.
- Kelly, R.T., II, Henriques, I.D.S., Love, N.G., 2004. Chemical Inhibition of Nitrification in Activated Sludge. *Biotechnology and Bioengineering*, 85 (6), 683-693.
- Kelly, R.T., II, Love, N.G., 2004. A Critical Comparison of Methods Used to Determine Nitrification Inhibition. *Water Environmental Federation*, (15) 166-180.
- Khin, T., Gheewala, S.H., Annachhatre, A.P. 2002. Modeling of Nitrification Inhibition with Aniline in Suspended-Growth Processes. *Water Environment Research*, 74 (6) 531-539.
- Kim, H.S., Pei, R., Gunsch, C., Gellner, J.W., Boltaz, J.P., Freudenberg, B., Dodson, R., Cho, K.D., Schuler, A.J. 2009. Trace Organic Chemical Profiles in Nutrient Removal Systems With and Without Integrated Fixed Film Activated Sludge. *WEFTEC 09*, 704-711.
- Kochany, J., Lipczynska-Kochany, E., Smith, W. 2007. Modeling Strategy for SBR Recovery After Upset. *WEFTEC 07*. Session 44, 3242-3251.
- Kong, Z., Vanrolleghem, P., Willems, P., Verstraete, W. 1996. Simultaneous Determination of Inhibition Kinetics of Carbon Oxidation and Nitrification with a Respirometer. *Water Research*, 30 (4), 825-836.
- Kreuzinger, N., Farnleitner, A., Wandl, G., Hornek, R., Mach, R. 2003. Molecular biological methods (DGGE) as a tool to investigate nitrification inhibition in wastewater treatment. *Water Science and Technology*, 47 (11), 165-172.
- Kreuzinger, N., Fuerhacker, M., Scharf, S., Uhl, M., Gans, O., Grillitsch, B., 2007. Methodological approach towards the environmental significance of uncharacterized substances—quaternary ammonium compounds as an example. *Desalination*. 215, 209-222.
- Laopaiboon, L., Hall, S.J., Smith, R.N. 2002. The Effect of Quaternary Ammonium Biocide on the Performance and Characteristics of Laboratory-scale Rotating Biological Contactors. *Journal of Applied Microbiology*, 93, 1051-1058.
- Li, X.Z., Zhao, Q.L. 1999. Inhibition of Microbial Activity of Activated Sludge by Ammonia in Leachate. *Environmental International*, 25 (8), 961-968.
- Love, N.G., Bott, C.B. 2000. A Review and Needs Survey of Upset Early Warning Devices. *Water Environment Research Federation*.
- Madoni, P., Davoli, D., Guglielmi, L. 1999. Response of SOUR and AUR to Heavy Metal Contamination in Activated Sludge. *Water Research*, 33 (10), 2459-2464.

Mamais, D., Noutsopoulos, C., Stasinakis, A.S., Kouris, N., Andreadakis, A.D. 2008. Comparison of Bioluminescence and Nitrification Inhibition Methods for Assessing Toxicity to Municipal Activated Sludge. *Water Environment Research*, 80 (6), 484-488.

McMahon, K.D., Jenkins, D., Keasling, J.D. 2001. Polyphosphate Kinase Genes From Activated Sludge Carrying Out Enhanced Biological Phosphorus Removal. *WEFTEC 01*. Session 1.

Mota, C.R., Batista, J.R., Unz, R.R., Buch, H., Johnson, W. 2001 Characterization Profiles and Correlations of Major Parameters in a Full-Scale Biological Phosphorus Removal System. *WEFTEC 01*.

Neethling, J.B., Bakke, B., Benisch, M., Gu, A., Stephens, H., Stensel H.D., Moore, R. 2005. Factors Influencing the Reliability of Enhanced Biological Phosphorus Removal. *WERF Report*. 01-CTS-3.

Nishiyama, N., Toshima, Y., Ikeda, Y. 1995. Biodegradation of Alkyltrimethylammonium Salts in Activated Sludge. *Chemosphere*, 30 (3), 593-603.

Nowak, O., Svardal, K. 1993. Observation on the Kinetics of Nitrification Under Inhibiting Conditions Caused by Industrial Wastewater Compounds. *Water Science and Technology*, 28 (2), 115-123.

Neufeld, R., Greenfield, J., Rieder, B., 1986. Temperature, Cyanide, and Phenolic Nitrification Inhibition. *Water Research*, 20 (5), 633-642.

Pagga, U. 1997. Testing Biodegradability with Standardized Methods. *Chemosphere*, 35 (12), 2953-2972.

Pagga, U., Bachner, J., Strotmann, U. 2006. Inhibition of nitrification in laboratory tests and model wastewater treatment plants. *Chemosphere*, 65, 1-8.

Perles, C.E., Matheus, A., Volpe, P.L.O. 2008. The effect of quaternary ammonium surfactants on the inhibition of the aerobic metabolism of *Saccharomyces cerevisiae*—A calorimetric study. *Thermochimica Acta*, 479, 28-31.

Schweighofer, P., Nowak, O., Svardal, K., Kroiss, H. 1996. Steps Towards The Upgrading of a Municipal WWTP Affected by Nitrification Inhibiting Compounds – A Case Study. *Water Science and Technology*, 33 (12), 39-46.

Scott, Z., Olson, B.H., Esmond, S., Maleki, N., Scherfig, J. 2008. Monitoring and Modeling the Anaerobic Digestion of Manure with qPCR in a Mesophilic Anaerobic Digester. *WEFTEC 08*, 534-553.

Semerci, N., Cecen, F. 2007. Importance of cadmium speciation in nitrification inhibition. *Journal of Hazardous Materials*, 147, 503–512.

- Stasinakis, A. S., Thomaidis, N. S., Mamais, D., Papanikolaou, E. C., Tsakon, A., Lekkas, T. D. 2003. Effects of chromium (VI) addition on the activated sludge process. *Water Research*, 37 (9), 2140-2148.
- Stricker, A., Lishman, L., Barrie, A. 2008. Effects of Fluctuating Iron dosage on Nitrification in Integrated Fixed Film and Conventional Activated Sludge Processes. *Water Environment Federation: WEFTEC 08*, 5022-5043.
- Sütterlin, H., Alexy, R., Coker, A., Kümmerer, K. 2008. Mixtures of quaternary ammonium compounds and anionic organic compounds in the aquatic environment: Elimination and biodegradability in the closed bottle test monitored by LC–MS/MS. *Chemosphere*, 72, 479-484.
- Sütterlin, H., Alexy, R., Coker, A., Kümmerer, K. 2008. The toxicity of the quaternary ammonium compound benzalkonium chloride alone and in mixtures with other anionic compounds to bacteria in test systems with Vibrio fischeri and Pseudomonas putida. *Ecotoxicology and Environmental Safety*, 71, 498-505.
- Tezel, U., Pierson, J. A., Pavlostathis, S.G. 2006. Fate and effect of quaternary ammonium compounds on a mixed methanogenic culture. *Water Research*, 40, 3660-3668.
- Tezel, U., Pierson, J. A., Pavlostathis, S.G. 2007. Effect of polyelectrolytes and quaternary ammonium compounds on the anaerobic biological treatment of poultry processing wastewater. *Water Research*, 41, 1334-1342.
- Tezel, U., Pierson, J. A., Pavlostathis, S.G. 2008. Effect of didecyl dimethyl ammonium chloride on nitrate reduction in a mixed methanogenic culture. *Water Science & Technology*, 57.4, 541-546
- Thomas, W.A., Bott, C.B., Regmi, P., Schafran, G., McQuarrie, J., Rutherford, B., Baulmer, R., Waltrip, D. 2009. Evaluation of Nitrification Kinetics for a 2.0 MGD IFAS Process Demonstration. *Water Environment Federation: Nutrient Removal 2009*.
- Tubbing, D.M.J., Admiraal, W. 1991. Inhibition of Bacterial and Phytoplanktonic Metabolic Activity in the Lower River Rhine by Ditallowdimethylammonium Chloride. *Applied and Environmental Microbiology*, 57 (12), 3616-3622.
- Weber, S.A., Sherrard, J.H. 1980. Effects of Cadmium on the Completely Mixed Activated-Sludge Process. *Water Pollution Control Federation*, 52 (9), 2378-2388.
- Wee, V.T., Kennedy, J.M. 1982. Determination of Trace Levels of Quaternary Ammonium Compounds in River Water by Liquid Chromatography with Conductometric Detection. *Analytical Chemistry*, 54 (9), 1631-1633.
- Whang, L.M., Park, J.K. 2001. Competition Between Polyphosphate- And Glycogen-Accumulating Organisms in Enhanced Biological Phosphorus Removal Systems Effect of Temperature and Sludge Age, *WEFTEC 01*

Wikipedia. Quaternary Ammonium Cation. Accessed: 16 March 2009. Last Modified: 17 February 2009 http://en.wikipedia.org/wiki/Quaternary_ammonium_compounds

Wikipedia Chemical Toilet. Accessed: 17 March 2009. Last Modified: 25 February 2009. http://en.wikipedia.org/wiki/Chemical_toilet

Yang, J., Li, K., Tezel, U., Pierson, J.A., Pavlostathis, S.G. 2008. Effect of Alkyl Benzyl Dimethyl Ammonium Chloride and Temperature on Nitrification. *WEFTEC 08*, 151-168.

Zarnovsky, L., Derco, J., Kuffa, R., Drtil, M. 1994. The Influence of Cadmium on Activated Sludge Activity. *Water Science and Technology*, 30 (11), 235-242.

Zilles, J.L., Peccia, J., Hung, C.H., Noguera, D.R. 2001. Molecular Microbial Ecology of Enhanced Biological Phosphorus Removal in Aerated-Anoxic Orbal Processes. *WEFTEC 01*.

3. Methodology

3.1 Bench-Scale Batch Reactor Experimentation

3.1.1 Sample Collection

Batch reactor experimentation was conducted by collecting biomass samples from the aeration basin effluent of several full-scale wastewater treatment plants. Grab samples were collected unless otherwise arranged with HRSD to collect composites. All samples were collected and stored in polypropylene carboys or similar containers. These samples were then transported to the lab and stored until use with continuous aeration for no longer than approximately 48 hours at room temperature. Biomass samples were collected from NTP; the HRSD VIP plant, which uses the same BNR process as NTP but does not experience inhibition; the Henrico County WWTP, which has a fully nitrifying 5-stage BNR process; and the HRSD York River plant from the sequencing batch reactor system.

Wastewater sources for experimentation were also collected via grab sample or composite sampling by HRSD based on the source being tested. Wastewater samples were collected including NTP and VIP primary effluent (PE); NTP and VIP secondary effluent (SE); raw wastewater from isolated branches of the NTP collection system; and industrial waste samples from a hog processing plant and a landfill leachate stream. Waste samples were also tested from other suspected sources such as chemical toilet disinfectants (quaternary ammonium compounds).

3.1.2 Batch Reactor Construction

Four parallel batch reactors were constructed of plexiglass and configured with vinyl and polypropylene fittings and appurtenances. The first reactor served as the control while operating conditions of the other three reactors were varied. Each reactor had a volume of 3 liters with

individual ports for sampling and chemical addition (acetate, ammonia, nitrite, and phosphate) during experimentation as needed. A diffuser stone was incorporated into each reactor to provide air/oxygen or nitrogen based on the experiment. This allowed the reactors to be operated under aerobic, anoxic, or anaerobic conditions for nitrification, denitrification, and phosphorus release/uptake measurements. The reactors were configured to be air tight with a water seal on the headspace for anaerobic conditions. Mixing was provided through magnetic stir-bars and stir plates. Dissolved oxygen (DO) was kept constant using HACH LDO probes with HQ40d meters. Two probes were connected to a single HQ40d meter (total of four probes and two meters). National Instruments LabView 7.0 DO Controller was used to provide on and off control of four solenoid air supply valves with associated relays. pH was controlled and logged using Cole-Parmer pH meters/controllers connected to acid (1M H₂SO₄) and base (3M Na₂CO₃) pumps. The reactors were placed in a water bath with a circulating chiller and immersion heaters to maintain a constant temperature during experimentation. Refer to Figure 3.1, 3.2, and 3.3 for a schematic and image of an individual reactor and for a complete schematic and image of the entire reactor configuration.

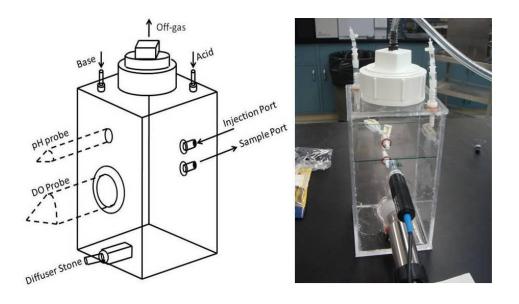


Figure 3.1 Reactor Schematic and Image

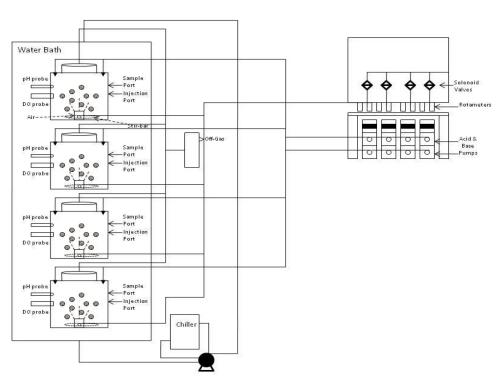


Figure 3.2 Complete Reactor System Schematic

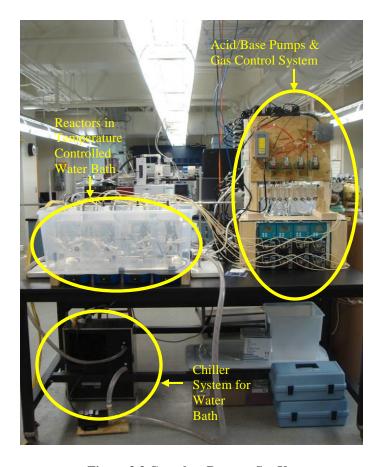


Figure 3.3 Complete Reactor Set-Up

3.1.3 Batch Reactor Operation

Experimentation consisted of the evaluation AOB, NOB, and Bio-P activity. The method for evaluating each type of experiment is discussed further in subsequent sections. Temperature for experimentation was based on the temperature of the biomass sample when it was collected onsite via grab sample. The matrix of experiments developed and completed is summarized in Table 3.1. The abbreviations in the table are as follows: Nansemond (NS), VIP (VIP), Henrico (HR), York River (YR), primary effluent (PE), secondary effluent (SE), raw water influent (RWI), primary clarifier influent (PCI), primary clarifier effluent (PCE), branches of collection system (Branch 1, 2, 3), iron addition (Fe).

Table 3.1 Batch Experiment Matrix

Week #	Biomass Source	Table 3.1 Batci	Experiments		
1	Nansemond	a. NS/PE	b. VIP/PE	AOB, NOB, Bio-	
	VIP	c. VIP/PE	d. NS/PE	P	
2	Nansemond	a. NS/PE	b. VIP/PE	AOB, NOB, Bio-P	
	VIP	c. VIP/PE	d. NS/PE		
3	Nansemond	a. VIP/SE + Leachate	b. VIP/SE	AOB, NOB, Bio-P	
	VIP	c. VIP/SE + Leachate	d. VIP/SE		
4	Nansemond	a. NS/PE	b. HR/PE	AOB, NOB	
	Henrico	c. HR/PE	d. NS/PE		
	Nansemond	a. VIP/SE + Hog Plant	b. VIP/SE	AOB, NOB, Bio-	
5	VIP	c. VIP/SE + Hog Plant	d. VIP/SE	P	
	Nansemond	a. VIP/SE + Leachate	b. VIP/SE		
6	VIP	c. VIP/SE + Leachate	d. VIP/SE	AOB, NOB	
	Nansemond	a. VIP/SE	b. VIP/SE + Branch 1	LOP YOU	
7	VIP	c. VIP/SE + Branch 2	d. VIP/SE + Branch 3	AOB, NOB	
0	Nansemond	a. VIP/SE	b. VIP/SE + Branch 2a	AOB, NOB	
8	VIP	c. VIP/SE + Branch 2b	d. VIP/SE + Branch 3		
0	Nansemond	a. VIP/SE	b. VIP/SE + Cedar Ln PS	AOB, NOB	
9	VIP	c. VIP/SE + Gum Rd. PRS	d. VIP/SE + Pughsville PRS		
10	Nansemond	a. VIP/PE	b. NTP RWI	AOB, NOB	
10	VIP	c. NTP PCI	d. NTP PCE		
11	Nansemond	a. VIP/SE	b. VIP/SE + 20 mg/L Fe	AOB, NOB	
11	VIP	c. VIP/SE + 35 mg/L Fe	d. $VIP/SE + 50 \text{ mg/L Fe}$		
12	Nansemond	a. VIP/SE	b. VIP/SE + 20 mg/L Fe	AOD NOD	
12	VIP	c. VIP/SE + 35 mg/L Fe	d. $VIP/SE + 50 \text{ mg/L Fe}$	AOB, NOB	
13	York River (YR)	a. YR/SE	b. YR/SE + 20 mg/L Fe	AOB, NOB	
13	York River (YR)	c. YR/SE + 35 mg/L Fe	d. YR/SE + 50 mg/L Fe		
14	York River (YR)	a. YR/PE	b. NTP RWI	- AOB	
	York River (YR)	c. NTP PCI	d. NTP PCE		
15	York River (YR)	a. YR/SE	b. YR/SE + 0.574 mL Chem Toilet Additive*	AOB, NOB	
	York River (YR)	c. YR/SE + 13.96 mL Chem. Toilet Additive*	d. YR/SE + 62.84 mL Chem Toilet Additive*		
16	York River (YR)	a. YR Mixed Liquor (ML)	b. YR ML + 15 mL Product B	- AOB, NOB	
10	York River (YR)	c. YR ML + 30 mL Product B	d. YR ML + 60 mL Product B		
17	York River (YR)	a. YR Mixed Liquor (ML)	b. YR ML + 60 mL Product B	AOB, NOB	
	York River (YR)	c. YR ML + 120 mL Product B	d. YR ML + 180 mL Product B		

*NOTE: Week 15 Experimentation – Two different Chemical Toilet Additives used for the two different days of experimentation (NatureFresh – Day 1 & Blue Works – Day 2)

3.1.3.1 AOB Experimentation

AOB activity was evaluated by spiking the reactors with ammonia and monitoring NO_x-N and NO₃-N generation rate over three hours (Kelly et al, 2004). During an AOB experiment the collected biomass was concentrated to approximately 8000 mg/L MLSS prior to experimentation. Then the reactor was filled 1/3 by volume with biomass and 2/3 by volume with a diluent (primary effluent, waste sample, etc.). Additional ammonia was added to achieve a total ammonia concentration of approximately 40 mg/L NH₄-N to provide sufficient ammonia during experimentation to prevent depletion. Dissolved oxygen (DO) and pH were constantly monitored and maintained within desired limits using online control systems. DO was maintained between 5-7 mg/L during experiments with a pH between 7.0-7.2. Each AOB experiment was conducted for 3 hours with 5 samples collected evenly throughout this period of time. Samples were immediately filtered through 0.45μm membrane filters after collection. AOB activity was based on NO_x-N generation rate.

3.1.3.2 NOB Experimentation

NOB activity was assessed by adding in nitrite (ensuring ammonia concentrations were low) and monitoring the rate of NO₂-N depletion and NO₃-N generation. NOB experimentation was conducted very similar except after the reactors were filled by volume as mentioned above they were aerated for approximately one hour to allow uptake of any residual ammonia. Samples were taken prior to starting the NOB Experimentation and analyzed for NH₄-N to ensure that the ammonia concentration in each reactor was <1 mg/L NH₄-N (detection limit of HACH TNT 831 NH₄-N analysis method). After it was determined that the ammonia concentration in each reactor was depleted, approximately 25 mg/L NO₂-N was added, and then 5 samples were collected evenly over a 3 hour test period. Samples were immediately filtered through 0.45μm membrane

filters after collection. DO and pH were maintained within the same parameters as was used for the AOB experiment. NOB activity was based on NO₃-N generation rate and NO₂-N uptake rate.

3.1.3.3 Bio-P Experimentation

Bio-P activity was determined by monitoring P uptake and release rates. Experimentation was conducted using the same concentrated biomass as in the AOB & NOB experimentation (~8000 mg/L MLSS). Experimentation was conducted in three phases: uptake, release, and a second uptake. During the first uptake (aerobic) phase, each reactor was filled with 1/3 by volume of biomass and 2/3 by volume of PE or another tested wastewater source. Then each reactor was spiked with 5 mg/L of PO₄-P to reach a target of 20 mg/L P in each reactor since it was presumed approximately 15 mg/L of PO₄-P would be present from the biomass and PE/wastewater sample. The first uptake phase was conducted for 1-2 hours and 3 samples were collected during this time with immediate filtration using 0.45µm membrane filters. During both uptake phases, pH was always controlled between 7.0-7.2 while the DO was maintained between 5-7 mg/L. After this initial uptake phase, the reactors were deaerated by sparging nitrogen (N₂) gas and reducing the DO to 0 mg/L which was maintained throughout the release phase. After the DO was reduced to 0 mg/L, each reactor was spiked with 200 mg/L acetate as COD. It was desired to have all NO₃-N and NO₂-N depleted, but this was not always the case. pH was controlled the same as in the uptake phase experiment and 5 samples were collected over a 1 hour period with immediate filtration using 0.45µm membrane filters. After the 1 hour release phase nitrogen was shut-off and then air/oxygen was sparged into the reactors to once again raise the DO and maintain the DO between 5-7 mg/L. As soon as the air/oxygen sparge began, 5 samples were collected over a 1-2 hour period with immediate filtration using 0.45µm membrane filters. All the samples from both uptake phases and the release phase were analyzed using the

same methods used for the AOB and NOB experiments noted in the subsequent 3.3 Analytical Methods section. Rates based on PO₄-P release and uptake was normalized to MLVSS concentrations for each experiment.

All experimentation was conducted in this manner except for that conducted during weeks 16 & 17, during which the entire reactor volume was filled with biomass due to the biomass from the York River plant at that time being very dilute.

3.2 Profile Sampling

Profile sampling was conducted from the primary effluent to the secondary effluent with samples collected along the length of the BNR process. A total of 15 samples were collected. Analysis of soluble COD (sCOD), Ammonia (NH₄-N), Nitrate (NO₃-N), Nitrite (NO₂-N), orthophosphate (PO₄-P) at all sampling locations.

- Primary Clarifier Effluent (PE) 1 Sample
- Anaerobic/Anoxic Tanks (AA1, AA2, AA3, AA4, AA5, AA6) 6 Samples, 1 sample
 per cell for a single train
- Aeration Tank (AE1, AE1.25, AE1.5, AE3, AE5) 5 samples, 1 at the beginning, 3 intermediate, 1 at the end for a single tank. The aeration tank was originally divided into 5 equal sample points along the length of the tank. AE1, AE2, AE3, AE4, AE5. Based on the data results two of the intermediate points were changed. Instead of taking 5 equally spaced points two were taken between the distance of AE1 and AE2, which became AE1.25 and AE1.5 to better capture ammonia uptake, nitrite consumption, and nitrate production. AE4 was eliminated as values were similar to AE5 (Figure 3.4).
- Clarifier Recycle (RAS) 1 sample

- Secondary Clarifier Influent Distribution Channel (IDC) 1 sample
- Secondary Clarifier Effluent (SE) 1 sample

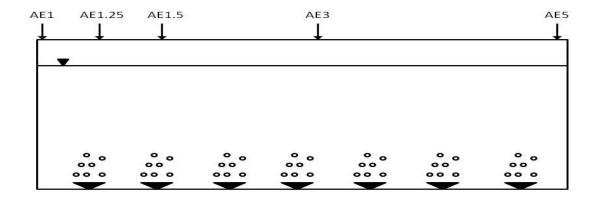


Figure 3.4 Aeration Tank Profile Sample Points

Each sample was collected via grab sample from the surface of the tank using a bucket or sample dipper. The sample was allowed to settle for approximately 30 seconds to allow a supernatant layer to form. The supernatant was then aspirated and immediately filtered through 0.45µm membrane filters using a portable vacuum filtration system. DO and pH were measured and recorded when each grab sample was collected. Each sample was also collected in accordance with corresponding HRTs associated with each process. Samples were collected in two different containers; (1) 15 mL centrifuge tube for on-site analysis using HACH colorimetric test kits and (2) 40 mL VOA vial for soluble COD analysis by the HRSD Central Environmental Laboratory (CEL). All samples collected were kept chilled in a cooler on ice until they could be refrigerated to <6°C. Samples which were collected and placed into the 40 mL VOA vials were preserved with H₂SO₄ acid as soon as samples were collected and stored at a temperature of <6°C in a refrigerator until they were transported for analysis by CEL the following morning.

This process for profile sampling was performed twice a week during normal plant operation hours (0600 to 1500) for a period of approximately 2 and half months until the plant stopped nitrifying.

3.3 Analytical Methods

3.3.1 AOB, NOB, and Bio-P Analysis

Samples were analyzed for NH₄-N, NO₂-N, NO₃-N, and PO₄-P using HACH colorimetric test kits, ion chromatography (IC), and flow injection analysis (FIA) (see subsections 3.3.3, 3.3.6, 3.3.7, 3.3.8). MLSS and MLVSS were performed at the end of the experimentation per *Standard Methods* (APHA et al, 1995). Nitrate production rates, phosphorus uptake and release rates were calculated by normalizing the slope of each chemical constituent by the MLVSS concentration and adjusting units to mg/g MLVSS/hr.

3.3.2 Profile Sampling Analysis

Sample analysis for profile sampling was conducted both on-site and off-site. Analytes measured at NTP for each profile include: PO₄-P, NH₄-N, NO₃-N, NO₂-N which were conducted using various HACH colorimetric test kits (see subsections 3.3.3 – 3.3.6) and a HACH DR2800 spectrophotometer. Analysis of NO₂-N was performed the same day as sample collection, while analysis of PO₄-P, NH₄-N, and NO₃-N were conducted the following day. Samples were preserved by refrigeration overnight after NO₂-N analysis was complete. Off-site analysis by CEL consisted of soluble COD and quality assurance and quality control samples (QA/QC) of the same analytes measured on-site using *Standard Methods*.

3.3.3 Ammonia (NH₄-N) HACH Test Kit

Ammonia was analyzed using HACH Test N' Tube (TNT) 831 kit and HACH DR2800 spectrophotometer. This method uses the salicylate method, whereby ammonium ions react with

hypochlorite and salicylate ions in the presence of sodium nitroprusside to as a catalyst form indophenol. The amount of color formed is directly proportional to the NH₄-N present.

3.3.4 Nitrate (NO₃-N) HACH Test Kit

Nitrate was analyzed using HACH TNT 835 kit. This kit incorporates the dimethylphenol method where nitrate ions in solution with sulfuric and phosphoric acids react with 2,6-dimethyphenol to form 4-nitro-2,6-dimethyphenol.

3.3.5 Nitrite (NO₂-N) HACH Test Kit

Nitrite was analyzed using NitriVer3 Nitrite Reagent Powder Pillows and 10 mL sample vials. This kit uses the diazotization method where nitrite in the sample reacts with sulfanilic acid to form a intermediate diazonium salt. This salt combined with chromotropic acid forms a pink color which is directly proportional to the amount of nitrite present.

3.3.6 Ortho-Phosphate (PO₄-P) HACH Test Kit

Ortho-Phosphate was analyzed using the HACH Reactive Phosphate TNT Reagent Kit.

This test kit uses the USEPA-approved PhosVer3 method where orthophosphate reacts with molybdate in an acid to produce a mixed complex. Ascorbic acid then reduces this complex, producing an intense blue color.

3.3.7 Ion Chromatography

IC analysis was conducted for measuring nitrate only for the batch experimentation work. The IC was used in conjunction with conductivity detection (Dionex ICS-1000), an AS14A analytical column and AG14A guard column, an ASRS conductivity suppressor, and an eluent flow of 1.0 mL/min of 1.0 mM NaHCO3 and 8.0 mM Na2CO3.

3.3.8 Flow Injection Analysis

FIA was conducted for measuring nitrite, nitrate, and phosphate for batch experimentation work using a SEAL Analytics auto-analyzer system. FIA analysis incorporated a volume of 2 mL of sample in sample vials and placed into a sample tray. The tray was then placed into the FIA instrument along with various reagents required for the different analyte test methods. A schedule was then configured using the associated computer software with the instrument and analyzed. QA/QC samples were measured every 10 samples and test methods were always standardized prior to analysis of actual samples. Data output was provided in concentrations in mg/L as NO₂-N, NO₃-N, and PO₄-P.

3.4 Biowin Modeling

Modeling was performed using Biowin version 3.1, a biological wastewater treatment simulation package developed by EnviroSim Ltd (Flamborough, Ontario, Canada) and based on the IWA activated sludge models. Biowin was incorporated into this work to compare the data generated from the profile sampling with previous work performed by Hazen and Sawyer, P.C. (H&S). The Biowin model was calibrated using recent plant performance data and previous simulation modeling work from H&S. There are various wastewater fractions (Table 3.2) which must be adjusted by the user based on the wastewater source (Primary Effluent). Wastewater fraction inputs for this work were derived from the NTP model created by H&S. The H&S NTP model was simulated with the raw influent characteristics that were generated from historical plant data and a two week special sampling study specified by H&S. The effluent data from the primary clarifier of this simulation was used to calculate wastewater fractions for the input of the simulation for the profile sampling work. Primary clarification was not simulated as part of this effort. Kinetic parameters such as maximum specific growth rate ($\mu_{max, AOB}$) and nitrite half saturation concentration were changed based on the work done by H&S in addition to changes

made to fit the simulation model to the profile sampling data. A Garret wasting configuration was incorporated to simplify modeling of secondary clarification (Figure 3.5), and the waste rate was then adjusted accordingly to match the solids wasted (lbs/day) from the plant performance data within a reasonable range. Diurnal load variations were not considered as part of this modeling.

Table 3.2 Biowin Influent Wastewater Fractions

Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.310
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.180
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.610
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.155
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.169
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.740
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.300
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.030
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.820
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.009
FZbh - Non-poly-P heterotrophs [gCOD/g of total COD]	0.000	0.0001
FZbm - Anoxic methanol utilizers [gCOD/g of total COD]	0.000	0.0001
FZaob - Ammonia oxidizers [gCOD/g of total COD]	0.000	0.0001
FZnob - Nitrite oxidizers [gCOD/g of total COD]	0.000	0.0001
FZamob - Anaerobic ammonia oxidizers [gCOD/g of total COD]	0.000	0.0001
FZbp - PAOs [gCOD/g of total COD]	0.000	0.0001
FZbpa - Propionic acetogens [gCOD/g of total COD]	0.000	0.0001
FZbam - Acetoclastic methanogens [gCOD/g of total COD]	0.000	0.0001
FZbhm - H2-utilizing methanogens [gCOD/g of total COD]	0.000	0.0001

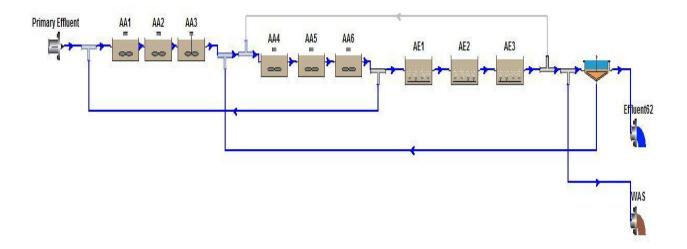


Figure 3.5 Biowin Simulation Model (Garrett Wasting Configuration)

The Biowin simulation was also reconfigured for dynamic simulation. During the profile sample period one anaerobic/anoxic (AA) train was taken offline and so the model had to be adjusted to be able to model this during the entire profile period. For this reason a separate set of AA tanks were added and the flow split between the two trains in the model. One train received 4/5 of the flow and train B received 1/5 of the flow. When all tanks were online the flow was split to each of the trains to simulate the actual 5 trains that were online in the full-scale. When the one train was shut down in the full-scale the flow split to train B (1/5 of the flow) was stopped to mimic what actually occurred (Figure 3.6).

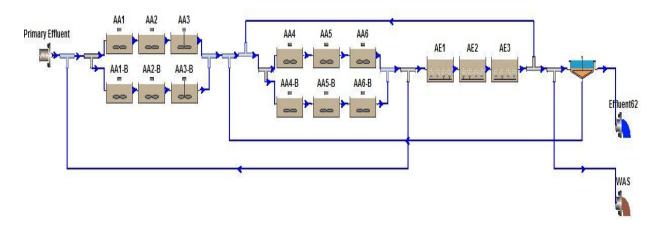


Figure 3.6 Biowin Model for Dynamic Simulation

3.4.1 Steady-State Simulations

Simulations were created for individual periods of the profile sampling as well as a simulation for the entire profile sampling period. Individual periods were created based on changes in mixed liquor suspended solids, changes in process; 5 anaerobic/anoxic trains to 4 anaerobic/anoxic trains; the Nitrate recycle (NRCY) being active and inactive, and the cease of nitrification. Steady-state simulations incorporated the use of constant input values for recycle streams, WAS, and other inputs. The entire profile sampling event was divided into 5 different periods and each period was modeled using steady-state conditions to calibrate the simulation except for the 4th period.

3.4.2 Dynamic Simulations

During the 4th period the plant was falling out of nitrification and therefore this period had to be modeled dynamically to calibrate it properly. The model for the entire profile sampling event from start to finish was modeled dynamically as well. Itineraries for influent (Primary Effluent) characteristics, temperature, recycle rates, waste rates, and clarifier removal rates were all set based on plant performance data and the steady-state & dynamic calibration models.

3.5 References

American Public Health Association, American Water Works Association, Water Environment Federation. (1998). *Standard Methods for the Examination of Water and Wastewater*; 20th ed.: Washington D.C.

Kelly, R.T., II, Love, N.G., 2004. A Critical Comparison of Methods Used to Determine Nitrification Inhibition. *Water Environmental Federation*, (15) 166-180.

4. Manuscript 1 – Evaluation of Nitrification Inhibition Using Bench-Scale Rate Measurements

Abstract

The Nansemond Treatment Plant (NTP) operated by the Hampton Roads Sanitation District (HRSD) was originally designed as a 3-stage VIP biological nutrient removal (BNR) process (Bilyk et al., 2008). NTP is currently being upgraded to a 5-stage Bardenpho process to achieve improved total nitrogen (TN) removal. NTP has experienced unexplained sporadic nitrification upsets for a number of years and some indication of continuous nitrification inhibition, as demonstrated by calibration of a process simulation model to historical data. A preliminary engineering report by Hazen and Sawyer, P.C., suggested that the ammonia oxidizing bacteria (AOB) maximum specific growth rate (μ_{max}) be reduced from 0.90 to 0.57 days⁻¹ to account for high effluent ammonia data during the calibration period (Hazen & Sawyer, 2007). This has significant implications in terms of the required aeration volume for consistent nitrification at cold temperatures. A study was undertaken using a wide variety of NTP, targeted industry (hog processing facility and landfill leachate), and control wastewater and biomass samples to attempt to identify possible sources of inhibition through batch rate measurements using bench-scale reactors. Biomass samples were collected from various well nitrifying facilities to compare nitrification kinetics. Experiments were carried out using four batch reactors in parallel. This batch rate testing independently evaluated AOB and NOB activity based on NO_x-N generation and NO₃-N generation/NO₂-N depletion, respectively. Experimental results to date have confirmed the sporadically inhibitory nature of NTP primary effluent when combined with other nitrifying biomasses; however, no specific source has been determined. NTP receives large quantities of chemical toilet waste that are discharged to the plant in slug doses via tank truck to

the septage receiving station. Investigation into quaternary ammonium compounds (QACs) which were contained in the chemical toilet waste suggested that QACs at higher concentrations caused some inhibition to NOB activity, but no significant impact on AOB activity.

4.1 Introduction

The Hampton Roads Sanitation District (HRSD) operates thirteen treatment plants in the Hampton Roads, Virginia, area with a combined capacity of 231 million gallons per day (mgd) (Bilyk et al., 2008). The Nansemond Treatment Plant (NTP) is one of the larger facilities operated by HRSD and was designed to treat 30 mgd (max monthly) using a 3-stage Virginia Imitative Process (VIP) biological nutrient removal (BNR) process (Figure 4.1 and 4.2) (Bilyk et al., 2008). The majority of the influent is domestic, but there is also a large industrial contribution, particularly from a pig processing facility, landfill leachate, and significant loads from septage and grease deliveries (Bilyk et al, 2008). NTP has had a long history of issues with nitrification after its initial upgrade to a BNR facility in 1998.

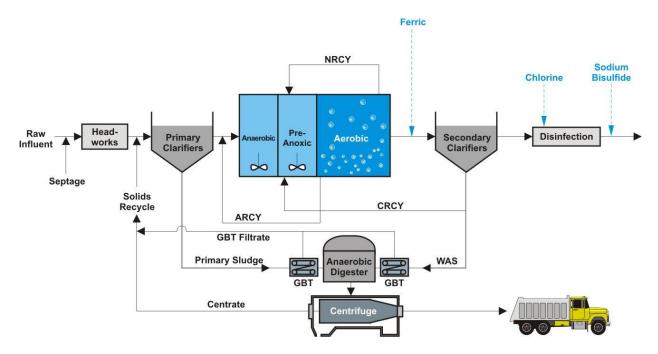


Figure 4.1 3-Stage VIP Process at NTP in 2008 (Bilyk et al., 2008).

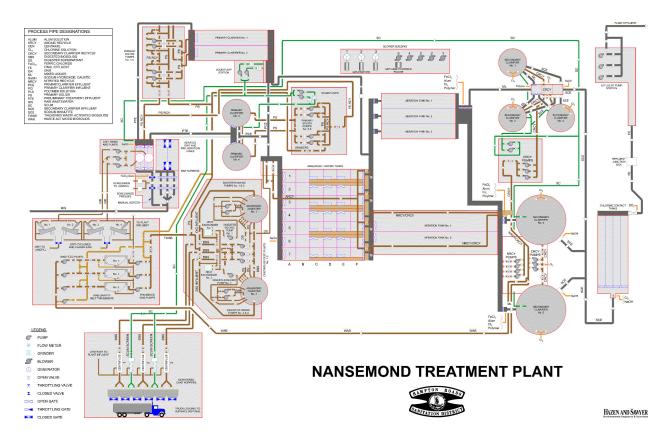


Figure 4.2 NTP 3-Stage VIP Process Plant Layout

Starting in 2001 until the present the plant has experienced continuous and sporadic nitrification upsets without much explanation. This determination was based upon the plant's efficiency in removing nitrogen. The VIP plant in Norfolk, VA which also employs the VIP process began BNR operations approximately seven (7) years prior to similar Nansemond operations (Balzer et al., 2005). Nitrogen removal was similar at both facilities (Table 4.1) during the first two years of side-by-side operation (1999 & 2000) (Balzer et al., 2005). Nansemond experienced an unexpected decline in nitrogen removal efficiency starting in 2001 which has continued to the present (Balzer et al., 2005). This deficiency in performance has been variable and has not been consistent from 2001 to the present time.

Table 4.1 Percent TN Removal for HRSD's Nansemond and VIP Treatment Plants (1999-2009)

PLANT	NP	VIP
YEAR	[% Removal]	[% Removal]
1999	69.9	71.5
2000	64.2	67.0
2001	53.1	66.3
2002	64.4	67.1
2003	45.0	62.5
2004	55.6	71.9
2005	50.6	67.83
2006	70.8	69.66667
2007	68	65.83333
2008	70.3	72.58333
2009	50	60
MEAN	60.2	67.5

NTP achieves continuous complete nitrification during certain periods consistently meeting desired effluent limits. In spite of this, NTP has also experienced unexplained sporadic nitrification upsets for a number of years when compared to the VIP plant (Figure 4.3 and 4.4) and some indication of continuous nitrification inhibition, as demonstrated by calibration of a process simulation model to historical plant performance data. The previous target total nitrogen (TN) treatment objective was 12 mg/L on a seasonal basis. This has changed more recently to a permitted limit of 8 mg/L TN on an annual average basis (Bilyk et al., 2008).

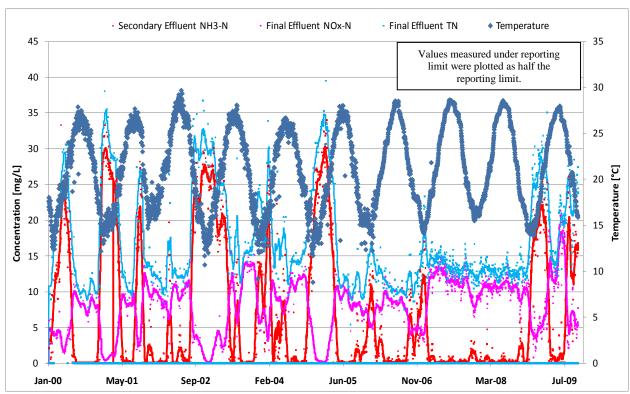


Figure 4.3 NTP Historical Effluent Ammonia, NOx-N, and Total Nitrogen Profile 2000-2009 (Lines represent 30-day rolling averages).

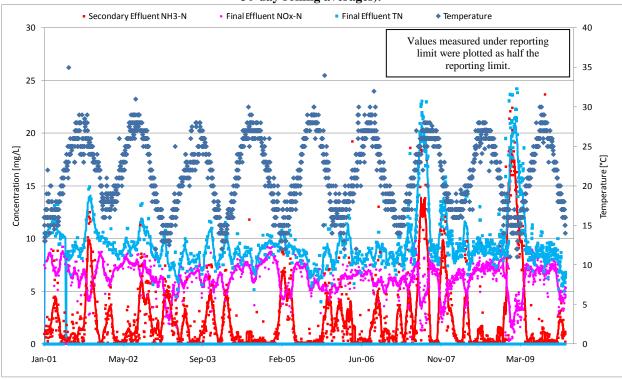


Figure 4.4 VIP Historical Effluent Ammonia, NOx-N, and Total Nitrogen Profile 2001-2009 (Lines represent 30-day rolling averages).

In addition the plant also experienced regular biological phosphorus removal upsets (Figure 4.5) which forced NTP to add large quantities of ferric chloride to avoid permit violations for effluent TP (Bilyk et al., 2008). It was the goal of NTP to meet a treatment objective of 1 mg/L of total phosphorus (TP) on an annual average basis (Bilyk et al., 2008).

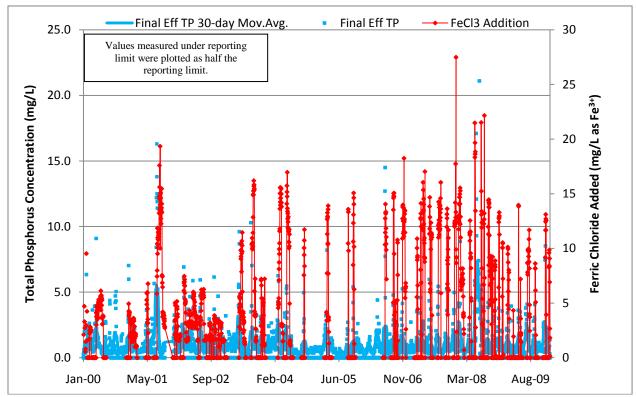


Figure 4.5 NTP Historical Effluent TP and Ferric Chloride Addition 2000-2009

This presented an issue as NTP along with other treatment facilities which discharge to the James River are required starting in 2011 to meet a combined annual discharge limit of 6 million pounds of total nitrogen (TN) (Balzer et al., 2005). Investigation as to possible contributors of nitrification inhibition has attributed the sources including industrial discharges (hog processing plant or landfill leachate) and truck-delivered waste received at the facility septage, grease, and chemical toilet waste. NTP receives large quantities of chemical toilet waste that are discharged to the plant in slug doses via tank truck to the septage receiving station (see

Tables 4.4 and 4.5). It was hypothesized that the quaternary ammonium compounds (QACs) contained in this chemical toilet waste could be a possible source for the sporadic nitrification inhibition that is experienced at the NTP.

Quaternary ammonium compounds (QACs) or quaternary ammonium salts (quaternary ammines) are salts of quaternary ammonium cations with a coordinating anion (e.g. chloride). They are organic compounds that contain four functional groups attached covalently to a positively charged central nitrogen atom (R₄N+). These functional groups (R) include at least one long chain alkyl group, and the rest are either methyl or benzyl groups. QACs are extensively used in domestic and industrial applications as surfactants, emulsifiers, fabric softeners, disinfectants and corrosion inhibitors (Tezel et al, 2007). QACs are of importance because it has been determined that acute inhibition of both heterotrophic COD removal and nitrification, especially the nitrite oxidation process, can be inhibited at high concentrations (Kreuzinger et al, 2007).

In order to meet future permit limits NTP is currently being upgraded to a 5-stage Bardenpho process (Figure 4.6). This will increase aeration volume/capacity and enhance both nitrification and Bio-P removal. The new upgrades will also incorporate a full-scale proprietary technology developed by Ostara that uses a fluidized bed reactor to recover phosphorus and ammonia through struvite precipitation from the centrate being generated at NTP. The harvested struvite can then be utilized as a slow release fertilizer (Ostara, 2007).

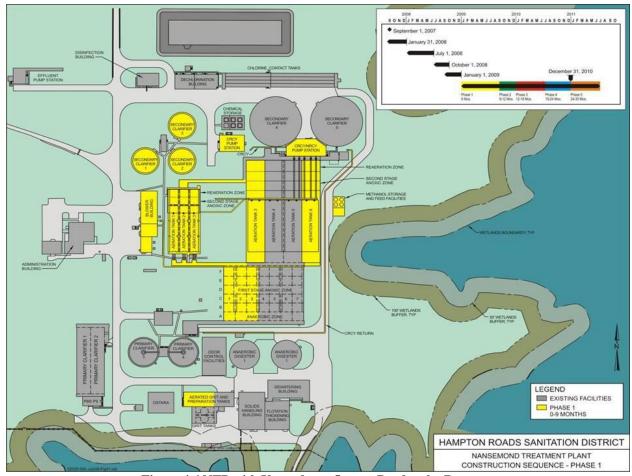


Figure 4.6 NTP with Upgrades to 5-stage Bardenpho Process.

A preliminary engineering report by Hazen and Sawyer, P.C. for the current nutrient removal upgrade, suggested that the ammonia oxidizing bacteria (AOB) maximum specific growth rate $(\mu_{max, AOB})$ be reduced from the default value of 0.90 days⁻¹ to 0.57 days⁻¹ to account for high effluent ammonia data during the calibration period (Hazen & Sawyer, 2007). This has significant implications in terms of the required aeration volume for consistent nitrification at cold temperatures. It was originally assumed that NTP would nitrify year-round with this background inhibition. This would only be made possible however, by reconfiguring the BNR process to allow the 2^{nd} anoxic zone to operate aerobically (swing zone). Removing an anoxic zone would reduce overall TN removal because the process would essentially change from the 5-stage Bardenpho process to an A2O Process. If the inhibition can be eliminated then there would

no longer be a need to configure the 2nd anoxic zone to run aerobic in cold temperature conditions, providing improved annual average TN removal. This would ensure that NTP and the other James River HRSD facilities meet the TN bubble permit limit with more certainty. The objectives of this study were the following:

- Establish the inhibitory characteristics of the NTP influent wastewater by bench-scale experiments independently evaluating ammonia oxidizing bacteria (AOB) and nitrite oxidizing bacteria (NOB) rates.
- Study and evaluate possible sources of inhibition from the industrial loads and chemical toilet waste received suspected of containing QACs and their affect on nitrification kinetics.

4.2 Methodology

4.2.1 Sample Collection

Batch reactor experiments were conducted by collecting biomass samples from the aeration basin effluent of several full-scale wastewater treatment plants. Grab samples were collected unless otherwise arranged with HRSD to collect composites. All samples were collected and stored in polypropylene carboys or similar containers. These samples were then transported to the lab and stored until use with continuous aeration for no longer than approximately 48 hours at room temperature. Biomass samples were collected from NTP; the HRSD VIP plant, which uses the same BNR process as NTP but does not experience inhibition; the Henrico County WWTP, which has a fully nitrifying 5-stage BNR process; and the HRSD York River plant from the sequencing batch reactor system.

Wastewater sources for experiments were also collected via grab sample or composite sampling by HRSD based on the source being tested. Wastewater samples were collected

including NTP and VIP primary effluent (PE); NTP and VIP secondary effluent (SE); raw wastewater from isolated branches of the NTP collection system; and industrial waste samples from a hog processing plant and a landfill leachate stream. Waste samples were also tested from other suspected sources such as chemical toilet disinfectants (quaternary ammonium compounds).

4.2.2 Batch Reactor Construction

Four parallel batch reactors were constructed of plexiglass and configured with vinyl and polypropylene fittings and appurtenances. The first reactor served as the control while operating conditions of the other three reactors were varied. Each reactor had a volume of 3 liters with individual ports for sampling and chemical addition (acetate, ammonia, nitrite, and phosphate) during experiments as needed. A diffuser stone was incorporated into each reactor to provide air/oxygen or nitrogen based on the experiment. This allowed the reactors to be operated under aerobic, anoxic, or anaerobic conditions for nitrification, denitrification, and phosphorus release/uptake measurements. The reactors were configured to be air tight with a water seal on the headspace for anaerobic conditions. Mixing was provided through magnetic stir-bars and stir plates. Dissolved oxygen (DO) was kept constant using HACH LDO probes with HQ40d meters. Two probes were connected to a single HQ40d meter (total of four probes and two meters). National Instruments LabView 7.0 DO Controller was used to provide on and off control of four solenoid air supply valves with associated relays. pH was controlled and logged using Cole-Parmer pH meters/controllers connected to acid (1M H₂SO₄) and base (3M Na₂CO₃) pumps. The reactors were placed in a water bath with a circulating chiller and immersion heaters to maintain a constant temperature during experiments. Refer to Figure 4.7, 4.8, and 4.9 for a schematic and

image of an individual reactor and for a complete schematic and image of the entire reactor configuration.

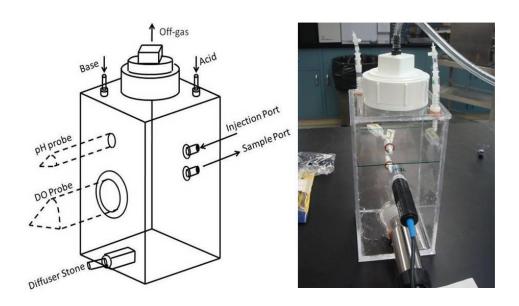


Figure 4.7 Reactor Schematic and Image

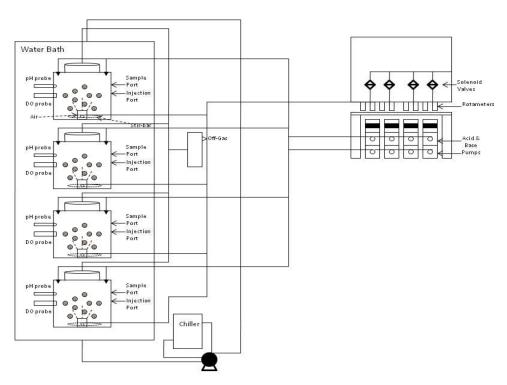


Figure 4.8 Complete Reactor System Schematic

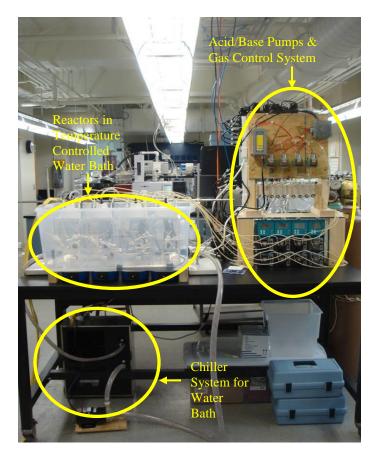


Figure 4.9 Complete Reactor Set-Up

4.2.3 Batch Reactor Operation

Experiments consisted of the evaluation AOB, NOB, and Bio-P activity. The method for evaluating each type of experiment is discussed further in subsequent sections. Temperature for experiments was based on the temperature of the biomass sample when it was collected onsite via grab sample. The matrix of experiments developed and completed is summarized in Table 4.2.

Table 4.2 Batch Experiment Matrix

Week #	Biomass Source	Di	Experiments		
1	Nansemond	a. NTP/PE	b. VIP/PE	AOB, NOB, Bio-P	
	VIP	c. VIP/PE	d. NTP/PE		
2	Nansemond	a. NTP/PE	b. VIP/PE	AOB, NOB, Bio-P	
	VIP	c. VIP/PE	d. NTP/PE		
3	Nansemond	a. VIP/SE + Leachate	b. VIP/SE	AOB, NOB, Bio-P	
	VIP	c. VIP/SE + Leachate	d. VIP/SE		
4	Nansemond	a. NTP/PE	b. HR/PE	AOB, NOB	
	Henrico	c. HR/PE	d. NTP/PE		
	Nansemond	a. VIP/SE + Hog Plant	b. VIP/SE	AOB, NOB, Bio-P	
5	VIP	c. VIP/SE + Hog Plant	d. VIP/SE		
	Nansemond	a. VIP/SE + Leachate	b. VIP/SE	AOB, NOB	
6	VIP	c. VIP/SE + Leachate	d. VIP/SE		
	Nansemond	a. VIP/SE	b. VIP/SE + Branch 1	AOB, NOB	
7	VIP	c. VIP/SE + Branch 2	d. VIP/SE + Branch 3		
8	Nansemond	a. VIP/SE	b. VIP/SE + Branch 2a	AOB, NOB	
	VIP	c. VIP/SE + Branch 2b	d. VIP/SE + Branch 3		
	Nansemond	a. VIP/SE	b. VIP/SE + Cedar Ln PS	AOB, NOB	
9	VIP	c. VIP/SE + Gum Rd. PRS	d. VIP/SE + Pughsville PRS		
	Nansemond	a. VIP/PE	b. NTP RWI	AOB, NOB	
10	VIP	c. NTP PCI	d. NTP PCE		
1.1	Nansemond	a. VIP/SE	b. VIP/SE + 20 mg/L Fe	AOB, NOB	
11	VIP	c. VIP/SE + 35 mg/L Fe	d. VIP/SE + 50 mg/L Fe		
12	Nansemond	a. VIP/SE	b. VIP/SE + 20 mg/L Fe	AOB, NOB	
12	VIP	c. VIP/SE + 35 mg/L Fe	d. VIP/SE + 50 mg/L Fe		
12	York River (YR)	a. YR/SE	b. YR/SE + 20 mg/L Fe	AOB, NOB	
13	York River (YR)	c. YR/SE + 35 mg/L Fe	d. YR/SE + 50 mg/L Fe		
14	York River (YR)	a. YR/PE	b. NTP RWI	AOB	
	York River (YR)	c. NTP PCI	d. NTP PCE		
1.5	York River (YR)	a. YR/SE	b. YR/SE + 0.574 mL Chem Toilet Additive*	AOB, NOB	
15	York River (YR)	c. YR/SE + 13.96 mL Chem. Toilet Additive*	d. YR/SE + 62.84 mL Chem Toilet Additive*		
16	York River (YR)	e. YR Mixed Liquor (ML)	f. YR ML + 15 mL Product B	AOB, NOB	
16	York River (YR)	g. YR ML + 30 mL Product B	h. YR ML + 60 mL Product B		
17	York River (YR)	e. YR Mixed Liquor (ML)	f. YR ML + 60 mL Product B	- AOB, NOB	
	York River (YR)	g. YR ML + 120 mL Product B	h. YR ML + 180 mL Product B		

^{*}NOTE: Week 15 Experiments – Two different Chemical Toilet Disinfectants (Product A & B) used for the two different days of experiments.

Table Abbreviations:

NTP – Nansemond Treatment Plant HR – Henrico County Treatment Plant PE Primary Effluent

PE – Primary Effluent ML – Mixed Liquor

VIP – Virginia Initiative Process Treatment Plant YR – York River Treatment Plant SE – Secondary Effluent RWI – Raw wastewater influent

4.2.3.1 AOB Experiments

AOB activity was evaluated by spiking the reactors with ammonia and monitoring NO_x-N and NO₃-N generation rate over three hours (Kelly et al, 2004). During an AOB experiment the collected biomass was concentrated to approximately 8000 mg/L MLSS prior to experiments. Then the reactor was filled 1/3 by volume with biomass and 2/3 by volume with a diluent (primary effluent, waste sample, etc.). Additional ammonia was added to achieve a total ammonia concentration of approximately 40 mg/L NH₄-N to provide sufficient ammonia during experiments to prevent depletion. Dissolved oxygen (DO) and pH were constantly monitored and maintained within desired limits using online control systems. DO was maintained between 5-7 mg/L during experiments with a pH between 7.0-7.2. Each AOB experiment was conducted for 3 hours with 5 samples collected evenly throughout this period of time. Samples were immediately filtered through 0.45μm membrane filters after collection. AOB activity was based on NO_x-N generation rate.

4.2.3.2 NOB Experiments

NOB activity was assessed by adding in nitrite (ensuring ammonia concentrations were low) and monitoring the rate of NO₂-N depletion and NO₃-N generation. NOB experiments was conducted very similar except after the reactors were filled by volume as mentioned above they were aerated for approximately one hour to allow uptake of any residual ammonia. Samples were taken prior to starting the NOB Experiments and analyzed for NH₄-N to ensure that the ammonia concentration in each reactor was <1 mg/L NH₄-N (detection limit of HACH TNT 831 NH₄-N analysis method). After it was determined that the ammonia concentration in each reactor was depleted, approximately 25 mg/L NO₂-N was added, and then 5 samples were collected evenly

over a 3 hour test period. Samples were immediately filtered through 0.45µm membrane filters after collection. DO and pH were maintained within the same parameters as was used for the AOB experiment. NOB activity was based on NO₃-N generation rate and NO₂-N uptake rate.

4.2.3.3 Bio-P Experiments

Bio-P activity was determined by monitoring P uptake and release rates. Experiments was conducted using the same concentrated biomass as in the AOB & NOB experiments (~8000 mg/L MLSS). Experiments were conducted in three phases: uptake, release, and a second uptake. During the first uptake (aerobic) phase, each reactor was filled with 1/3 by volume of biomass and 2/3 by volume of PE or another tested wastewater source. Then each reactor was spiked with 5 mg/L of PO₄-P to reach a target of 20 mg/L P in each reactor since it was presumed approximately 15 mg/L of PO₄-P would be present from the biomass and PE/wastewater sample. The first uptake phase was conducted for 1-2 hours and 3 samples were collected during this time with immediate filtration using 0.45µm membrane filters. During both uptake phases, pH was always controlled between 7.0-7.2 while the DO was maintained between 5-7 mg/L. After this initial uptake phase, the reactors were deaerated by sparging nitrogen (N₂) gas and reducing the DO to 0 mg/L which was maintained throughout the release phase. After the DO was reduced to 0 mg/L, each reactor was spiked with 200 mg/L acetate as COD. It was desired to have all NO₃-N and NO₂-N depleted, but this was not always the case. pH was controlled the same as in the uptake phase experiment and 5 samples were collected over a 1 hour period with immediate filtration using 0.45µm membrane filters. After the 1 hour release phase nitrogen was shut-off and then air/oxygen was sparged into the reactors to once again raise the DO and maintain the DO between 5-7 mg/L. As soon as the air/oxygen sparge began, 5 samples were collected over a 1-2 hour period with immediate filtration using 0.45µm membrane filters. All the samples from

both uptake phases and the release phase were analyzed using the same methods used for the AOB and NOB experiments noted in the subsequent 3.3 Analytical Methods section. Rates based on PO₄-P release and uptake was normalized to MLVSS concentrations for each experiment.

All experiments was conducted in this manner except for that conducted during weeks 16 & 17, during which the entire reactor volume was filled with biomass due to the biomass from the York River plant at that time being very dilute.

4.2.4 Analytical Methods

Samples were analyzed for NH₄-N, NO₂-N, NO₃-N, and PO₄-P using HACH colorimetric test kits, ion chromatography (IC), and flow injection analysis (FIA) (see subsections 3.3.3, 3.3.6, 3.3.7, 3.3.8). MLSS and MLVSS were performed at the end of the experiments per *Standard Methods* (APHA et al., 1998). Nitrate production rates, phosphorus uptake and release rates were calculated by normalizing the slope of each chemical constituent by the MLVSS. Normalizing the slope refers to the calculated slope being divided by the mixed liquor volatile suspended solids and converted into the appropriate units of mg/g MLVSS/hr.

4.2.4.1 Ammonia (NH₃-N) HACH Test Kit

Ammonia was analyzed using HACH Test N' Tube (TNT) 831 kit and HACH DR2800 spectrophotometer. This method uses the salicylate method, whereby ammonium ions react with hypochlorite and salicylate ions in the presence of sodium nitroprusside to as a catalyst form indophenol. The amount of color formed is directly proportional to the ammonia nitrogen present.

4.2.4.2 Nitrate (NO₃-N) HACH Test Kit

Nitrate was analyzed using HACH TNT 835 kit. This kit incorporates the dimethylphenol method where nitrate ions in solution with sulfuric and phosphoric acids react with 2,6-dimethyphenol to form 4-nitro-2,6-dimethyphenol.

4.2.4.3 Nitrite (NO₂-N) HACH Test Kit

Nitrite was analyzed using NitriVer3 Nitrite Reagent Powder Pillows and 10 mL sample vials. This kit uses the diazotization method where nitrite in the sample reacts with sulfanilic acid to form a intermediate diazonium salt. This salt combined with chromotropic acid forms a pink color which is directly proportional to the amount of nitrite present.

4.2.4.4 Ortho-Phosphate (PO₄-P) HACH Test Kit

Ortho-Phosphate was analyzed using the HACH Reactive Phosphate TNT Reagent Kit.

The kit analyzed 5 mL of sample in a vial. This test kit uses the USEPA PhosVer3 method where orthophosphate reacts with molybdate in an acid to produce a mixed complex. Ascorbic acid then reduces this complex, producing an intense blue color.

4.2.4.5 Ion Chromatography

IC analysis was conducted for measuring nitrate only for the batch experiments work. The IC was used in conjunction with conductivity detection (Dionex ICS-1000), an AS14A analytical column and AG14A guard column, an ASRS conductivity suppressor, and an eluent flow of 1.0 mL/min of 1.0 mM NaHCO₃ and 8.0 mM Na₂CO₃.

4.2.4.6 Flow Injection Analysis

FIA was conducted for measuring nitrite, nitrate, and phosphate for batch experiments work using a SEAL Analytics auto-analyzer system. FIA analysis incorporated a volume of 2 mL of sample in sample vials and placed into a sample tray. The tray was then placed into the FIA instrument along with various reagents required for the different analyte test methods. A schedule was then configured using the associated computer software with the instrument and

analyzed. QA/QC samples were measured every 10 samples and test methods were always standardized prior to analysis of actual samples. Data output was provided in concentrations in mg/L as NO_2 -N, NO_3 -N, and PO_4 -P.

4.3 Results and Discussion

4.3.1 AOB and NOB Batch Rate Results

AOB and NOB kinetics were independently evaluated based on NOx-N production (AOB), NO₃-N production (NOB), and NO₂-N consumption rates (NOB). These rates were calculated based on a linear regression of the change in nitrogen species concentration over the experimental period (Figure 4.10). These slopes were divided by the mixed liquor volatile suspended solids (MLVSS) to normalize the rate to the biomass concentration, referred to here as the "specific" rate. The representative experiment shown below in Figure 4.10 plots the measured nitrogen species, NH₄-N, NO₂-N, NO₃-N, NO_x-N (NO₂-N + NO₃-N), versus time. Batch experiments were conducted for a total of 17 weeks. Results from the initial batch-rate testing (Weeks 1-4) confirmed the inhibitory characteristics of NTP wastewater. Whenever NTP wastewater was combined with a stable nitrifying biomass there was an observed decrease in NO_x-N production, nitrate production, and nitrite consumption rates (Figure 4.11).

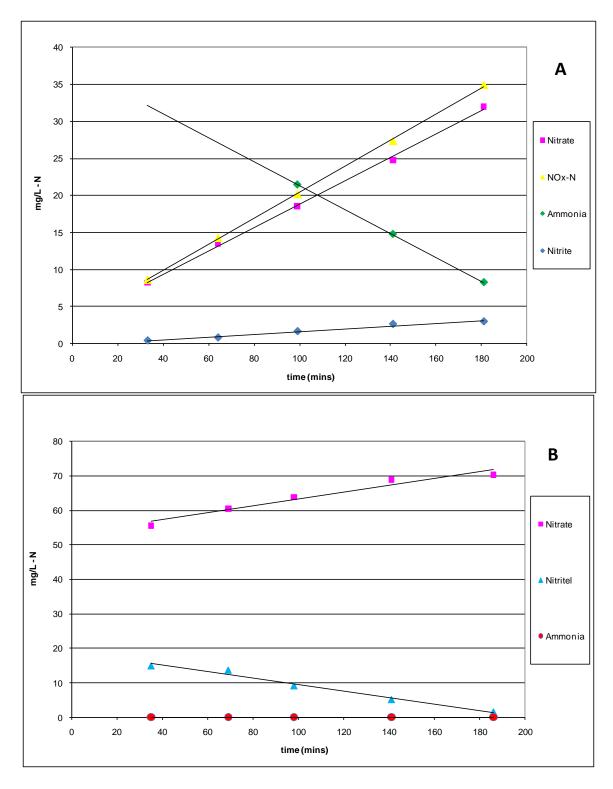


Figure 4.10 Representative Experiment – a. AOB experiment, b. NOB experiment

The data presented from the Week 4 experiment showed some inhibition when the Henrico County (HR) WWTP biomass (control) was combined with NTP PE compared to HR

PE. Although this was true, evaluation of the 95% confidence intervals between the regression slopes revealed that it was not statistically significant as only five data points were collected. This did not provide enough data points to evaluate the regression slopes, but data such as that presented in Figure 4.11 was repeatedly seen through multiple experiments. This being the case, it was determined that NTP PE exhibited some inhibitory characteristics. When NTP biomass was combined with HR PE there was little observed improvement in AOB and NOB activity, likely because the NTP biomass had already been exposed to some degree of inhibition.

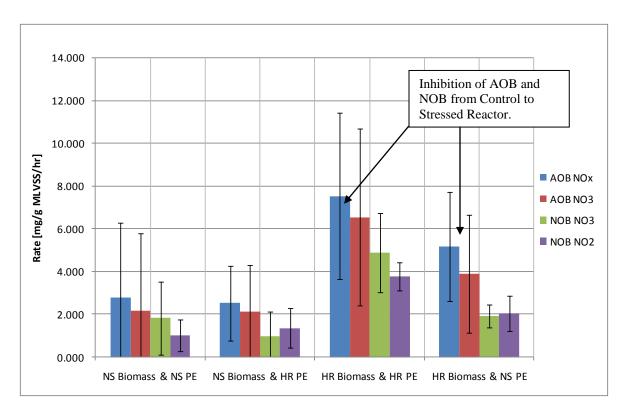


Figure 4.11 Comparisons of AOB & NOB Rates with Control (Week 4 – NTP and Henrico County). (Error bars represent the 95% confidence interval of the regression slope.)

There were several suspected wastewater sources were tested (Table 4.4). Initial testing of inhibitors was for the waste received from a hog processing facility and leachate from a landfill. Results from the testing of both wastewater sources provided no significant differences in AOB and NOB activity compared to the control (Figure 4.12 and 4.13). This was determined

based on the difference between the 95% confidence intervals of the regression slopes. Since only five data points were collected during the experiments, the error in the slopes were much greater; however, through multiple experiments of other wastewater sources similar results were observed. This supported that there was no significant difference in AOB and NOB activity from these suspected sources of inhibition. It should be noted that experiments on both of these waste streams were repeated producing similar results. In testing the two waste streams, biomass from the VIP plant was used as the control to compare to the performance of NTP biomass.

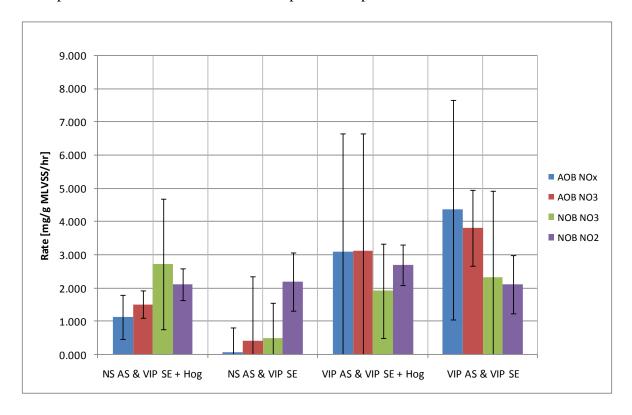


Figure 4.12 Hog Processing Plant Rate Comparison. (Error bars represent the 95% confidence interval of the regression slope.)

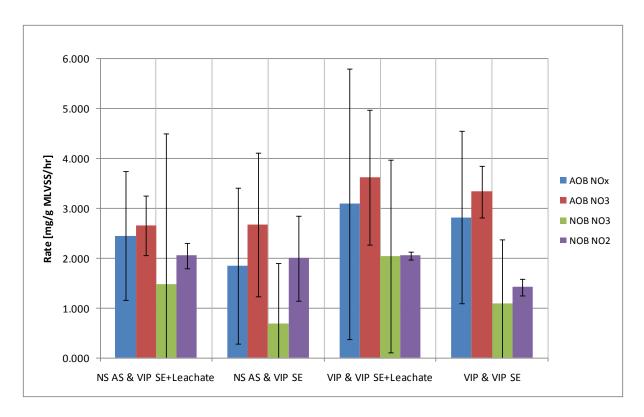


Figure 4.13 Landfill Leachate Rate Comparisons. (Error bars represent the 95% confidence interval of the regression slope.)

Experiments with other wastewater sources generated similar results as the landfill leachate and hog processing plant. There were no significant signs of inhibition from any of the tested waste streams listed in Table 4.2 based on similar results from multiple experiments, in spite of the lack of difference in the 95% confidence interval of the slopes.

Week 14 experiments were conducted to evaluate the impact of internal recycle streams within NTP. The tested sources included the raw wastewater influent (RWI), primary clarifier influent (PCI), and primary clarifier effluent (PCE). The tested samples were composite samples collected over a four day period (Monday – Thursday) by HRSD. Two duplicate AOB experiments were conducted for each day of composite sample to examine NO_x-N (NO₂-N + NO₃-N) and NO₃-N generation for a total of eight experiments (Figure 4.14). Some mild signs of inhibition were observed between the control and other reactors. The changes in AOB rates between the different tested sources (RWI, PCI, PCE) however, were not significantly different.

This suggested that the inhibitory source was in the plant raw influent and not an internal recycle stream issue. Graphical representation suggested that the York River Biomass was affected negatively by the addition of the Nansemond RWI when compared to the Nansemond PCI and PCE. For this reason a linear regression was performed on the NOx-N rate for one of the days to examine the 95% confidence in the slopes from the different reactors (Table 4.3). The highlighted values represent the 95% confidence interval on the regression slopes between Reactor A (York River Biomass & York River PE) and Reactor B (York River Biomass & Nansemond RWI). The results suggested that the differences in slopes were not statistically different at the 95% confidence level, but were very close to this limit. This inferred that it may not have been a recycle stream issue, but it could have been an insignificant amount of truck delivered septage or grease received during the sampling week.

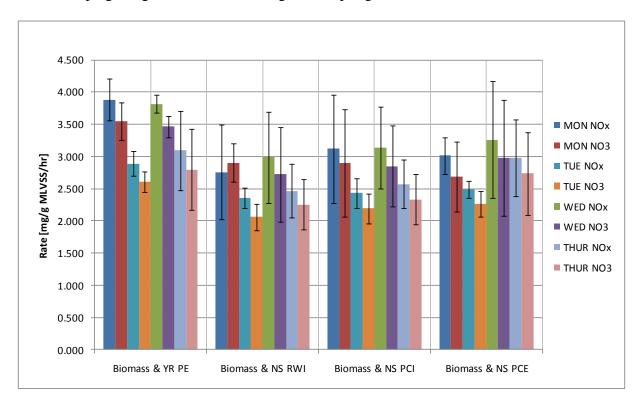


Figure 4.14 AOB Comparisons for Composite Samples of NTP Internal Recycle Streams (Week 14). (Error bars represent the 95% confidence interval of the regression slope.)

Table 4.3 Linear Regression for Calculation of 95% Confidence

MONDAY COMPOSITE SAMPLE							
Reactor	Lower 95%	Slope (Regression)	Upper 95%				
_							
Α	0.0544	0.0593	0.0643				
В	0.0334	0.0457	0.0579				
С	0.0371	0.0507	0.0643				
D	0.0449	0.0495	0.0541				
	TUESDAY C	OMPOSITE SAMPLE					
Reactor	Lower 95%	Slope (Regression)	Upper 95%				
_							
Α	0.0412	0.0441	0.0470				
В	0.0364	0.0390	0.0416				
С	0.0358	0.0396	0.0433				
D	0.0388	0.0409	0.0431				
	WEDNESDAY	COMPOSITE SAMPLE					
Reactor	Lower 95%	Slope (Regression)	Upper 95%				
Α	0.0435	0.0452	0.0468				
В	0.0310	0.0407	0.0503				
С	0.0326	0.0408	0.0490				
D	0.0305	0.0424	0.0542				
	THURSDAY COMPOSITE SAMPLE						
Reactor	Lower 95%	Slope (Regression)	Upper 95%				
Α	0.0293	0.0366	0.0439				
В	0.0279	0.0336	0.0392				
С	0.0285	0.0335	0.0384				
D	0.0309	0.0387	0.0465				

4.3.2 Bio-P Batch Rate Results

Bio-P experiments were only conducted in conjunction with AOB and NOB experiments for four of the seventeen weeks. Bio-P experiments generated varied results. The varied results can be attributed to the fact that NTP was recovering from Bio-P upsets (FeCl₃ addition) when samples were collected. There was a noticeable effect on Bio-P performance of other biomass sources when combined with NTP PE. It was observed that when a stable biomass, in this case

VIP biomass, was combined with NTP PE there was a small decrease in phosphate uptake rate (Figure 4.15).

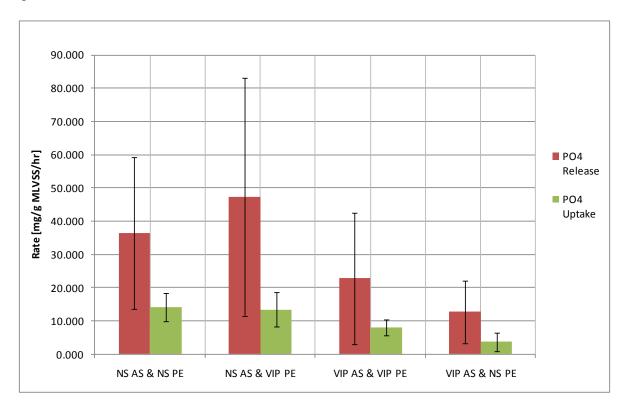


Figure 4.15 Bio-P Experiment with NTP and VIP Biomass and Primary Effluent. (Error bars represent the 95% confidence interval of the regression slope.)

In some cases when biomass samples were collected ferric chloride was being added to assist in phosphorus removal at NTP. For this reason experiments were also conducted examining varying ferric chloride concentrations on nitrification (Figure 4.16 and 4.17).

Experiments with ferric chloride addition were conducted for a total of three weeks (Week 11, 12, and 13). Week 11 and 12 were carried out using VIP biomass and week 13 using York River biomass. With appropriate control of pH after Fe³⁺ addition, there were no significant changes between the different concentrations of ferric chloride added (as Fe³⁺). This suggested that there was no impact of ferric chloride addition on nitrification rates for both AOB and NOB, even at very high concentrations.

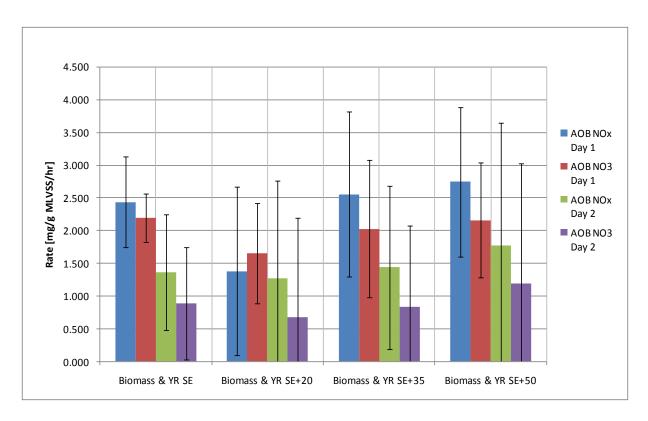


Figure 4.16 AOB Comparison with Ferric Chloride Addition (Week 13). (Error bars represent the 95% confidence interval of the regression slope. 20, 35, and 50 = mg/L as Fe³⁺added to each reactor.)

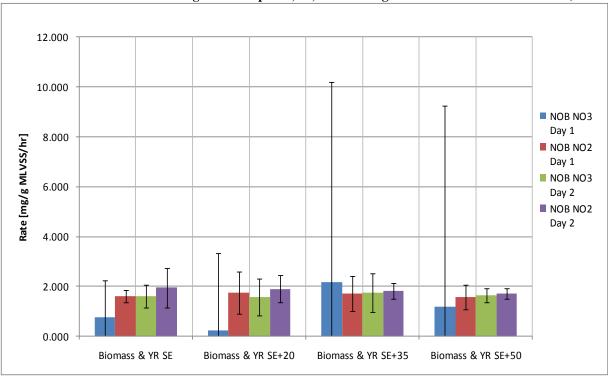


Figure 4.17 NOB Comparisons with Ferric Chloride Addition (Week 13). (Error bars represent the 95% confidence interval of the regression slope. 20, 35, and 50 = mg/L as Fe^{3+} added to each reactor.)

4.3.3 QAC Batch Rate Testing

Three experiments were performed examining chemical toilet disinfectants suspected of containing QACs. Two different chemical toilet disinfectants were used for testing, Product A and Product B; however, Product A was found to contain no QACs. Various assumptions were made when determining concentrations of chemical toilet disinfectant to add to each stressed reactor. Based on the large quantity of chemical toilet waste delivered to NTP (Table 4.4 and 4.5) it was decided to use three different assumptions for calculating experiment doses.

- NTP received trucked waste deliveries all during one month of the year, Monday –
 Friday only, during the normal 8 hour workday.
- 2. NTP received waste during two months of the summer and received the waste over 40 days, Monday Friday, in 5 hour periods throughout the 8 hour workday.
- 3. NTP received waste during summer months (June, July, August) when use of portable toilets was at a peak, receiving the waste over 60 days in 6 hour periods throughout the 8 hour workday.

Table 4.4 Septage Waste Received for Several HRSD WWTPs

Wastewater	Residential Septage	Chemical Toilet Waste	Total Delivered Waste	Average Daily Flow	Chemical Toilet Waste	Total Septage
Treatment Plant	[gallons/year]	[gallons/year]	[gallons/year]	[MGD]	[gallons/day]	[gallons/day]
Atlantic	501,937	560,258	1,885,655	40	1,535	5,166
Boat Harbor	90,762	852,656	1,618,515	14.9	2,336	4,434
Chesapeake-Elizabeth	454,906	369,426	1,284,278	13.2	1,012	3,519
Nansemond	1,800,999	1,291,602	4,427,292	18.5	3,539	12,130
Williamsburg	1,549,416	272,326	4,000,171	15.7	746	10,959
York River	2,592,031	76,418	2,730,400	12.9	209	7,481

Table 4.5 Septage Waste Received as a Fraction of Average Daily Plant Flow

WWTP Loading	Chemical Toilet Waste	Chemical Toilet Waste	
Highest to Lowest	[gallons/day]	[% Avg Day Flow]	
Nansemond	3,539	0.019	
Boat Harbor	2,336	0.016	
Chesapeake-Elizabeth	1,012	0.008	
Williamsburg	746	0.005	
Atlantic	1,535	0.004	
York River	209	0.002	

Results from the experiments showed no significant signs of inhibition or variability in concentration of Product A (Figure 4.18).

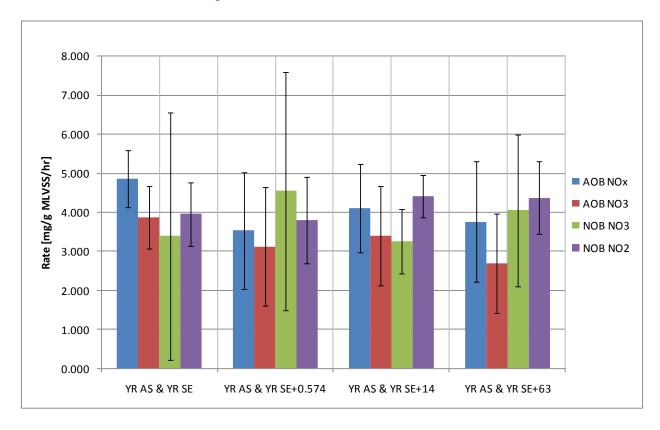


Figure 4.18 QAC Experiment with Product A (Week 15). (Error bars represent the 95% confidence interval of the regression slope. 0.574, 14, 63 = mL of Product A added to reactor.)

It was observed in Product B experiments containing QACs, that there was a mild reduction in NOB activity at higher concentrations (Figure 4.19). This was true for the first day of experiments, but was not replicated in the second day of experiments. The decrease in NOB activity however, was consistent with literature which suggested that NOB exhibit greatest inhibition in response to QAC loading (Kreuzinger et al, 2007). Therefore it was proposed that Product B at higher concentrations may cause some inhibition of nitrification.

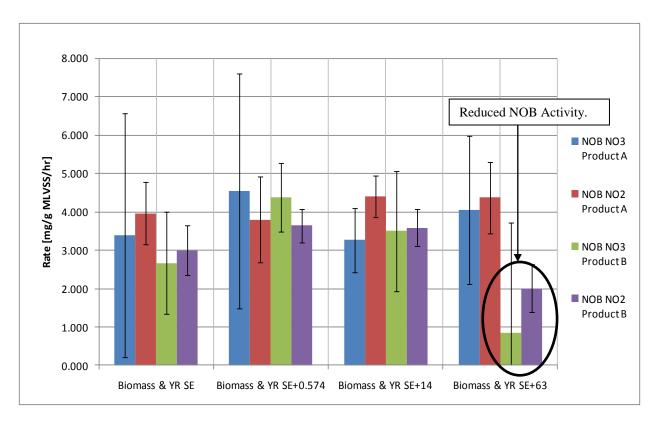


Figure 4.19 Chemical Toilet Disinfectants containing QACs (Week 15). (Error bars represent the 95% confidence interval of the regression slope. 0.574, 14, 63 = mL of Product added to reactor.)

Further experiments in the later two weeks (week 16 and 17) using Product B generated different results from week 15 experiments. It was desired in these experiments to use Product B at higher concentrations to show more significant inhibition of NOB or some reduction in AOB rates. Results however, from week 16 and 17 showed very slight inhibition in rates at higher concentrations, but with AOB and not NOB rates (Figure 4.20). In some cases performance was better or similar with the addition of QACs.

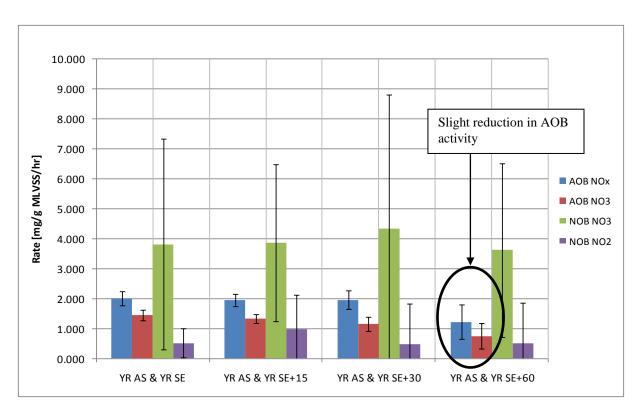


Figure 4.20 Experiments with Chemical Toilet Additive containing QACs (Week 16). (Error bars represent the 95% confidence interval of the regression slope. 15, 30, and 60 = mL of Product B added to reactor.)

4.4 Conclusion

Based on the batch experiments, no significant inhibitor was identified as the cause of sporadic nitrification problems at NTP; the batch-rate testing did show that NTP PE was consistently inhibitory on a seemingly continuous basis. The H&S model calibration based on historical plant data suggested a reduction in AOB μ_{max} from 0.9 days⁻¹ to 0.57 days⁻¹, which supported the finding that there was some degree of continuous inhibition. It was also determined that wastewater derived from the hog processing plant and the previously suspected landfill leachate did not cause inhibition of nitrification or bio-P at least at the time the samples were collected. This does not exclude the possibility of an upset or unusual shock load event at one of these industries impacting nitrification at NTP. Evaluation of the various wastewater sources provided an opportunity to study each source independently and its inhibitory

characteristics on both AOB and NOB activity. Although there were large margins of error for some of the tests based on the 95% confidence interval of the regression slopes, data generated from the repeated experiments suggested similar findings. It was also determined that batch experiments were not sufficient to determine the source of sporadic inhibition. For this reason it was planned to reconfigure the bench-scale experiments to continuous sequencing batch reactors (SBRs) which would be run in parallel to the full-scale plant.

4.5 References

American Public Health Association, American Water Works Association, Water Environment Federation. (1998). *Standard Methods for the Examination of Water and Wastewater*; 20th ed.: Washington D.C.

Balzer, B., Cox, M., Deluna, J., Ghosn, S., Hogg, P., Kennedy, G., Pletl, J. 2005. The Status of Biological Nutrient Removal at HRSD's Nansemond Treatment Plant. NITRO Team Report.

Bilyk, K., Cubbage, L., Stone, A., Pitt, P., Dano, J., Balzer, B., 2007. Unlocking the Mystery of Biological Phosphorus Removal Upsets and Inhibited Nitrification at a 30 mgd BNR Facility. *Water Environmental Federation*, 5133-5151.

Boethling, R.S. 1984 Environmental fate and toxicity in wastewater treatment of quaternary ammonium surfactants, *Water Research*, 18 (9), 1061–1076.

Hazen and Sawyer. 2007. Nansemond Treatment Plant Nutrient Reduction Improves Technical Manual.

Kelly, R.T., II, Love, N.G., 2004. A Critical Comparison of Methods Used to Determine Nitrification Inhibition. *Water Environmental Federation*, (15) 166-180.

Kreuzinger, N., Fuerhacker, M., Scharf, S., Uhl, M., Gans, O., Grillitsch, B., 2007. Methodological approach towards the environmental significance of uncharacterized substances — quaternary ammonium compounds as an example. *Desalination*. 215, 209-222.

Ostara. 2007. Proposal: Full Scale Struvite Recovery Project at the Nansemond Wastewater Treatment Facility. *Ostara Nutrient Recovery Technologies*.

Tezel, U., Pierson, J. A., Pavlostathis, S.G. 2007. Effect of polyelectrolytes and quaternary ammonium compounds on the anaerobic biological treatment of poultry processing wastewater. *Water Research*, 41, 1334-1342.

5. Manuscript 2 – Evaluation of the Nansemond Treatment Plant Biological Nutrient Removal Process through Baseline Profile Sampling and Modeling.

Abstract

During the initial upgrades to the Nansemond Treatment Plant (NTP), nitrification was maintained; however, due to available blower capacity during construction it was planned that nitrification would not be maintained during the fall of 2009. In an effort to stop nitrification, the solids retention time (SRT) was purposely reduced over a period of about one month (as wastewater temperature cooled) until additional blower capacity was available. This provided an excellent opportunity to study baseline nitrification kinetics and to evaluate the presence of a source of continuous inhibition. This involved profile sampling via grab samples along the biological nutrient removal (BNR) process starting from the primary effluent to secondary effluent. Samples were immediately filtered upon collection and analyzed on-site for ammonia, nitrite, nitrate, and orthophosphate. Profile sampling was conducted until the plant completely ceased nitrifying in the fall of 2009. The profile sampling data were modeled using *Biowin 3.1* (EnviroSim, Ltd.). Using plant performance data, a calibrated dynamic model was generated over the period of profile samples. These simulations were then compared to data collected during the profile sampling to better estimate the level of continuous nitrification inhibition. Results from the profile sampling showed no apparent indication of sporadic nitrification inhibition. Evaluation of the BNR process also demonstrated that there was little impact on nitrification from nitrite accumulation or sporadic addition of ferric chloride. Modeling of the profile data generated similar profiles; however, there were slight variations as the model predicted nitrification to stop earlier than what was observed. From the modeling, it was also

estimated that the maximum specific growth rate (μ_{max}) of ammonia oxidizing bacteria (AOB) was approximately in the range of 0.50-0.60 days⁻¹, similar to the design value of $0.57d^{-1}$.

5.1 Introduction

An initial investigation of nitrification inhibition was conducted at the Nansemond Treatment Plant (NTP) using bench-scale batch reactor experiments to confirm inhibition of nitrification and then identify possible sources of inhibition in the summer 2008 to summer 2009 (Yi et al, 2010). Through several weeks of batch experiments testing of various industrial sources (hog processing plant and landfill leachate), plant recycle streams, ferric chloride addition, and quaternary ammonium compounds (QACs) was conducted. Batch reactors were used to evaluate nitrification kinetics, both AOB and NOB independently; however, identification of an inhibitor for the nitrification was still inconclusive. NTP was undergoing upgrades from a 3-stage VIP process to a 5-stage Bardenpho process to achieve more stringent nutrient limits. During the initial upgrades to the facility, nitrification was maintained; however, due to available blower capacity during construction it was planned that nitrification would not be maintained during the fall of 2009. In an effort to stop nitrification, the solids retention time (SRT) was purposely reduced over a period of about one month (as wastewater temperature cooled) until additional blower capacity was available. This coincided with the profile sampling creating an opportunity to study nitrification kinetics as the plant stopped nitrifying.

NTP began operations as a 10 mgd secondary treatment plant in 1983. Expansions and upgrades were completed in May 1998, which converted the facility into a 30 mgd BNR facility. Since the upgrade the facility has experienced mixed success in the BNR mode (Balzer et al., 2005). This determination was based upon the plant's efficiency in removing nitrogen. The VIP plant in Norfolk, VA which also employs the VIP process began BNR operations approximately

seven (7) years prior to similar Nansemond operations (Balzer et al., 2005). Nitrogen removal was similar at both facilities during the first two years of side-by-side operation (1999 & 2000) (Balzer et al., 2005). Nansemond experienced an unexpected decline in nitrogen removal efficiency starting in 2001 which has continued to the present (Balzer et al., 2005). This deficiency in performance has been variable and has not been consistent from 2001 to the present time.

Profile sampling of wastewater treatment plants provides an effective method for evaluating the performance of a BNR processes by sampling for species such as ammonia, nitrate, nitrite, orthophosphate, soluble COD, etc. through the treatment train. Profiles are most commonly incorporated with DO and nutrient profiles (NH₃-N, NO₃-N, etc.), but can be applied to different applications; examining different bacterial activities in a treatment process. Profile sampling is often used in conjunction with modeling work and evaluating existing and future nutrient removal processes (Kochany et al, 2007; Scott et al, 2008; Kim et al, 2009).

The profile sampling data were combined with plant operating data and modeled using *Biowin 3.1* (EnviroSim, Ltd.). The profile sampling period (7-23-09 to 11-5-09) was divided into 5 separate periods to generate a calibrated model. The modeling for four of the periods was first performed in steady-state conditions. The period in which NTP stopped nitrifying had to be calibrated with a dynamic model. A calibrated simulation was then generated over the entire period of profile samples. These simulations were compared to data collected during the profile sampling to better understand the level of continuous nitrification inhibition.

The purpose of the baseline profile sampling and modeling was to address the following issues:

• Conduct baseline sampling prior to the more detailed nitrification inhibition study estimated to begin in March 2010, which will include influent sampling and the

operation of bench-scale sequencing batch reactors. This will be used to establish "normal" COD, nutrient and DO profiles though the VIP process without (and possibly with) the impact of inhibitory conditions, specifically with respect to N conversions and P release and uptake along the process.

- Evaluate the potential for nitrite accumulation in the process and its potential effect on aerobic phosphate uptake.
- Evaluate the impact of sporadic ferric chloride addition to the biological process as a means of preventing effluent TP exceedances.
- Evaluate the design μ_{max} to the actual observed μ_{max} for AOB through simulation modeling.
- Compare modeling and observed profile data for signs of continuous nitrification inhibition.

5.2 Methodology

Under normal circumstances, samples were collected during morning hours for 2 days of a normal business week from 7/23/09 to 11/5/09 via grab samples from the surface of each tank or reactor with a bucket. After the sample was collected it was allowed to settle for approximately 30 seconds until a supernatant layer was evident. Supernatant was then aspirated from the surface of the sample bucket using a 60 ml syringe. The sample would then be immediately filtered through the vacuum filtration system incorporating 0.45 µm membrane filters and transferred to the appropriate sample vial (see 5.2.1 Sample Preservation and Containers). DO and pH were measured and recorded when each grab sample was collected. This process was repeated for a total of two profiles per week. During each day of sampling the first sample collection point would be from the primary clarifier effluent. All successive

collection points were taken along the length of the BNR treatment process (refer to Figure 5.2) according to the actual hydraulic retention time presented in Table 5.1 (including the impact of recycle streams). The clarifier influent distribution channel sample was collected immediately following the last aeration basin sample and the RAS sample immediately after the secondary effluent sample was taken. A total of 15 samples were collected. Analysis of soluble COD (sCOD), Ammonia (NH₄-N), Nitrate (NO₃-N), Nitrite (NO₂-N), ortho-phosphate (PO₄-P) at all sampling locations.

- Primary Clarifier Effluent (PE) 1 Sample
- Anaerobic/Anoxic Tanks (AA1, AA2, AA3, AA4, AA5, AA6) 6 Samples, 1 sample
 per cell for a single train
- Aeration Tank (AE1, AE1.25, AE1.5, AE3, AE5) 5 samples, 1 at the beginning, 3 intermediate, 1 at the end for a single tank. The aeration tank was originally divided into 5 equal sample points along the length of the tank. AE1, AE2, AE3, AE4, AE5. Based on the results, two of the intermediate points were changed. Instead of taking 5 equally spaced points, two were taken between the distance of AE1 and AE2, which became AE1.25 and AE1.5 to better capture ammonia uptake, nitrite consumption, and nitrate production. AE4 was eliminated as values were similar to AE5 (Figure 5.1).
- Clarifier Recycle (RAS) 1 sample
- Secondary Clarifier Influent Distribution Channel (IDC) 1 sample
- Secondary Clarifier Effluent (SE) 1 sample

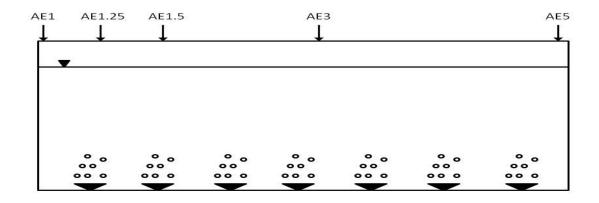


Figure 5.1 Aeration Tank Profile Sample Points

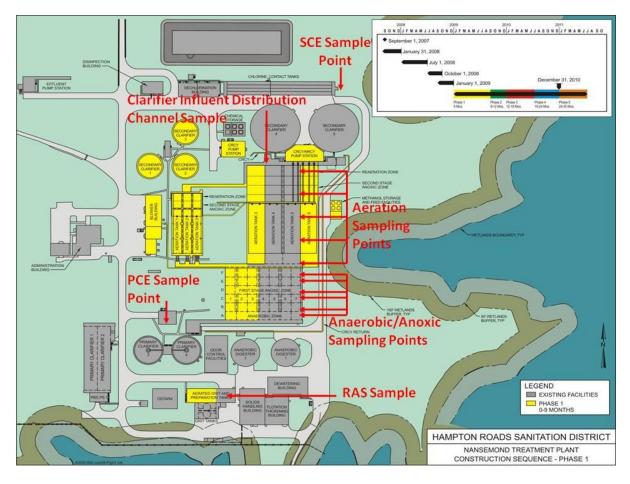


Figure 5.2 Profile Sampling Collection Points

As part of the process to stop nitrification, the nitrate recycle pump was also turned off converting all the anaerobic/anoxic cells to an anaerobic environment. Also due to upgrade efforts, the number of anaerobic/anoxic trains was reduced from 5 to 4 during the sampling period. This slightly changed the HRT in regards to sample collection; however, sample collection was still obtained based on the initial calculation of HRT from the start of the profile sampling. Profile sampling was conducted for a period of approximately 2 and half months until the plant stopped nitrifying.

5.2.1 Sample Preservation and Containers

After samples were collected and immediately filtered through 0.45µm membrane filters by vacuum pumps and associated filtration equipment they were placed into various containers for preservation before analysis. Table 5.2 provides a summary of the various containers, quantity of samples, and analysis methods which were used for the baseline profile sampling.

Table 5.1 Sample Preservation and Container Summary

Table 5.1 Sample 1 reservation and Container Summary					
Analysis Location:	NP	CEL	QA/QC by CEL		
Sample Container	15 ml centrifuge	40 ml VOA vial	(1) 40 ml VOA vial		
	tube	+ H ₂ SO ₄	(2) 40 ml VOA vial + H_2SO_4		
Analytes	PO ₄ -P	sCOD	(1) PO ₄ -P		
	NH ₄ -N		NO ₃ -N		
	NO ₃ -N		NO ₂ -N		
	NO ₂ -N		(2) NH ₄ -N		
Probe/Equipment	DO, pH				
No. Samples/week	15 x 2 = 30	15 x 2 = 30	2 x 2 = 4		
No. Samples/month	120	120	16		
Total No. Samples	480	480	64		
(duration of profiling)					

Note: The CEL sample vial used for sCOD analysis could also be used for the QA/QC sample for $NH_{\mathcal{F}}N$ analysis.

5.2.2 Analytical Methods

Sample analysis for profile sampling was conducted both on-site and off-site. Analytes measured at NTP for each profile include: PO₄-P, NH₄-N, NO₃-N, NO₂-N which were conducted using various HACH colorimetric test kits (see subsections 5.2.2.1 – 5.2.2.4) and a

HACH DR2800 spectrophotometer. Analysis of NO₂-N was performed the same day as sample collection, while analysis of PO₄-P, NH₄-N, and NO₃-N were conducted the following day. Samples were preserved by refrigeration overnight after NO₂-N analysis was complete. Off-site analysis by CEL consisted of soluble COD and quality assurance and quality control samples (QA/QC) of the same analytes measured on-site using *Standard Methods*.

5.2.2.1 Ammonia (NH₃-N) HACH Test Kit

Ammonia was analyzed using HACH Test N' Tube (TNT) 831 kit and HACH DR2800 spectrophotometer. This method uses the salicylate method, whereby ammonium ions react with hypochlorite and salicylate ions in the presence of sodium nitroprusside to as a catalyst form indophenol. The amount of color formed is directly proportional to the NH₄-N present.

5.2.2.2 Nitrate (NO₃-N) HACH Test Kit

Nitrate was analyzed using HACH TNT 835 kit. This kit incorporates the dimethylphenol method where nitrate ions in solution with sulfuric and phosphoric acids react with 2,6-dimethyphenol to form 4-nitro-2,6-dimethyphenol.

5.2.2.3 Nitrite (NO₂-N) HACH Test Kit

Nitrite was analyzed using NitriVer3 Nitrite Reagent Powder Pillows and 10 mL sample vials. This kit uses the diazotization method where nitrite in the sample reacts with sulfanilic acid to form a intermediate diazonium salt. This salt combined with chromotropic acid forms a pink color which is directly proportional to the amount of nitrite present.

5.2.2.4 Ortho-Phosphate (PO₄-P) HACH Test Kit

Ortho-Phosphate was analyzed using the HACH Reactive Phosphate TNT Reagent Kit.

This test kit uses the USEPA-approved PhosVer3 method where orthophosphate reacts with

molybdate in an acid to produce a mixed complex. Ascorbic acid then reduces this complex, producing an intense blue color.

5.2.3 Biowin Modeling Methodology

Modeling was performed using Biowin version 3.1, a biological wastewater treatment simulation package developed by EnviroSim Ltd (Flamborough, Ontario, Canada) and based on the IWA activated sludge models. Biowin was incorporated into this work to compare the data generated from the profile sampling with previous work performed by Hazen and Sawyer, P.C. (H&S). The Biowin model was calibrated using plant performance data from 7/23/09 to 11/5/09 period and previous simulation modeling work from H&S. Primary effluent wastewater fraction inputs for this work were derived from the NTP model created by H&S. The H&S NTP model was simulated with the raw influent characteristics that were generated from historical plant data and a two week special sampling study specified by H&S. The effluent data from the primary clarifier of this simulation was used to calculate wastewater fractions for the input of the simulation for the profile sampling work. Primary clarification was not simulated as part of this effort. Kinetic parameters such as maximum specific growth rate ($\mu_{max, AOB}$) and nitrite half saturation concentration were changed based on the work done by H&S in addition to changes made to fit the simulation model to the profile sampling data. A Garrett wasting configuration was incorporated to simplify modeling of secondary clarification (Figure 5.3), and the waste rate was then adjusted accordingly to match the solids wasted (lbs/day) from the plant performance data within a reasonable range. A Garrett wasting configuration directly wastes activated sludge from the aeration effluent, rather than wasting sludge from the secondary clarifier underflow (typical configuration). Diurnal load variations were not considered as part of this modeling.

Table 5.2 Biowin Influent Wastewater Fractions

Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.310
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.180
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.610
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.155
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.169
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.740
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.300
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.030
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.820
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.009
FZbh - Non-poly-P heterotrophs [gCOD/g of total COD]	0.000	0.0001
FZbm - Anoxic methanol utilizers [gCOD/g of total COD]	0.000	0.0001
FZaob - Ammonia oxidizers [gCOD/g of total COD]	0.000	0.0001
FZnob - Nitrite oxidizers [gCOD/g of total COD]	0.000	0.0001
FZamob - Anaerobic ammonia oxidizers [gCOD/g of total COD]	0.000	0.0001
FZbp - PAOs [gCOD/g of total COD]	0.000	0.0001
FZbpa - Propionic acetogens [gCOD/g of total COD]	0.000	0.0001
FZbam - Acetoclastic methanogens [gCOD/g of total COD]	0.000	0.0001
FZbhm - H2-utilizing methanogens [gCOD/g of total COD]	0.000	0.0001

5.2.3.1 Steady-State Simulation

Simulations were created for individual periods of the profile sampling as well as a simulation for the entire profile sampling period. Individual periods were created based on changes in mixed liquor suspended solids, changes in process; 5 anaerobic/anoxic trains to 4 anaerobic/anoxic trains; the Nitrate recycle (NRCY) being active and inactive, and the cease of nitrification. Steady-state simulations incorporated the use of constant input values. The entire profile sampling event was divided into 5 different periods and each period was modeled using steady-state conditions to calibrate the model except for the 4th period. Steady-state simulations were performed using the configuration shown in Figure 5.3.

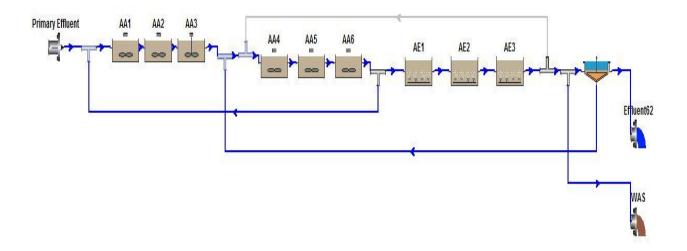


Figure 5.3 Biowin Simulation Model (Garrett Wasting Configuration)

5.2.3.2 Dynamic Simulation

Dynamic simulations were used to calibrate the model for the entire profile sampling event from start to finish. This required a reconfiguration of the model to account for dynamic simulation of one train being removed from service. This was done by creating a second anaerobic/anoxic train (train B) which only received 1/5 of the flow while the main train received 4/5 of the flow. During the period when all 5 trains were in operation the flow would be split to the 1/5 and 4/5 trains for a total of 5/5 (5 trains). Once the simulation reached the time period when one train was taken off-line, the simulation stopped the split of flow to train B (1/5) (Figure 5.4). This diverted all of the flow to the main anaerobic/anoxic train (4/5), simulating the system going from 5 to 4 anaerobic/anoxic trains. Dynamic simulations were first brought to steady-state using constant values from period 1 steady-state simulation and then simulated dynamically for the entire period. Itineraries for the various inputs were all based on plant performance data (WAS, NRCY, ARCY, RAS, Temperature, etc.).

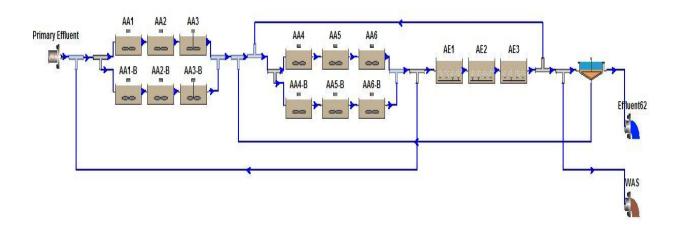


Figure 5.4 Biowin Model for Dynamic Simulation

5.3 Results and Discussion

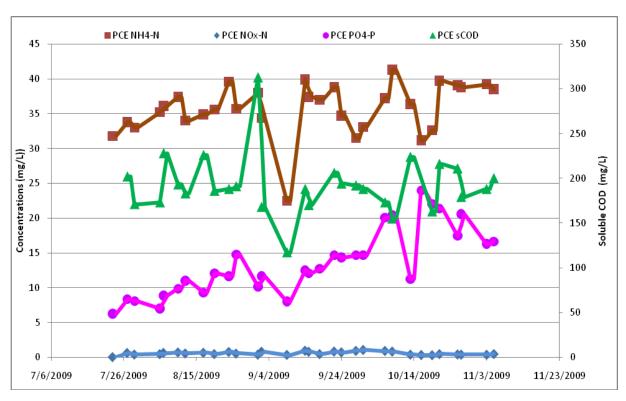
5.3.1 Profile Sampling

During the start of the profile sampling, Anaerobic/Anoxic (AA) trains 3-7, Aeration Tanks (AE) 4 and 5, and Secondary Clarifiers 4 and 5 were being operated at NP. Based on daily plant data as of July 23, 2009, the average daily flow rate over the entire profile sampling period was 18.11 MGD (final effluent). There was one nitrate recycle (NRCY) pump active at a flow of approximately 19.5 MGD, the anoxic recycle (ARCY) flow was 19.0 MGD, and the clarifier recycle (RAS) flow for secondary clarifiers 4 and 5 was 5.7 MGD each. Traditionally NRCY rates tend to be two or three times the influent flow. However, since the plant only had one recycle pump running at a rate of about the average daily flow this suggests that denitrification performance in the anoxic tanks is limited by nitrate availability. The plant added ferric chloride on an as needed basis to the aeration basin when the effluent TP approached the treatment objective. The hydraulic retention times (HRT) at the flow during the initial profile sampling for the different treatment processes are presented in Table 5.4.

Table 5.3 Hydraulic Retention Times at Current Average Day Flow Rate

	No. Units	Vol. Each Unit	Total Vol.	HRT w/out Recycle	HRT w/Recycle	Cumm. HRT w/RCY
	In Service	[MG]	[MG]	[hr]	[hr]	[hr]
Anaerobic Cell 1	5	0.174	0.87	1.21	0.58	0.58
Anaerobic Cell 2	5	0.174	0.87	1.21	0.58	1.15
Anaerobic Cell 3	5	0.174	0.87	1.21	0.58	1.73
Anoxic Cell 1	5	0.174	0.87	1.21	0.31	2.04
Anoxic Cell 2	5	0.174	0.87	1.21	0.31	2.35
Anoxic Cell 3	5	0.174	0.87	1.21	0.31	2.66
Aeration Tank	2	2.7	5.4	7.50	2.69	5.35
Secondary Clarifier	2	2.25	4.5	6.25	3.77	9.11

Profiling sampling started July 7, 2009 and ended on November 5, 2009. The data collected during this time period provided an overview of a BNR process coming out of nitrification as the SRT was intentionally lowered to stop nitrification. Figures 5.5 and 5.6 provide an overview of the influent (primary effluent) and secondary effluent characteristics during the entire profile sampling period. NH₄-N and NO_x-N data demonstrate that the plant began to stop nitrifying in early October, as the NH₄-N increased and the NO_x-N decreased. There was an observed steady increase in the influent phosphate which was attributed to centrate from the centrifuges and filtrate from the gravity belt thickeners.



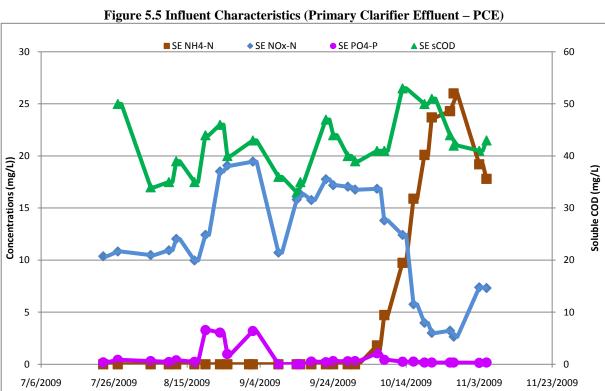


Figure 5.6 Secondary Effluent (SE) Characteristics

Profile sampling data were compared to the monthly plant performance data (composite samples) collected by HRSD. Results from the comparison proved to be very similar (Figure 5.7). This similarity supports the accuracy of the profile sampling data and the use of this data as a means for modeling nitrification performance to determine a maximum specific growth rate (μ_{max}) for AOB to compare to design μ_{max} used for the upgrade of the facility.

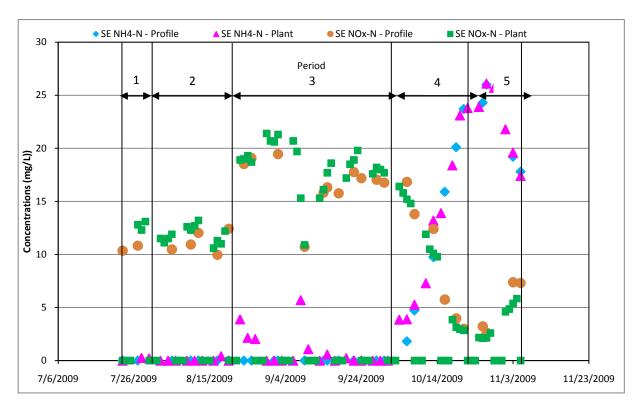


Figure 5.7 Comparison of Profile to Plant Performance Data

Temperature was very consistent from day to day sampling (Figure 5.8). It was observed that temperature slowly decreased as the weather changed. The coldest temperature that was observed was 22.5°C in November. Figure 5.9 demonstrates the relationship between the WAS rate and the SRT. It was observed that as the SRT was reduced throughout the profile sampling, the WAS rate correspondingly increased, until nitrification began to fail. At this point, the SRT was increased once again and the WAS decreased. The mixed liquor suspended solids (MLSS) data are consistent with the WAS and SRT data (Figure 5.8). Figure 5.10 was generated to show

the correlation between AOB washout SRT as a function of temperature assuming DO and pH were not limiting. The μ_{max} values used for calculating the AOB washout SRT were 0.90, 0.70, 0.60, and 0.50 day⁻¹ (see equation (1) below). Figures 5.8 showed that nitrification began to fail around a temperature of 22°C and based on Figure 5.9 the SRT during this failure was approximately 3 to 4 days. Based on the AOB washout SRT curve, at a temperature of 22°C, the μ_{max} which predicted an SRT in the range of 3 to 4 days was approximately 0.45 to 0.41 day⁻¹ based on equation (2) below. In calculating the above approximately μ_{max} for AOB an SRT of 3.5 days was used at a temperature 22°C. The calculated value suggested that the approximate range of μ_{max} for AOB to be between 0.40 – 0.60 days⁻¹.

$$SRT = \frac{1}{\left(\mathbf{A}_{\text{max}} \bullet \theta_{\text{max}} \right)^{T-20} - \left(\mathbf{A}_{i} \bullet \theta_{i}^{T-20} \right)}$$
 (1)

$$\mu_{\text{max}} = \frac{\frac{1}{SRT} + \mathbf{\Phi}_i \bullet \theta_i^{T-20}}{\theta_{\text{max}}^{T-20}}$$
 (2)

Where:

SRT = solids retention time

 $\mu_{max} = maximum \ specific \ growth \ rate$

 θ_{max} = Arrhenius temperature coefficient for growth

 $b_i = decay rate$

 θ_i = Arrhenius temperature coefficient for decay

T = temperature

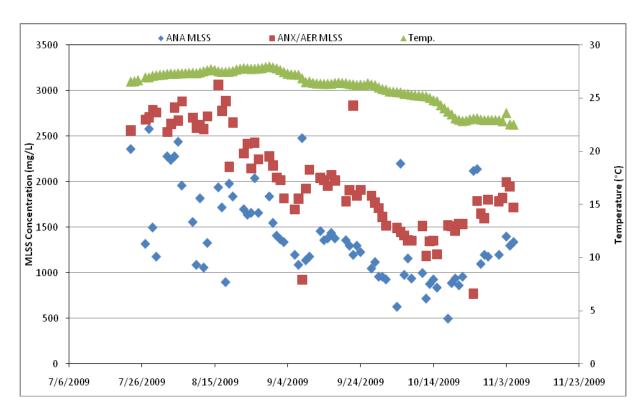


Figure 5.8 Mixed Liquor Suspended Solids (MLSS) and Temperature During Profile Sampling

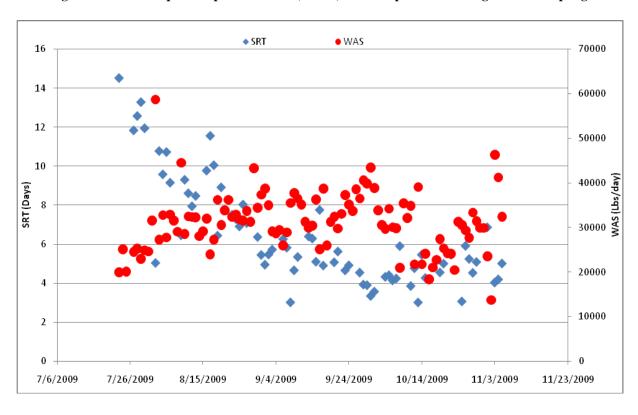


Figure 5.9 WAS and SRT During Profile Sampling

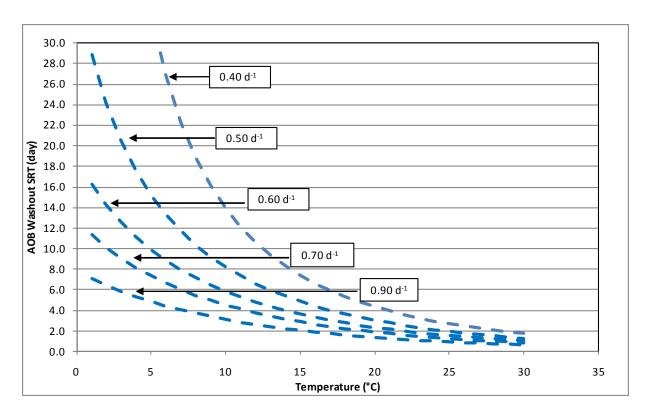


Figure 5.10 AOB Washout SRT as a Function of Temperature - μ_{max} = 0.40 to 0.90 days⁻¹.

Profile sampling data were also used to calculate ammonia uptake rates (AUR), specific ammonia uptake rates (SAUR), nitrate production rates (NPR), and specific nitrate production rates (SNPR). Slope regressions were performed of ammonia and nitrate concentrations over the hydraulic residence time (HRT) in the aeration tanks to determine uptake and production rates. In order to calculate the specific rates, the AUR and NPR were normalized to the mixed liquor volatile suspended solids (MLVSS). Nitrification performance was stable and rates were increasing until the SRT was purposely reduced to stop nitrification. This loss of nitrification was observed in the specific rates as time progressed from the start of the profile sampling to the end (Figure 5.11).

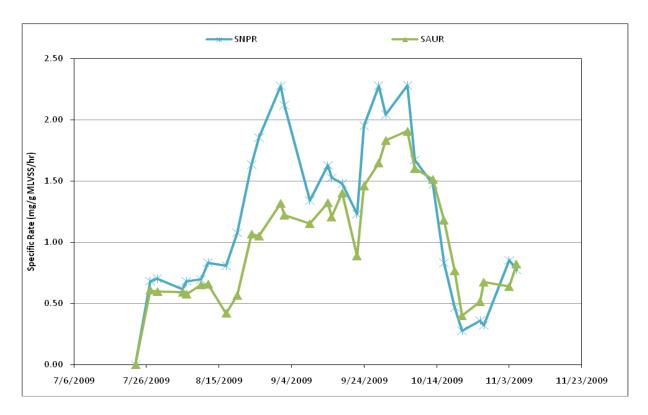


Figure 5.11 Specific Ammonia Uptake Rate (SAUR) and Specific Nitrate Production Rate (SNPR)

It was of interest to evaluate the potential for nitrite accumulation in the process and its potential effect on aerobic phosphate uptake by PAOs. Figure 5.12 shows aerobic nitrite concentration over the profile sampling period. The data shown represent individual profile experiments that were representative of the sampling period as a whole. It was evident that nitrite was present and accumulated to some degree as nitrification stopped. Although this was true, Figure 5.13 shows continued uptake of phosphate in the aerobic zone even after nitrification had stopped.

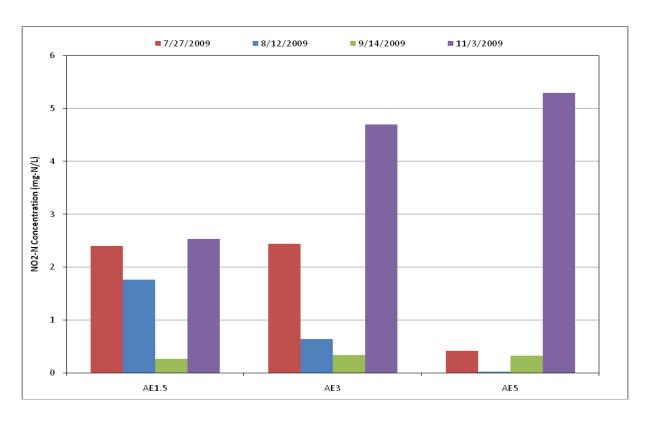


Figure 5.12 Aerobic NO₂-N Concentrations

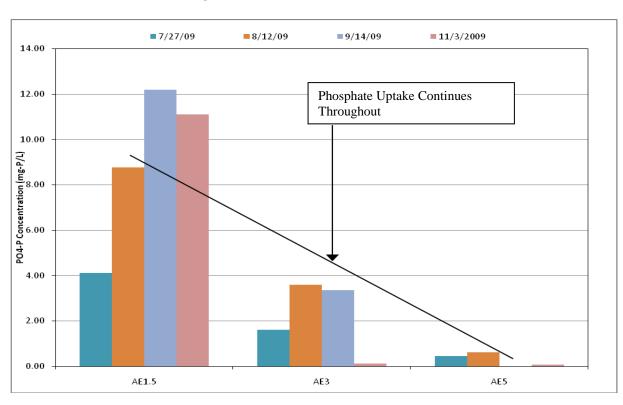


Figure 5.13 Aerobic PO₄-P Uptake

Data for late September and most of October were not shown on these figures due to the variability of this time period. Between September 30 and October 21 nitrification rates began to decrease and eventually ceased in late October/early November. This period of time was evaluated independently; however, aerobic phosphate uptake continued throughout the period of time when nitrification stopped (Figure 5.14). This suggested that there was no significant effect of nitrite accumulation on the aerobic phosphate uptake.

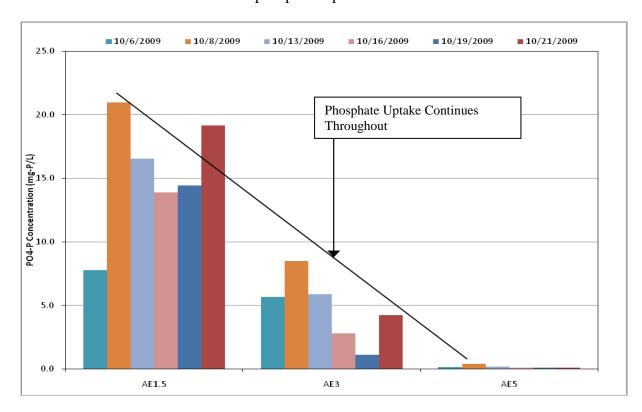


Figure 5.14 Aerobic PO4-P Uptake (10/6/09 - 10/21/09)

Ferric chloride was sporadically added during certain periods of the profile sampling to prevent effluent TP exceedances. Table 5.4 shows the dates during the profile sampling period in which ferric chloride was added to assist in TP removal. During these dates there was no observed change in nutrient profiles between days with and without the addition of ferric chloride. This suggested that there was no significant impact of ferric chloride on nitrogen removal.

Table 5.4 Wastewater Characteristics during Ferric Chloride Addition

		N	1LSS	SE Chem	. Addition	SC EFF	FIN EFF	FIN EFF	FIN EFF	FIN EFF
	WAS	ANA	ANX/AER	Chem	Dose	NH3	NOx	T-N	T-P	OPO4
Date	[lbs/day]	[mg/L]	[mg/L]	Туре	[mg/L]	[mg/L]	[mg/L]	[mg/L]	[mg/L]	[mg/L]
7/23/09	19957	2360	3940	Fe3	4.00	< 0.20				
7/24/09	25136			Fe3	4.84					
7/25/09	20080			Fe3	2.46					
7/26/09										
7/27/09	24512	1320	4060				12.8	15.4	0.55	0.258
7/28/09	25240	2580	4100			0.26	12.3	15.3	0.54	0.395
7/29/09	22932	1500	4220				13.1	15.6	0.49	0.334
7/30/09	24902	1180	4180			0.22				
7/31/09	24585									
8/22/09	36228									
8/23/09	32336	1700	3120			3.89	18.9	21.1	2.79	2.51
8/24/09	32935	1640	3180				19	21.1	1.71	1.54
8/25/09	31741	1660	2860	Fe3	0.02	2.15	19.3	21.4	0.96	0.74
8/26/09	31562	2040	3280	Fe3	0.02		18.7	21.9	0.85	0.578
8/27/09	33687	1660	3160	Fe3	0.02	2.04				
8/28/09	31233									
8/29/09	43327									
8/30/09	34395	1840	3160			< 0.20	21.4	23.5	2.18	1.9
8/31/09	37346	1550	2980				20.7	22.8	3.08	2.59
9/1/09	38706	1410	2800			< 0.20	20.6	22.4	2.93	2.73
9/2/09	34972	1370	2800				21.3	23.3	4.06	3.52
9/3/09	29091	1340	2420			< 0.20				
9/4/09	28724									
9/5/09	29398			Fe3	8.17					
9/6/09	25968	1200	2320	Fe3	8.62	< 0.20	20.7	22.1	0.69	0.514
9/7/09	28892	1090	2480	Fe3	8.17		19.7	22.3	0.67	0.399
9/8/09	35490	2480	1100	Fe3	6.16	5.7	15.3	21.0	0.63	0.277
9/9/09	37745	1140	2600	Fe3	5.08		10.9	13.2	0.44	0.092
9/10/09	36473	1180	2880	Fe3	2.07	1.08				
9/11/09	35126									
9/12/09	31251									
9/13/09	29903	1460	2760			< 0.20	15.3	16.9	0.48	0.21

5.3.2 Biowin Modeling

5.3.2.1 Steady-State Simulations

Model calibration was first conducted for periods 1, 2, 3, and 5 using steady-state conditions. Periods 1, 2, 3, and 5 are individual time periods during the profile sampling where MLSS, SRT, primary effluent TKN, and other wastewater characteristics were similar in value (refer to Figure 5.7). The results presented from the steady-state calibration were comparable to actual profile and plant performance data in regards to nutrient and specific rate profiles (Figures 5.15 – 5.19). The results suggested that modeling kinetics were similar in value to the actual observed.

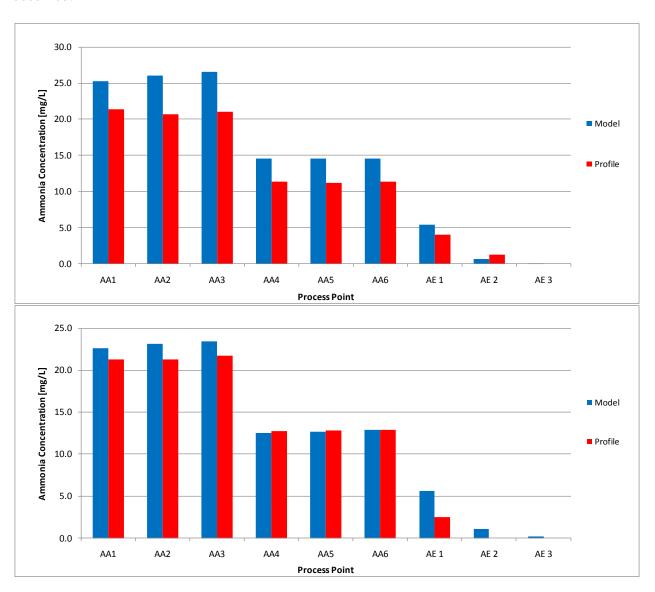


Figure 5.15 Ammonia Comparisons for Period 1-Top (7/23/09 – 7/31/09) & 2-Bottom (8/1/09 – 8/21/09)

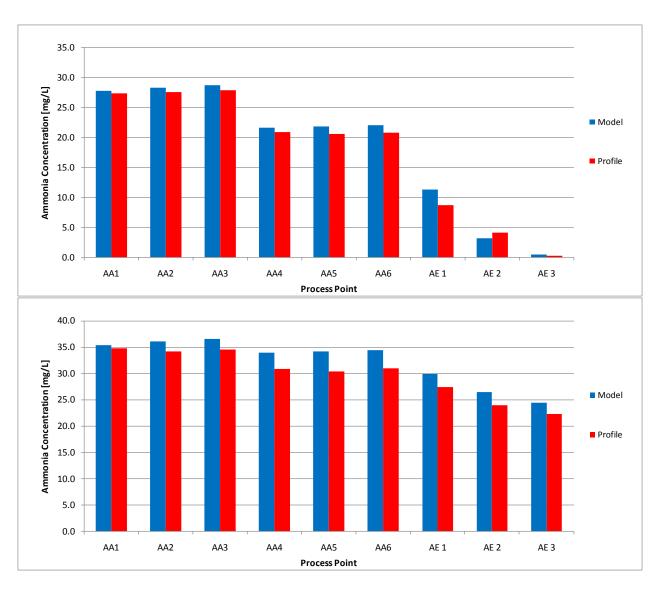


Figure 5.16 Ammonia Comparisons for Period 3-Top (8/22/09 – 10/2/09) & 5-Bottom (10/23/09 – 11/5/09)

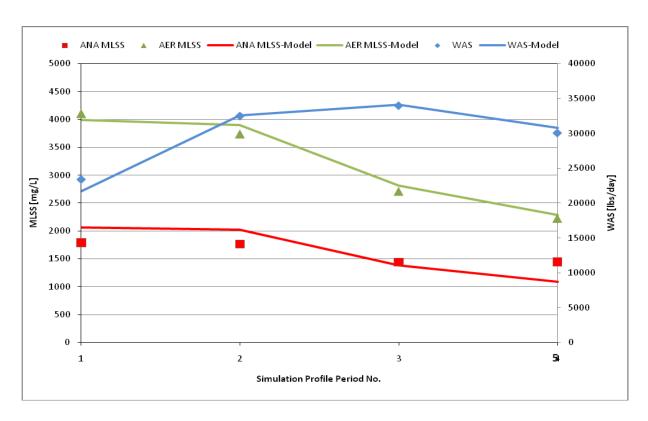


Figure 5.17 Mixed Liquor and WAS Comparison for Periods 1, 2, 3, and 5

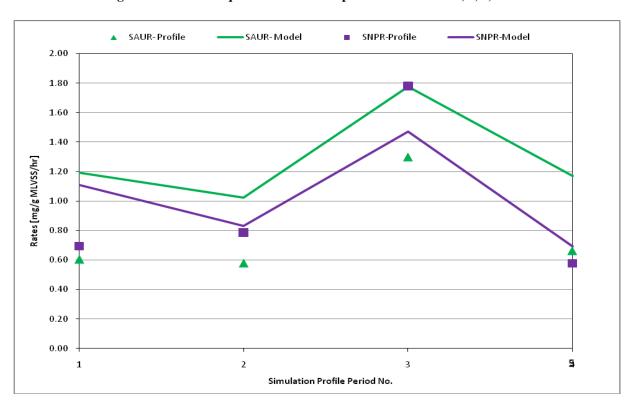


Figure 5.18 Nitrification Rate Comparisons for Periods 1, 2, 3, and 5

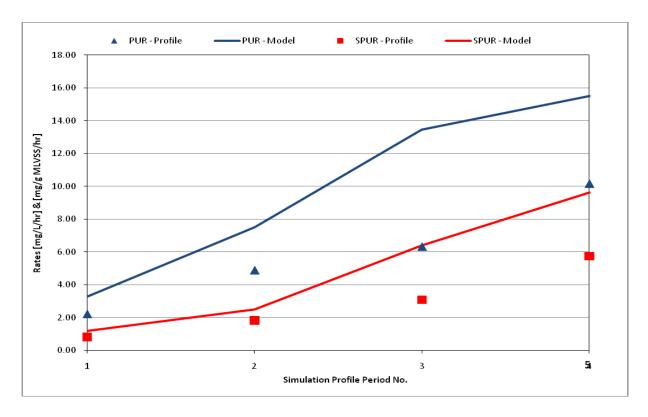


Figure 5.19 Phosphate Uptake Comparisons for Periods 1, 2, 3, and 5

Although similar, the specific rates calculated from the model were generally greater than what was actually observed. This suggested either that the model predicted better performance than the observed rates or the calibration was off, thus creating the differential.

5.3.2.2 Dynamic Simulations

Dynamic simulation generated similar results to profile and plant performance data; however, some of the data were not matched perfectly to the actual observed data. Period 4 (10/3/09 to 10/22/09) was modeled dynamically, as well, since it was the period where nitrification rates declined; however, the model could not be calibrated properly for this period alone. It was speculated that periods of high PE COD and TKN caused by removing primary clarifiers from service, which were not captured by the WAS and SRT, were the main cause of the offset between the model and the observed data. Due to the high COD and TKN during this period, the MLSS concentrations in the reactors increased; however, the plant data suggested a

decline in the WAS rather than an increase. The model continued to match the SRT, but the WAS and MLSS were not well correlated due the high PE COD and TKN (Figure 5.20). The figure demonstrated the offset between the model and observed plant data. The model, in the case of all the nutrients, failed to match the observed data and constantly over predicted the effluent concentrations. There were also unknown spikes observed starting at 10/14/09 believed to be caused by the high PE COD and TKN.



Figure 5.20 Nutrient Profile of Period 4 Dynamic Simulation

Therefore, instead of modeling period 4 separately, dynamic simulations of the entire profile sampling period were developed. There were noticeable differences between the model and observed data during the period from 10/3/09 to 10/22/09 for NH₄-N, NO₃-N, MLSS, SRT, and WAS (Figure 5.21, 5.22, 5.23, 5.24). From 7/23/09 to 10/2/09 MLSS slowly decreased as the plant reduced the SRT, thus increasing the WAS. The model followed this pattern until approximately 10/3/09. After this point, it was noted that the model SRT continued to match the

plant SRT; however, large differences were observed between the MLSS and WAS in comparison to plant data. Nutrient profiles behaved similarly as they initially matched plant data and then a large variation was seen. These unknown spikes predicted by the model in solids wasting, mixed liquor, and nutrient concentrations were not observed in the actual profile and plant data. The model also continued to predict failure of nitrification sooner than the actual data, which was followed by a steady recovery. It was suspected that the origin of the issue was input data taken from the plant, as there was a large increase in the inorganic suspended solids and total kheldajl nitrogen (TKN) casued by removing primary clarifiers from service.

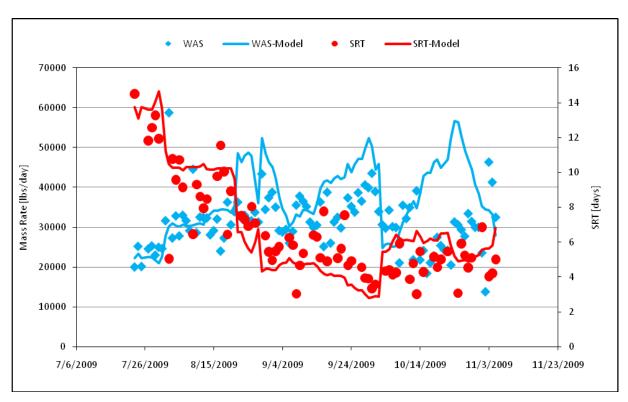


Figure 5.21 Comparison of WAS and SRT Plant Data to Model Data – $\mu_{max} = 0.62 days^{-1}$

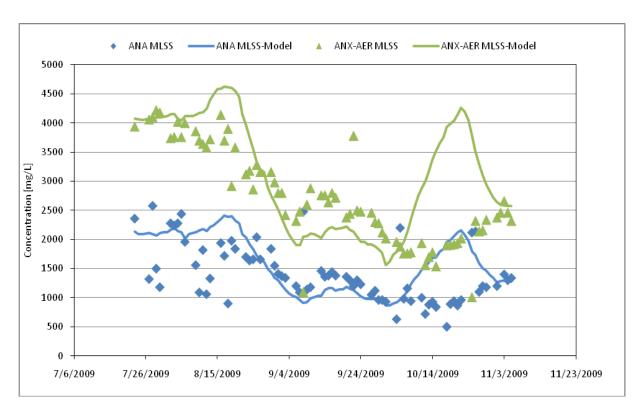


Figure 5.22 Mixed Liquor Suspended Solids Comparison – μ_{max} = 0.62days⁻¹

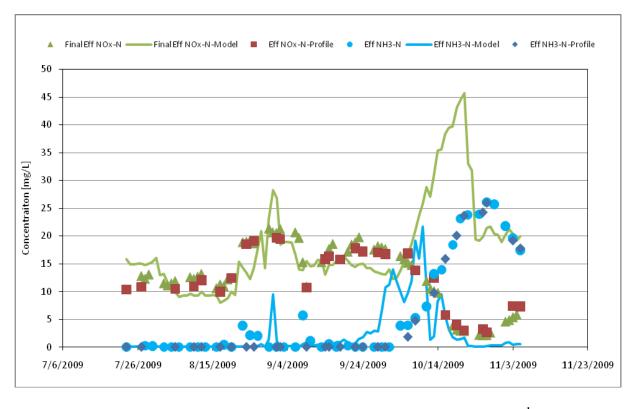


Figure 5.23 Final Effluent Nitrogen Species Comparison – $\mu_{max} = 0.62 days^{\text{-}1}$

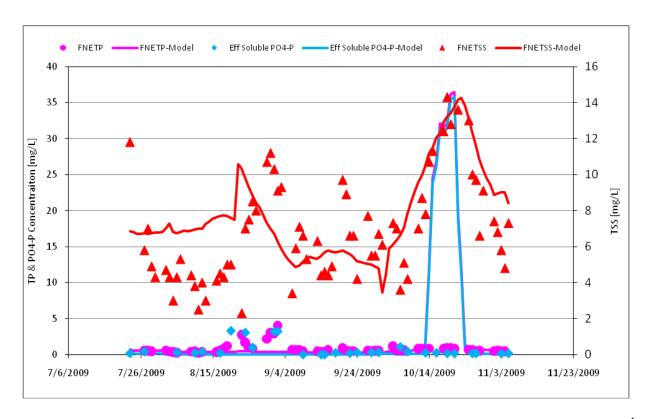


Figure 5.24 Final Effluent TP / TSS and Secondary Effluent ortho-Phosphate Comparison – μ_{max} = 0.62days⁻¹

Profiles of the secondary effluent ammonia concentration across the BNR process for certain dates demonstrated similar trends (Figure 5.25). The model and profile data were comparable until about 10/3/09 when a reduction in nitrification performance was observed. At this point the model predicted nitrification failure sooner than the observed data and then recovery.

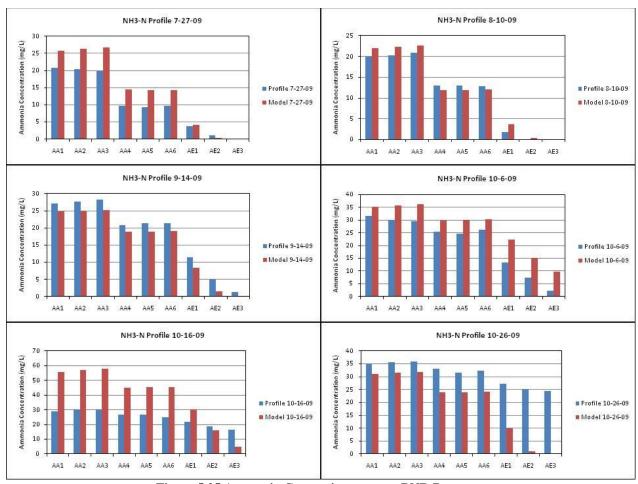


Figure 5.25 Ammonia Comparisons across BNR Process

Biowin modeling suggested that nitrification would stop sooner than what the actual profile data showed. The plant had intentionally stopped the NRCY and reduced the SRT to stop nitrification; however, the plant continued to nitrify for several more weeks.

Modeling provided a means of comparing the actual observed μ_{max} for AOB to the original design μ_{max} . The AOB μ_{max} value used for design was 0.57 days⁻¹. Modeling was carried out using various μ_{max} values to determine which value best fit the profile and plant data. Despite the variation between the model and the observed, the modeling suggested an approximate μ_{max} value close to a value in the range of 0.50 to 0.60 days⁻¹ (Figure 5.26 and 5.27). The suggested range of μ_{max} values was determined based on the effluent NH₄-N and NOx-N comparisons between generated data from Biowin and profile data. Although Biowin continued to show

failure of nitrification to occur sooner than the profile data, simulations generally followed the pattern that was seen in the profile data. Diurnal load variations were not considered as part of the modeling. This suggested that the actual in situ μ_{max} for AOB might be slightly greater than the projected μ_{max} . Regardless, it appeared that the suggested μ_{max} value and the original design value were similar.

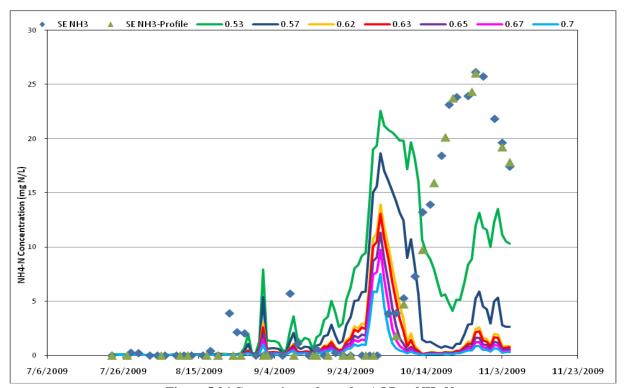


Figure 5.26 Comparison of μ_{max} for AOB to $NH_4\text{-}N$

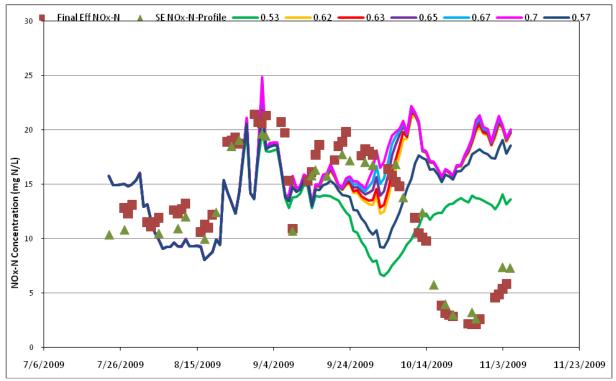


Figure 5.27 Comparison of μ_{max} for AOB to NOx-N

5.4 Conclusion

Based on results from the profile sampling and Biowin modeling, it was suggested that there was no significant sporadic inhibition event during the sampling period. Profile and plant data was collected which demonstrated a fully nitrifying plant ceasing to nitrify as the SRT was reduced to values below which one would not expect nitrification to be normally maintained (Figure 5.7). This allowed for the use of simulation modeling to estimate a μ_{max} for AOB which correlates with the observed data. This estimated value could then be compared to the original design value. Modeling suggested a μ_{max} for AOB to be approximately in the range of 0.50 – 0.60 days⁻¹, based primarily on the effluent ammonia concentrations, which was similar to the original design value of 0.57 days⁻¹. This supported previous batch-rate testing which suggested that there was some form of continuous inhibition present in the wastewater at NTP. Though model and profile/plant data were somewhat similar, the model did not match the profile data within a reasonable difference towards the end of the sampling. Reasoning behind the difference

was high PE COD and TKN concentrations recorded in the plant performance data. These high concentrations were caused by bad samples collected during this period where primary clarifiers were being taken in and out of service.

5.5 References

American Public Health Association, American Water Works Association, Water Environment Federation. (1998). *Standard Methods for the Examination of Water and Wastewater*; 20th ed.: Washington D.C.

Balzer, B., Cox, M., Deluna, J., Ghosn, S., Hogg, P., Kennedy, G., Pletl, J. 2005. The Status of Biological Nutrient Removal at HRSD's Nansemond Treatment Plant. NITRO Team Report.

Hazen and Sawyer. 2007. Nansemond Treatment Plant Nutrient Reduction Improvement Technical Memorandum.

Kim, H.S., Pei, R., Gunsch, C., Gellner, J.W., Boltaz, J.P., Freudenberg, B., Dodson, R., Cho, K.D., Schuler, A.J. 2009. Trace Organic Chemical Profiles in Nutrient Removal Systems With and Without Integrated Fixed Film Activated Sludge. *WEFTEC 09*, 704-711.

Kochany, J., Lipczynska-Kochany, E., Smith, W. 2007. Modeling Strategy for SBR Recovery After Upset. *WEFTEC 07*. Session 44, 3242-3251.

Scott, Z., Olson, B.H., Esmond, S., Maleki, N., Scherfig, J. 2008. Monitoring and Modeling the Anaerobic Digestion of Manure with qPCR in a Mesophilic Anaerobic Digester. *WEFTEC 08*, 534-553.

Yi, P.H., Balzer, B., Bilyk, K., Bott, C.B. (In Preperation - 2010). Evaluation of Nitrification Inhibition Using Bench-Scale Rate Measurements. *Water Environment Research*.

6. Engineering Significance

The purpose of this project was to identify a specific source of inhibition which would cause failure of nitrification and determine the inhibitory characteristics of NTP influent wastewater. The first objective was to attempt to find the cause of sporadic nitrification upset events at NTP, and at the outset of this work, it was hypothesized that possible sources included the hog processing facility, landfill leachate, and truck delivered septage/FOG/chemical toilet waste could be the cause. AOB and NOB batch rate testing using a range of biomass and wastewater sources suggested that there was some level of continuous inhibition, however no significant source of sporadic inhibition was observed.

The profile sampling and modeling, combined with the calibrated model developed from historical plant data by H&S seemed to indicate this as well. Based on this work, there was definitely some degree of continuous nitrification inhibition characterized by an apparent AOB μ_{max} reduction from a typical value of 0.9 days⁻¹ to a value approximately in the range of 0.50 – 0.60 days⁻¹. An evaluation of the nitrification process at the treatment plant was performed as the SRT was reduced to stop nitrification (13 days to 3-4 days). Evaluation of the profile sampling data showed that the plant ceased nitrification at an approximate SRT of 3 to 4 days and a temperature around 22°C. It was suggested that for these parameters, the AOB μ_{max} at which washout occurred would be close to a value of 0.45 to 0.41 day⁻¹. This was based on an AOB washout curve developed as a function of the temperature (not including rate reduction due to pH and DO).

Although the suspected inhibitors such as the hog processing facility and landfill leachate were eliminated as causes of inhibition based on this work, these sources should be reconsidered in the future. Sporadic nitrification inhibition was not observed using samples collected on one

of several days. For this reason, it was proposed in future work to configure continuous batch reactors to examine if the nitrification failure is caused over a period of time or capture the sporadic failure.

The next step was to determine the extent of the continuous nitrification inhibition. This was carried out through profile sampling of the BNR process. This provided a means for looking at nutrient concentrations through the treatment process to determine if any part of the treatment process was operating inefficiently or where issues may arise. The plant sporadically added ferric chloride to assist in meeting effluent TP limits; however, there were no observed affects on nitrification. No trends were observed in the influent received at the plant or plant process operations based on the profile sampling which would indicate an inhibition issue. The loss of nitrification was captured as the plant purposely reduced the SRT to stop nitrification.

Biowin modeling was used to calculate the μ_{max} value based on the correlation of effluent nutrient concentrations between the model and profile data. The model failed to perfectly match the observed data towards the end of the profile sampling, where there were noticeable differences. It was believed that the cause was due to primary clarifiers being taken out of service. This caused the collected samples to contain higher concentrations of COD and TKN which created a false representative sample. In spite of this, the model was still used to calculate a range of μ_{max} values which best correlated with the profile data. After several iterations with different μ_{max} values, it was determined that the range of values which best fit the observed effluent ammonia and NO_x -N concentrations were between 0.50-0.60 day⁻¹.

Appendices

7.1 Appendix A

PERIOD 1 (7/23/09 - 7/31/09)

SIMULATION INPUTS

Primary Effluent Fractions		
Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.270
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.220
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.615
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.157
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.171
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.744
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.148
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.033
FupN - N: COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.849
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.011

Primary Ef	Primary Effluent Itinerary				ANT					
Flow	Total COD mgCOD/L	TKN mgN/L	TP mgP/L	NO3- N mgN/L	pН	Alk. mmol/L	ISS mgISS/L	Ca mg/L	Mg mg/L	DO mg/L
18.36	422.10	46.2	9.56	0.1	7	5.4	21.84	12	6	0
	↑						↑			
Increased	0%					Increased	0%			
by =						by =				
Original =	422.1					Original =	21.84			

*NOTE: Values are averages of effluent data taken from Nansemond MPOR

	Total Vol.	HRT w/out recycle	DO-Set Point	Tanks in Service	% Removal	Sludge Blanket	Underflow - Constant
Process	[MG]	[hr]	[mg/L]	[No.]	[%]		[MGD]
Anaerobic Tanks (3 cells per Train)	2.606193	1.17	un- aerated	5	-	-	-
Anoxic Tanks (3 cells per Train)	2.606193	1.17	un- aerated	5	-	-	-
Aeration Tank 1	1.800	1.453	2	2	-	-	-
Aeration Tank 2	1.800	1.453	2	2	-	-	-
Aeration Tank 3	1.800	1.453	2	2	-	1	-
Secondary Clarifier	4.510	3.640	-	2	99.9	0.2	0.00

	Rate in Side	Desired Avg. Solids		Avg. Over	
	(S)	Waste	Flow Data	Period	SRT
SPLITTERS	[MGD]	[lbs/d]	ARCY =	18.90	[days]
WAS	0.65	23418	NRCY ON =	19.5	12.82
NRCY	19.5		CRCY =	11.37	
ARCY	18.90		Daily =	18.35	
			Temperature =	26.89	
PARAMETER	S (KINETIC)				
AOB					
Name			Default	Value	Arrhenius
Max. spec. grov	vth rate [1/d]		0.9	0.53	1.072
Substrate (NH4)) half sat. [mgN/L]		0.7	0.7	1
Aerobic decay r	ate [1/d]		0.17	0.17	1.029
Anoxic/anaerob	ic decay rate [1/d]		0.08	0.08	1.029
KiHNO2 [mmo	1/L]		0.005	0.005	1
NOB					
Name			Default	Value	Arrhenius
Max. spec. grov	vth rate [1/d]		0.7	0.7	1.06
) half sat. [mgN/L]		0.1	0.1	1
Aerobic decay r			0.17	0.17	1.029
•	ic decay rate [1/d]		0.08	0.08	1.029
KiNH3 [mmol/I	•		0.075	0.075	1
SWITCHES					
Name				Def	ault Value
Heterotrophic D	OO half sat. [mgO2	/L]		0.	0.05
Aerobic denit. I	OO half sat. [mgO2	/L]		0.	0.05
Ammonia oxidi	zer DO half sat. [m	ngO2/L]		0.	25 0.25
Nitrite oxidizer	DO half sat. [mgO	2/L]		0	.5 0.5
Anaerobic amm	onia oxidizer DO l	nalf sat. [mgO2/L]		0.	0.01
Anoxic NO3 ha	lf sat. [mgN/L]			0	.1 0.1
Anoxic NO2 ha	lf sat. (mgN/L)			0.	01 0.1
NH3 nutrient ha	ılf sat. [mgN/L]			1.00	E-04 0.001
PolyP half sat. [mgP/L]			0.	0.01
VFA sequestrati	ion half sat. [mgC0	DD/L]		4	5 5
P uptake half sa	t. [mgP/L]			0.	15 0.15
P nutrient half s	at. [mgP/L]			0.0	0.001
Autotroph CO2	half sat. [mmol/L]			0	.1 0.1
Heterotrophic H	lydrogen half sat. [mgCOD/L]			1 1
Propionic aceto	gens Hydrogen hal	f sat. [mgCOD/L]		4	5 5

SUMMARY OF INPUTS

										SC
					FIN		FIN		FIN	EFF
					EFF		EFF		EFF	NH4
Total		ANA	ANX	AER	TKN	FIN EFF	T-N	FIN EFF	TSS	-N
SRT	WAS	MLSS	MLSS	MLSS	[mg/	NOx	[mg/	T-P	[mg/	[mg/
[days]	[lbs/day]	[mg/L]	[mg/L]	[mg/L]	L]	[mg/L]	L]	[mg/L]	L]	L]
12.82	23418	1788	4100	4100	2.72	12.73	15.45	0.53	6.76	0.24

Profile Data

SC EFF OPO4-P [mg/L]	SC EFF NOx-N [mg/L]	SC EFF NH4-N [mg/L]	AUR [mg/L/ hr]	SAUR [mg/g MLVSS /hr]	NPR [mg/ L/hr]	SNPR [mg/g MLVSS/hr]	PUR [mg/ L/hr]	SPUR [mg/g MLVSS/hr]
0.30	10.59	<1.0	1.65	0.60	1.90	0.69	2.241	0.820

PERIOD 2 (8/1/09 - 8/21/09)

SIMULATION INPUTS

Primary Effluent Fractions		
Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.270
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.220
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.615
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.157
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.171
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.744
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.148
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.033
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.849
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.011

Primary Ef	Primary Effluent Itinerary				CONSTANT					
Flow	Total COD mgCOD/L	TKN mgN/L	TP mgP/L	NO3- N mgN/L	рН	Alk. mmol/L	ISS mgISS/L	Ca mg/L	Mg mg/L	DO mg/L
19.2	630.53	40.65	8.83	0.1	7	5.4	25.4	12	6	0
Increased by =	↑ 23%					Increased by =	↑ 23%			

			ansemond N	

	Total Vol.	HRT w/out recycle	DO-Set Point	Tanks in Service	% Removal	Sludge Blanket	Underflow - Constant
Process	[MG]	[hr]	[mg/L]	[No.]	[%]		[MGD]
Anaerobic Tanks (3 cells per Train)	2.606193	1.17	un- aerated	5	-	-	-
Anoxic Tanks (3 cells per Train)	2.606193	1.17	un- aerated	5	-	-	-
Aeration Tank 1	1.800	1.453	2	2	-	-	-
Aeration Tank 2	1.800	1.453	2	2	-	-	-
Aeration Tank 3	1.800	1.453	2	2	-	-	-
Secondary Clarifier	4.510	3.640	-	2	99.9	0.2	0.00

SPLITTERS	Rate in Side (S) [MGD]	Desired Avg. Solids Waste [lbs/d]	Flow Data ARCY =	Avg. Over Period	Avg. Calc. SRT [days]
	[MOD]		†		
WAS	1	23418	NRCY ON =	19.5	8.85
NRCY	19.5		CRCY =	11.80	
ARCY	19.01		Daily =	19.18	
			Temperature =	27.44	

PARAMETERS (KINETIC)

AOB

Name	Default	Value	Arrhenius
Max. spec. growth rate [1/d]	0.9	0.53	1.072
Substrate (NH4) half sat. [mgN/L]	0.7	0.7	1
Aerobic decay rate [1/d]	0.17	0.17	1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08	1.029
KiHNO2 [mmol/L]	0.005	0.005	1

NOB

Name	Default	Value	Arrhenius
Max. spec. growth rate [1/d]	0.7	0.7	1.06
Substrate (NO2) half sat. [mgN/L]	0.1	0.1	1
Aerobic decay rate [1/d]	0.17	0.17	1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08	1.029
KiNH3 [mmol/L]	0.075	0.075	1

SWITCHES

Name	Default	Value
Heterotrophic DO half sat. [mgO2/L]	0.05	0.05
Aerobic denit. DO half sat. [mgO2/L]	0.05	0.05

Ammonia oxidizer DO half sat. [mgO2/L]	0.25	0.25
Nitrite oxidizer DO half sat. [mgO2/L]	0.5	0.5
Anaerobic ammonia oxidizer DO half sat. [mgO2/L]	0.01	0.01
Anoxic NO3 half sat. [mgN/L]	0.1	0.1
Anoxic NO2 half sat. (mgN/L)	0.01	0.1
NH3 nutrient half sat. [mgN/L]	1.00E-04	0.001
PolyP half sat. [mgP/L]	0.01	0.01
VFA sequestration half sat. [mgCOD/L]	5	5
P uptake half sat. [mgP/L]	0.15	0.15
P nutrient half sat. [mgP/L]	0.001	0.001
Autotroph CO2 half sat. [mmol/L]	0.1	0.1
Heterotrophic Hydrogen half sat. [mgCOD/L]	1	1
Propionic acetogens Hydrogen half sat. [mgCOD/L]	5	5
Synthesis anion/cation half sat. [meq/L]	0.01	0.01

SUMMARY OF INPUTS

										SC
					FIN		FIN		FIN	EFF
					EFF		EFF		EFF	NH4
Total		ANA	ANX	AER	TKN	FIN EFF	T-N	FIN EFF	TSS	-N
SRT	WAS	MLSS	MLSS	MLSS	[mg/	NOx	[mg/	T-P	[mg/	[mg/
[days]	[lbs/day]	[mg/L]	[mg/L]	[mg/L]	L]	[mg/L]	L]	[mg/L]	L]	L]
8.85	32509	1763	3735	3735	1.81	11.83	13.63	.51	4.1	.42

Profile Data

SC EFF OPO4-P [mg/L]	SC EFF NOx-N [mg/L]	SC EFF NH4-N [mg/L]	AUR [mg/L/ hr]	SAUR [mg/g MLVSS /hr]	NPR [mg/ L/hr]	SNPR [mg/g MLVSS/hr]	PUR [mg/ L/hr]	SPUR [mg/g MLVSS/hr]
0.88	11.17	<1.0	1.55	0.58	2.12	0.79	4.91	1.82

PERIOD 3 (8/22/09 - 10/2/09)

SIMULATION INPUTS

Primary Effluent Fractions		
Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.270
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.220
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.615
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.157
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.171
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.744
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.148

Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.033
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.849
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.011

Primary Ef	Primary Effluent Itinerary				CONSTANT					
Flow	Total COD mgCOD/L	TKN mgN/L	TP mgP/L	NO3- N mgN/L	pН	Alk. mmol/L	ISS mgISS/L	Ca mg/L	Mg mg/L	DO mg/L
19.96	565.1	41.7	11.3	0.1	7	5.4	23.71	12	6	0
	↑						↑			
Increased	20%					Increased	20%			
by =						by =				
Original =	470.9					Original =	19.76			

*NOTE: Values are averages of effluent data taken from Nansemond MPOR

Process	Total Vol. [MG]	HRT w/out recycle [hr]	DO-Set Point [mg/L]	Tanks in Service [No.]	% Removal [%]	Sludge Blanket	Underflow - Constant [MGD]
Anaerobic Tanks (3 cells per Train)	2.085	0.91	un- aerated	4	-	-	-
Anoxic Tanks (3 cells per Train)	2.085	0.91	un- aerated	4	-	-	-
Aeration Tank 1	1.800	1.453	2	2	-	-	-
Aeration Tank 2	1.800	1.453	2	2	-	-	-
Aeration Tank 3	1.800	1.453	2	2	-	-	-
Secondary Clarifier	4.510	3.640	-	2	99.85	0.2	0.00

SPLITTERS	Rate in Side (S) [MGD]	Desired Avg. Solids Waste [lbs/d]	Flow Data ARCY =	Avg. Over Period	Avg. Calc. SRT [days]
SELITIENS	[MGD]	[IDS/U]	AKCI –	17.00	[uays]
WAS	1.45	33986	NRCY ON =	0	5.59
NRCY	0		CRCY =	11.92	
ARCY	17.86		Daily =	19.99	
			Temperature =	26.84	

PARAMETERS (KINETIC)

AOB			
Name	Default	Value	Arrhenius
Max. spec. growth rate [1/d]	0.9	0.53	1.072
Substrate (NH4) half sat. [mgN/L]	0.7	0.7	1
Aerobic decay rate [1/d]	0.17	0.17	1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08	1.029
KiHNO2 [mmol/L]	0.005	0.005	1

NOB				
Name	Default	Value	A	rrhenius
Max. spec. growth rate [1/d]	0.7	0.7		1.06
Substrate (NO2) half sat. [mgN/L]	0.1	0.1		1
Aerobic decay rate [1/d]	0.17	0.17		1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08		1.029
KiNH3 [mmol/L]	0.075	0.075		1
SWITCHES				
Name			Default	Value
Heterotrophic DO half sat. [mgO2/L]			0.05	0.05
Aerobic denit. DO half sat. [mgO2/L]			0.05	0.05
Ammonia oxidizer DO half sat. [mgO2/L]			0.25	0.25
Nitrite oxidizer DO half sat. [mgO2/L]			0.5	0.5
Anaerobic ammonia oxidizer DO half sat. [mgO2/L]			0.01	0.01
Anoxic NO3 half sat. [mgN/L]			0.1	0.1
Anoxic NO2 half sat. (mgN/L)			0.01	0.1
NH3 nutrient half sat. [mgN/L]			1.00E-04	0.001
PolyP half sat. [mgP/L]			0.01	0.01
VFA sequestration half sat. [mgCOD/L]			5	5
P uptake half sat. [mgP/L]			0.15	0.15
P nutrient half sat. [mgP/L]			0.001	0.001
Autotroph CO2 half sat. [mmol/L]			0.1	0.1
Heterotrophic Hydrogen half sat. [mgCOD/L]			1	1
Propionic acetogens Hydrogen half sat. [mgCOD/L]			5	5
Synthesis anion/cation half sat. [meq/L]			0.01	0.01

SUMMARY OF INPUTS

										SC
					FIN		FIN		FIN	EFF
					EFF		EFF		EFF	NH4
Total		ANA	ANX	AER	TKN	FIN EFF	T-N	FIN EFF	TSS	-N
SRT	WAS	MLSS	MLSS	MLSS	[mg/	NOx	[mg/	T-P	[mg/	[mg/
[days]	[lbs/day]	[mg/L]	[mg/L]	[mg/L]	L]	[mg/L]	L]	[mg/L]	L]	$\mathbf{L}]$
5.59	33986	1440	2710	2710	2.16	18.38	20.54	1.20	6.89	2.24

Profile Data

	SC	SC		SAUR		SNPR		SPUR
SC EFF OPO4-P	EFF NOx-N	EFF NH4-N	AUR [mg/L/	[mg/g MLVSS	NPR [mg/	[mg/g MLVSS/hr	PUR [mg/	[mg/g MLVSS/hr
[mg/L]	[mg/L]	[mg/L]	hr]	/hr]	L/hr]		L/hr]	
1.31	17	<1.0	2.64	1.3	3.65	1.78	6.35	3.09

PERIOD 5 (10/23/09 - 11/5/09)

SIMULATION INPUTS

Primary Effluent Fractions		
Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.270
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.220
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.615
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.157
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.171
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.744
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.148
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.033
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.849
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.011

Primary Ef	Primary Effluent Itinerary					CONSTANT						
Flow	Total COD mgCOD/L	TKN mgN/L	TP mgP/L	NO3- N mgN/L	pН	Alk. mmol/L	ISS mgISS/L	Ca mg/L	Mg mg/L	DO mg/L		
19.27	481.3	46.1	13.6	0.1	7	5.4	22	12	6	0		
	↑						↑					
Increased	0%					Increased	0%					
by =						by =						
Original =	481.3					Original =	21.97					

*NOTE: Values are averages of effluent data taken from Nansemond MPOR

Process	Total Vol. [MG]	HRT w/out recycle [hr]	DO-Set Point [mg/L]	Tanks in Service [No.]	% Removal [%]	Sludge Blanket	Underflow - Constant [MGD]
Anaerobic Tanks (3 cells per Train)	2.085	0.91	un- aerated	4	-	-	-
Anoxic Tanks (3 cells per Train)	2.085	0.91	un- aerated	4	-	-	-
Aeration Tank 1	1.800	1.453	2	2	-	-	-
Aeration Tank 2	1.800	1.453	2	2	-	-	-
Aeration Tank 3	1.800	1.453	2	2	-	-	-
Secondary Clarifier	4.510	3.640	-	2	99.8	0.2	0.00

Rate in Side	Desired Avg. Solids		Avg. Over	Avg. Calc.
(S)	Waste	Flow Data	Period	SRT

SPLITTERS	[MGD]	[lbs/d]	ARCY =	16.99	[days]
WAS	1.6	30075	NRCY ON =	0	4.90
NRCY	0		CRCY =	11.40	
ARCY	16.99		Daily =	19.13	
			Temperature =	22.49	
PARAMETERS	S (KINETIC)				·
AOB					
Name			Default	Value	Arrhenius
Max. spec. grow	th rate [1/d]		0.9	0.53	1.072
Substrate (NH4)	half sat. [mgN/L]		0.7	0.7	1
Aerobic decay ra	ate [1/d]		0.17	0.17	1.029
Anoxic/anaerobi	c decay rate [1/d]		0.08	0.08	1.029
KiHNO2 [mmol	/L]		0.005	0.005	1
NOB					
Name			Default	Value	Arrhenius
Max. spec. grow	th rate [1/d]		0.7	0.7	1.06
Substrate (NO2)	half sat. [mgN/L]		0.1	0.1	1
Aerobic decay ra	ate [1/d]		0.17	0.17	1.029
Anoxic/anaerobi	c decay rate [1/d]		0.08	0.08	1.029
KiNH3 [mmol/L	.]		0.075	0.075	1
SWITCHES					
Name				De	efault Value
Heterotrophic D	O half sat. [mgO2/L]		(0.05
Aerobic denit. D	O half sat. [mgO2/L	.]		(0.05
Ammonia oxidiz	er DO half sat. [mg	O2/L]		(0.25
Nitrite oxidizer I	DO half sat. [mgO2/	L]			0.5 0.5
Anaerobic ammo	onia oxidizer DO ha	If sat. [mgO2/L]		(0.01
Anoxic NO3 hal	f sat. [mgN/L]				0.1
Anoxic NO2 hal	f sat. (mgN/L)			(0.01 0.1
NH3 nutrient hal	If sat. [mgN/L]			1.0	0E-04 0.001
PolyP half sat. [1	mgP/L]			(0.01
VFA sequestration	on half sat. [mgCOI	D/L]			5 5
P uptake half sat	. [mgP/L]			(0.15
P nutrient half sa	at. [mgP/L]			0	.001 0.001
Autotroph CO2	half sat. [mmol/L]				0.1 0.1
Heterotrophic H	ydrogen half sat. [m	gCOD/L]			1 1
Propionic acetog	gens Hydrogen half s	sat. [mgCOD/L]			5 5
Synthesis anion/	cation half sat. [meq	/L]		(0.01

SUMMARY OF INPUTS

D C I I I I										
										SC
					FIN		FIN		FIN	EFF
					EFF		EFF		EFF	NH4
Total		ANA	ANX	AER	TKN	FIN EFF	T-N	FIN EFF	TSS	-N
SRT	WAS	MLSS	MLSS	MLSS	[mg/	NOx	[mg/	T-P	[mg/	[mg/
[days]	[lbs/day]	[mg/L]	[mg/L]	[mg/L]	L]	[mg/L]	L]	[mg/L]	L]	L]
4.89	30075	1442	2225	2225	24.35	3.71	28.06	0.56	8.05	22.4

Profile Data

SC EFF OPO4-P [mg/L]	SC EFF NOx-N [mg/L]	SC EFF NH4-N [mg/L]	AUR [mg/L/ hr]	SAUR [mg/g MLVSS /hr]	NPR [mg/ L/hr]	SNPR [mg/g MLVSS/hr]	PUR [mg/ L/hr]	SPUR [mg/g MLVSS/hr]
0.16	9.30	21.8	1.18	0.66	1.05	0.58	10.2	5.74

7.2 Appendix B

Complete Simulation Dynamic (7/23/09 - 11/5/09)

SIMULATION INPUTS

Primary Effluent Fractions		Input
Name	Default	Value
Fbs - Readily biodegradable (including Acetate) [gCOD/g of total COD]	0.270	0.310
Fac - Acetate [gCOD/g of readily biodegradable COD]	0.150	0.180
Fxsp - Non-colloidal slowly biodegradable [gCOD/g of slowly degradable COD]	0.500	0.610
Fus - Unbiodegradable soluble [gCOD/g of total COD]	0.080	0.155
Fup - Unbiodegradable particulate [gCOD/g of total COD]	0.080	0.169
Fna - Ammonia [gNH3-N/gTKN]	0.750	0.740
Fnox - Particulate organic nitrogen [gN/g Organic N]	0.250	0.300
Fnus - Soluble unbiodegradable TKN [gN/gTKN]	0.020	0.030
FupN - N:COD ratio for unbiodegradable part. COD [gN/gCOD]	0.035	0.030
Fpo4 - Phosphate [gPO4-P/gTP]	0.750	0.820
FupP - P:COD ratio for unbiodegradable part. COD [gP/gCOD]	0.011	0.009

Process	Total Vol.	HRT w/out recycle [hr]	DO-Set Point [mg/L]	Tanks in Service [No.]	% Removal [%]	Sludge Blanket	Underflow - Constant [MGD]
Anaerobic Tanks (3 cells per Train) Phase 1	2.606193	1.17	un-aerated	5	-	-	-
Anaerobic Tanks (3 cells per Train) Phase 2	2.084954	0.91	un-aerated	4			
Anoxic Tanks (3 cells per Train) Phase 1	2.606193	1.17	un-aerated	5	-	-	-

Anoxic Tanks (3 cells per Train) Phase 2	2.084954	0.91	un-aerated	4			
Aeration Tank 1	1.800	1.45	2	2	-	-	-
Aeration Tank 2	1.800	1.45	2	2	-	-	-
Aeration Tank 3	1.800	1.45	1	2	-	-	-
Secondary Clarifier	4.510	3.640	-	2	Variable	0.2	0.00

PARAMETERS (KINETIC)

AOB			
Name	Default	Value	Arrhenius
Max. spec. growth rate [1/d]	0.9	0.62	1.072
Substrate (NH4) half sat. [mgN/L]	0.7	0.7	1
Aerobic decay rate [1/d]	0.17	0.17	1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08	1.029
KiHNO2 [mmol/L]	0.005	0.005	1

NOB			
Name	Default	Value	Arrhenius
Max. spec. growth rate [1/d]	0.7	0.7	1.06
Substrate (NO2) half sat. [mgN/L]	0.1	0.1	1
Aerobic decay rate [1/d]	0.17	0.17	1.029
Anoxic/anaerobic decay rate [1/d]	0.08	0.08	1.029
KiNH3 [mmol/L]	0.075	0.075	1

SPLITTER ITINERARY

AR	CY	NRO	CY	CR	CY	W	AS	Temp	erature	Clarifier	ANA	'ANX
Time	Split	Time	Split	Time	Split	Time	Split	Time	Temp	% Removal	Time	Split
0	18.8	0	19.5	0	11.8	0	0.65	0	26.55	99.9	0	0.2
1	18.8	1	19.5	1	11.5	1	0.68	1	26.58	99.9	1	0.2
2	18.7	2	19.5	2	11.1	2	0.65	2	26.68	99.9	2	0.2
3		3		3		3	0.65	3		99.9	3	0.2
4	19.0	4	19.5	4	11.1	4	0.65	4	26.95	99.9	4	0.2
5	18.9	5	19.5	5	11.1	5	0.65	5	26.95	99.9	5	0.2
6	19.1	6	19.5	6	11.8	6	0.63	6	27.1	99.9	6	0.2
7	19.0	7	19.5	7	11.3	7	0.61	7	27.15	99.9	7	0.2
8	19.0	8	19.5	8	11.2	8	0.65	8	27.18	99.9	8	0.2
9	18.8	9	19.5	9	11.1	9	1.00	9	27.21	99.9	9	0.2
10	19.4	10	19.5	10	11.9	10	1.07	10	27.27	99.9	10	0.2
11	19.5	11	19.5	11	12.4	11	1.10	11	27.31	99.9	11	0.2
12	19.3	12	19.5	12	11.1	12	1.10	12	27.28	99.9	12	0.2
13	19.4	13	19.5	13	11.1	13	1.10	13	27.32	99.9	13	0.2

_													
	14	19.6	14	19.5	14	13.1	14	1.09	14	27.32	99.9	14	0.2
	15	19.1	15	19.5	15	12.0	15	1.08	15	27.34	99.9	15	0.2
	16	19.1	16	19.5	16	11.6	16	1.09	16	27.36	99.9	16	0.2
	17	19.0	17	19.5	17	11.7	17	1.09	17	27.37	99.9	17	0.2
	18	19.0	18	19.5	18	11.7	18	1.09	18	27.34	99.9	18	0.2
	19	19.0	19	19.5	19	11.5	19	1.09	19	27.42	99.9	19	0.2
	20	19.0	20	19.5	20	11.9	20	1.07	20	27.52	99.9	20	0.2
	21	18.9	21	19.5	21	12.4	21	1.10	21	27.62	99.9	21	0.2
	22	18.8	22	19.5	22	12.3	22	1.10	22	27.71	99.9	22	0.2
	23	18.9	23	19.5	23	12.5	23	1.10	23	27.64	99.9	23	0.2
	24	18.9	24	19.5	24	11.9	24	1.10	24	27.52	99.9	24	0.2
	25	18.8	25	19.5	25	11.6	25	1.10	25	27.46	99.9	25	0.2
	26	18.5	26	19.5	26	11.6	26	1.10	26	27.46	99.9	26	0.2
	27	18.9	27	19.5	27	11.5	27	1.10	27	27.48	99.9	27	0.2
	28	18.4	28	19.5	28	11.5	28	1.09	28	27.54	99.9	28	0.2
	29	18.7	29	19.5	29	11.5	29	1.17	29	27.68	99.9	29	0.2
	30	18.5	30	0	30	11.6	30	1.45	30	27.75	99.85	30	0.2
	31	18.3	31	0	31	11.6	31	1.45	31	27.81	99.85	31	0.2
	32	18.1	32	0	32	11.6	32	1.58	32	27.79	99.85	32	0.2
	33	18.2	33	0	33	11.5	33	1.70	33	27.74	99.85	33	0.2
	34	17.9	34	0	34	11.2	34	1.77	34	27.75	99.85	34	0.2
	35	18.3	35	0	35	11.5	35	1.60	35	27.79	99.85	35	0.2
	36	18.0	36	0	36	11.8	36	1.41	36	27.83	99.85	36	0.2
	37	18.3	37	0	37	11.7	37	2.21	37	27.92	99.85	37	0
	38	18.2	38	0	38	11.5	38	2.17	38	27.97	99.85	38	0
	39	18.0	39	0	39	11.8	39	2.16	39	27.89	99.85	39	0
	40	18.0	40	0	40	11.5	40	2.21	40	27.8	99.85	40	0
	41	17.8	41	0	41	11.4	41	2.21	41	27.64	99.85	41	0
	42	17.8	42	0	42	11.3	42	2.08	42	27.45	99.85	42	0
	43	17.9	43	0	43	11.2	43	2.04	43	27.3	99.85	43	0
	44	17.9	44	0	44	11.1	44	2.04	44	27.23	99.85	44	0
	45	18.2	45	0	45	10.8	45	1.95	45	27.21	99.85	45	0
	46	17.9	46	0	46	11.7	46	2.04	46	27.18	99.85	46	0
	47	17.7	47	0	47	14.1	47	2.03	47	26.86	99.85	47	0
	48	17.7	48	0	48	14.0	48	1.99	48	26.5	99.85	48	0
	49	17.6	49	0	49	13.9	49	2.04	49	26.55	99.85	49	0
	50	17.4	50	0	50	13.1	50	2.04	50	26.43	99.85	50	0
	51	17.5	51	0	51	12.4	51	2.04	51	26.37	99.85	51	0
	52	18.1	52	0	52	12.2	52	2.04	52	26.36	99.85	52	0
	53	18.0	53	0	53	12.1	53	2.12	53	26.33	99.85	53	0
	54	17.7	54	0	54	12.0	54	2.25	54	26.34	99.85	54	0
	55	17.8	55	0	55	11.9	55	2.32	55	26.34	99.85	55	0

		1										
56	17.6	56	0	56	12.0	56	2.37	56	26.42	99.85	56	0
57	17.6	57	0	57	11.9	57	2.32	57	26.46	99.85	57	0
58	17.8	58	0	58	12.0	58	2.38	58	26.42	99.85	58	0
59	18.2	59	0	59	11.8	59	2.39	59	26.4	99.85	59	0
60	17.7	60	0	60	11.1	60	2.39	60	26.32	99.85	60	0
61	17.8	61	0	61	11.2	61	2.45	61	26.28	99.85	61	0
62	17.9	62	0	62	11.5	62	2.75	62	26.21	99.85	62	0
63	17.6	63	0	63	11.4	63	2.74	63	26.26	99.85	63	0
64	17.5	64	0	64	12.6	64	2.88	64	26.21	99.85	64	0
65	17.5	65	0	65	12.3	65	3.03	65	26.39	99.85	65	0
66	17.4	66	0	66	12.5	66	3.03	66	26.28	99.85	66	0
67	17.3	67	0	67	12.2	67	3.25	67	26.21	99.85	67	0
68	17.4	68	0	68	12.2	68	3.52	68	26	99.85	68	0
69	17.4	69	0	69	11.7	69	3.47	69	25.85	99.85	69	0
70	17.6	70	0	70	5.9	70	3.43	70	25.78	99.85	70	0
71	17.6	71	0	71	11.5	71	3.42	71	25.62	99.85	71	0
72	17.7	72	0	72	11.7	72	1.80	72	25.59	99.8	72	0
73	17.6	73	0	73	11.5	73	1.80	73	25.56	99.8	73	0
74	17.5	74	0	74	11.2	74	1.75	74	25.53	99.8	74	0
75	17.7	75	0	75	11.2	75	1.62	75	25.4	99.8	75	0
76	17.5	76	0	76	11.2	76	1.51	76	25.39	99.8	76	0
77	17.2	77	0	77	11.2	77	1.55	77	25.31	99.8	77	0
78	17.2	78	0	78	11.2	78	1.57	78	25.26	99.8	78	0
79	17.0	79	0	79	11.4	79	1.54	79	25.22	99.8	79	0
80	17.5	80	0	80	11.2	80	1.55	80	25.2	99.8	80	0
81	17.2	81	0	81	11.2	81	1.55	81	25.16	99.8	81	0
82	17.2	82	0	82	10.9	82	1.40	82	25.02	99.8	82	0
83	16.7	83	0	83	11.3	83	1.48	83	24.8	99.8	83	0
84	17.1	84	0	84	12.0	84	1.56	84	24.71	99.8	84	0
85	17.2	85	0	85	11.9	85	1.53	85	24.32	99.8	85	0
86	17.2	86	0	86	11.7	86	1.49	86	24.03	99.8	86	0
87	17.3	87	0	87	11.8	87	1.52	87	23.72	99.8	87	0
88	17.2	88	0	88	11.6	88	1.52	88	23.46	99.8	88	0
89	17.2	89	0	89	11.2	89	1.43	89	23.08	99.8	89	0
90	17.1	90	0	90	11.2	90	1.42	90	22.95	99.8	90	0
91	17.1	91	0	91	11.3	91	1.41	91	22.84	99.8	91	0
92	16.9	92	0	92	11.4	92	1.60	92	22.89	99.8	92	0
93	16.9	93	0	93	11.5	93	1.79	93	23	99.8	93	0
94	16.8	94	0	94	11.8	94	1.89	94	23.07	99.8	94	0
95	17.0	95	0	95	11.1	95	1.90	95	23.09	99.8	95	0
96	16.6	96	0	96	11.4	96	1.89	96	22.97	99.8	96	0
97	16.6	97	0	97	11.3	97	1.92	97	22.92	99.8	97	0

98	17.2	98	0	98	11.4	98	1.92	98	22.92	99.8	98	0
99	17.1	99	0	99	11.5	99	1.84	99	22.93	99.8	99	0
100	17.1	100	0	100	11.5	100	1.80	100	22.92	99.8	100	0
101	17.1	101	0	101	11.5	101	1.70	101	22.92	99.8	101	0
102	17.1	102	0	102	11.4	102	1.69	102	22.82	99.8	102	0
103	17.1	103	0	103	11.8	103	1.69	103	23.59	99.8	103	0
104	17.1	104	0	104	11.9	104	1.64	104	22.53	99.8	104	0
105	17.1	105	0	105	10.3	105	1.39	105	22.49	99.8	105	0

7.3 Appendix C – Profile Sampling Data and Calculated Rates

Week	Profile	Date	Actual SRT [days]	Aerobic MLSS [mg/L]	Average AER DO [mg/L]	Min AER pH	AUR (NH4-N Slope) [mg/L/hr]	SAUR [mg/g MLVSS/hr]	NPR (NOx-N Slope) [mg/L/hr]	SNPR [mg/g MLVSS/hr]	Highest AER NO ₂ -N Concentration [mg/L]	PUR (PO ₄ -P Uptake Slope) [mg/L/hr]	SPUR [mg/g MLVSS/hr]	Clarifier IDC NH ₄ -N [mg/L]	Clarifier IDC NOx-N [mg/L]	Clarifier IDC PO ₄ -P Concentration [mg/L]
1	1	7/23/2009	12.6	3940	2.09	6.84	-	1	-	-	2.95	-	-	0.00	10.7	0.2
2	1	7/27/2009	14.3	4060	1.9	6.9	1.6	0.6	1.8	0.7	2.44	2.13	0.80	0.00	10.5	0.3
2	2	7/29/2009	12.6	4220	2.1	6.9	1.7	0.6	2.0	0.7	2.1	2.4	0.8	0.0	12.2	0.3
3	1	8/5/2009	9.8	3760	1.6	6.9	1.6	0.6	1.6	0.6	1.4	7.1	2.6	0.0	10.3	0.3
3	2	8/6/2009	9.9	4000	1.4	6.9	1.7	0.6	2.0	0.7	1.7	3.8	1.3	0.0	11.7	0.2
4	1	8/10/2009	8.1	3700	1.8	6.8	1.7	0.7	1.8	0.7	1.4	4.1	1.6	0.0	11.0	0.3
	2	8/12/2009	8.3	3640	2.3	6.9	1.7	0.7	2.2	0.8	1.8	3.8	1.5	0.0	11.8	0.4
5	1	8/17/2009	10.5	3700	1.3	6.8	1.2	0.4	2.2	0.8	0.9	6.1	2.2	0.0	10.2	0.4
3	2	8/20/2009	7.7	3580	1.9	6.8	1.5	0.6	2.9	1.1	0.7	4.7	1.8	0.0	12.1	3.9
6	1	8/24/2009	7.2	3180	1.3	6.7	2.6	1.1	4.0	1.6	0.4	6.9	2.8	0.0	18.5	4.1
0	2	8/26/2009	7.3	3280	1.3	6.7	2.6	1.1	4.5	1.9	0.5	6.9	2.9	1.0	18.9	3.0
7	1	9/1/2009	5.2	2800	1.3	6.7	2.7	1.3	4.7	2.3	1.3	5.4	2.7	0.0	19.7	4.1
	2	9/2/2009	5.4	2800	1.3	6.7	2.5	1.2	4.3	2.1	0.3	4.8	2.4	0.0	19.3	4.5
8	1	9/9/2009	4.3	2600	1.7	6.5	2.2	1.2	2.6	1.3	0.3	4.9	2.6	0.0	11.0	0.0
	1	9/14/2009	5.9	2760	1.8	6.7	2.7	1.3	3.3	1.6	0.3	6.3	3.1	0.0	15.6	0.0
9	2	9/15/2009	6.4	2640	2.0	6.6	2.4	1.2	3.1	1.5	0.4	7.1	3.5	0.0	16.5	0.0
	3	9/18/2009	4.4	2720	1.5	6.7	2.8	1.4	3.0	1.5	0.5	6.5	3.2	0.0	16.1	0.3
10	1	9/22/2009	5.9	3780	1.7	6.7	2.5	0.9	3.5	1.2	0.6	7.3	2.6	0.0	18.1	0.2
10	2	9/24/2009	4.8	2480	2.0	6.7	2.8	1.5	3.7	2.0	1.0	7.0	3.7	0.0	17.4	0.2
11	1	9/28/2009	4.1	2300	2.0	6.7	2.9	1.6	4.0	2.3	1.3	7.0	4.0	0.0	17.2	0.3
- 11	2	9/30/2009	3.6	2120	1.8	6.6	3.0	1.8	3.3	2.0	1.8	6.0	3.7	0.0	16.8	1.1

12	1	10/6/2009	5.0	1760	2.3	6.6	2.7	1.9	3.2	2.3	2.6	7.2	5.1	2.3	16.9	0.1
	2	10/8/2009	4.0	1780	1.7	-	2.2	1.6	2.3	1.7	4.0	7.3	5.4	5.8	13.3	0.4
13	1	10/13/2009	4.4	1700	1.8	6.8	2.0	1.5	2.0	1.5	6.3	8.4	6.2	10.9	11.6	0.1
	2	10/16/2009	4.9	1540	1.4	6.8	1.4	1.2	1.0	0.8	3.2	9.0	7.5	16.1	5.6	0.2
14	1	10/19/2009	4.9	1910	1.6	6.9	1.2	0.8	0.7	0.5	2.0	9.7	6.4	21.0	3.7	0.1
	2	10/21/2009	5.3	1940	1.2	6.9	0.6	0.4	0.4	0.3	1.5	11.0	7.2	24.9	2.9	0.2
15	1	10/26/2009	5.9	2320	1.4	6.9	0.9	0.5	0.6	0.4	1.9	10.3	5.8	24.2	3.3	0.1
	2	10/27/2009	5.2	2140	1.1	7.0	1.1	0.7	0.5	0.3	1.5	9.0	5.5	26.8	2.7	0.2
16	1	11/3/2009	4.0	2660	1.3	6.8	1.3	0.6	1.7	0.9	5.3	9.4	4.7	19.1	7.3	0.1
	2	11/5/2009	4.6	2320	1.0	6.8	1.4	0.8	1.3	0.8	5.3	12.0	7.0	18.8	7.0	0.2

Week	Profile	Date	Max ANA PO ₄ -P Release [mg/L]	Avg. ANA NO ₃ -N Concentration [mg/L]	AA4 NOx-N [mg/L]	AA6 NOx-N [mg/L]	AA1 PO ₄ -P Release Slope [mg/L/hr]	AA1 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA2 PO ₄ -P Release Slope [mg/L/hr]	AA2 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA3 PO ₄ -P Release Slope [mg/L/hr]	AA3 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA4 PO ₄ -P Release Slope [mg/L/hr]	AA4 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA5 PO ₄ -P Release Slope [mg/L/hr]	AA5 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA6 PO ₄ -P Release Slope [mg/L/hr]	AA6 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	(AAI) COD Uptake Slope [mg/L/hr]	(AA1) Specific COD Uptake Rate [mg/g MLVSS/hr]	(AA1) PO ₄ -P/COD [mg/mg]
1	1	7/23/2009	17.8	0.0	1.2	0.1	-	-	-	-	-	-	-	-	-	-	1	-	-	-	-
2	1	7/27/2009	28.4	0.7	2.9	0.5	13.5	14.4	10.5	11.2	2.16	2.31	-1.07	-0.40	-0.05	-0.02	1.03	0.39	50.4	53.8	13.5
2	2	7/29/2009	20.2	0.5	2.3	0.3	16.1	14.9	4.3	4.0	4.08	3.77	-0.14	-0.05	-0.01	0.00	0.58	0.21	40.4	37.5	16.1
3	1	8/5/2009	23.9	0.3	1.5	0.3	12.9	7.0	6.8	3.7	4.74	2.56	-0.30	-0.11	-0.30	-0.11	4.38	1.64	44.8	24.2	12.9
3	2	8/6/2009	28.6	0.3	1.5	0.3	18.5	12.8	9.3	6.4	6.36	4.38	-0.19	-0.06	0.05	0.02	1.07	0.37	67.2	46.3	18.5
4	1	8/10/2009	24.6	0.4	1.9	0.3	13.4	16.4	7.6	9.3	5.46	6.67	0.15	0.06	0.01	0.01	1.24	0.48	59.8	73.2	13.4
4	2	8/12/2009	23.4	0.4	1.7	0.0	13.4	9.1	8.7	5.9	5.25	3.56	0.14	0.05	0.00	0.00	1.03	0.39	46.6	31.6	13.4
5	1	8/17/2009	27.7	0.3	0.7	0.3	15.0	11.0	8.1	5.9	5.74	4.23	0.33	0.12	0.04	0.02	1.15	0.42	64.4	47.4	15.0
3	2	8/20/2009	27.3	0.6	2.5	0.3	14.5	10.1	8.8	6.1	5.98	4.16	-0.18	-0.07	0.26	0.10	1.17	0.44	45.6	31.8	14.5

	1	8/24/2009	35.0	0.2	0.3	0.3	10.9	8.4	11.1	8.5	6.81	5.26	0.98	0.40	0.39	0.16	2.11	0.87	43.1	33.3	10.9
6	2	8/26/2009	28.5	0.3	1.0	0.3	14.9	9.4	8.7	5.5	6.46	4.06	0.09	0.04	0.12	0.05	1.80	0.74	42.7	26.8	14.9
7	1	9/1/2009	27.7	0.1	1.0	0.0	11.8	10.6	11.0	9.8	7.16	6.43	0.08	0.04	0.26	0.13	2.23	1.09	113.5	101.9	11.8
/	2	9/2/2009	30.1	0.6	2.7	0.3	15.9	14.4	13.4	12.1	5.84	5.26	-0.36	-0.18	0.11	0.06	2.02	1.00	43.1	38.9	15.9
8	1	9/9/2009	12.7	0.3	0.4	0.3	8.6	9.8	6.8	7.7	4.27	4.87	-0.25	-0.13	0.12	0.06	1.62	0.84	32.5	37.0	8.6
	1	9/14/2009	20.3	0.4	0.5	0.3	7.7	7.0	9.9	9.0	6.33	5.74	0.38	0.19	0.43	0.21	2.24	1.11	35.3	32.0	7.7
9	2	9/15/2009	21.1	0.4	1.1	0.3	13.5	12.4	9.0	8.3	6.37	5.84	-0.16	-0.08	-0.07	-0.04	2.45	1.25	50.6	46.4	13.5
	3	9/18/2009	25.8	0.3	0.5	0.0	11.6	11.4	11.9	11.6	7.24	7.09	0.18	0.09	0.16	0.08	2.90	1.44	-	-	11.6
10	1	9/22/2009	25.3	0.4	0.4	0.4	7.1	7.9	10.7	11.9	7.08	7.86	1.42	0.50	0.45	0.16	3.25	1.15	41.8	46.4	7.1
	2	9/24/2009	27.1	0.3	0.8	0.3	16.2	17.1	10.9	11.5	7.82	8.25	-0.26	-0.14	0.30	0.16	3.02	1.58	52.6	55.5	16.2
11	1	9/28/2009	22.7	0.3	0.5	0.3	7.8	9.1	10.1	11.7	7.11	8.25	1.87	1.06	0.40	0.23	2.77	1.56	28.3	32.8	7.8
	2	9/30/2009	24.6	0.3	0.7	0.4	15.2	20.8	8.7	12.0	6.83	9.36	1.71	1.06	1.66	1.03	1.37	0.85	35.4	48.5	15.2
12	1	10/6/2009	20.2	0.3	0.4	0.3	4.0	4.6	9.0	10.4	6.92	8.02	2.95	2.09	0.22	0.16	2.58	1.83	16.3	18.9	4.0
	2	10/8/2009	20.6	0.3	0.7	0.3	11.0	13.5	10.5	12.8	7.50	9.17	-0.03	-0.02	-0.22	-0.16	3.36	2.49	23.8	29.1	11.0
13	1	10/13/200 9	31.0	0.4	0.4	0.4	11.2	14.9	10.6	14.1	7.12	9.52	1.96	1.46	0.47	0.35	3.54	2.63	35.3	47.2	11.2
13	2	10/16/200 9	17.6	0.3	0.3	0.3	-0.3	-0.4	12.6	16.9	4.48	5.99	3.21	2.67	0.69	0.57	2.64	2.20	-	-	-0.3
	1	10/19/200	21.0	0.3	0.3	0.3	0.1	0.1	9.2	11.6	6.45	8.14	2.98	1.97	1.15	0.76	2.87	1.90	0.8	1.0	0.1
14	2	10/21/200 9	21.7	0.4	0.4	0.4	5.0	6.5	11.3	14.5	7.28	9.35	2.21	1.44	0.73	0.47	5.19	3.39	44.6	57.2	5.0
	1	10/26/200	32.5	0.3	0.3	0.3	9.5	5.2	13.9	7.6	7.41	4.07	2.48	1.39	1.43	0.80	2.43	1.36	42.4	23.3	9.5
15	2	10/27/200 9	22.8	0.0	0.0	0.0	10.3	11.0	12.0	12.8	7.58	8.11	0.96	0.58	0.15	0.09	3.00	1.82	43.1	46.1	10.3
16	1	11/3/2009	31.7	0.1	0.0	0.0	14.6	12.4	13.2	11.2	8.79	7.48	0.11	0.06	0.48	0.24	2.97	1.49	43.5	37.0	14.6
10	2	11/5/2009	34.3	0.3	0.3	0.3	9.0	8.1	14.4	13.0	7.25	6.52	2.52	1.47	0.33	0.19	4.11	2.40	25.6	23.0	9.0

7.4 Appendix D - Plant Characteristics during Profile Sampling

Week	Profile	Date	# Chem Used [lbs]	FeCl3 Hrs Used [hr]	FeCl3 Addition [mg/L]	Primary Clarifier Effluent Flow [MGD]	ARCY Flow [MGD]	CRCY Flow [MGD]	NRCY Flow [MGD]	ANA MLTSS [mg/L]	ANA % VOL [%]	ANA MLVSS [mg/L]	ANX MLTSS [mg/L]	ANX % VOL [%]	ANX MLVSS [mg/L]	AER MLTSS [mg/L]	AER % VOL [%]	AER MLVSS [mg/L]
1	1	7/23/2009	1000.0	24.0	4.0	18	19	12	20	2360	69	1628	3940	65	2561	3940	65	2561
2	1	7/27/2009	0.0	0.0	0.0	18	19	11	20	1320	71	937	4060	66	2680	4060	66	2680
	2	7/29/2009	0.0	0.0	0.0	19	19	12	20	1500	72	1080	4220	66	2785	4220	66	2785
3	1	8/5/2009	0.0	0.0	0.0	18	19	11	20	2440	76	1854	3760	71	2670	3760	71	2670
	2	8/6/2009	0.0	0.0	0.0	22	20	13	20	1960	74	1450	4000	72	2880	4000	72	2880
4	1	8/10/2009	0.0	0.0	0.0	19	19	12	20	1090	75	818	3700	70	2590	3700	70	2590
	2	8/12/2009	0.0	0.0	0.0	20	19	12	20	1820	81	1474	3640	72	2621	3640	72	2621
5	1	8/17/2009	0.0	0.0	0.0	19	19	12	20	1720	79	1359	3700	75	2775	3700	75	2775
	2	8/20/2009	0.0	0.0	0.0	19	18	12	20	1840	78	1435	3580	74	2649	3580	74	2649
6	1	8/24/2009	0.0	0.0	0.0	19	18	12	0	1640	79	1296	3180	76	2417	3180	76	2417
	2	8/26/2009	0.0	0.0	0.0	19	18	11	0	2040	78	1591	3280	74	2427	3280	74	2427
7	1	9/1/2009	0.0	0.0	0.0	19	18	11	0	1410	79	1114	2800	73	2044	2800	73	2044
	2	9/2/2009	0.0	0.0	0.0	19	18	11	0	1370	81	1110	2800	72	2016	2800	72	2016
8	1	9/9/2009	1500.1	22.0	5.1	25	18	14	0	1140	77	878	2600	74	1924	2600	74	1924
9	1	9/14/2009	0.0	0.0	0.0	20	18	12	0	1360	81	1102	2760	73	2015	2760	73	2015
	2	9/15/2009	0.0	0.0	0.0	20	18	12	0	1380	79	1090	2640	74	1954	2720	74	2013
	3	9/18/2009	0.0	0.0	0.0	20	18	12	0	1380	74	1021	2720	74	2013	2720	74	2013
10	1	9/22/2009	0.0	0.0	0.0	19	18	11	0	1200	75	900	3780	75	2835	3780	75	2835
	2	9/24/2009	0.0	0.0	0.0	20	18	11	0	1230	77	947	2480	77	1910	2480	77	1910
11	1	9/28/2009	0.0	0.0	0.0	21	17	12	0	1120	77	862	2300	77	1771	2300	77	1771
	2	9/30/2009	0.0	0.0	0.0	20	17	12	0	960	76	730	2120	76	1611	2120	76	1611
12	1	10/6/2009	0.0	0.0	0.0	19	18	11	0	980	88	862	1760	80	1408	1760	80	1408
	2	10/8/2009	0.0	0.0	0.0	19	17	11	0	940	87	818	1780	76	1353	1780	76	1353

13	1	10/13/2009	0.0	0.0	0.0	19	17	11	0	880	85	748	1700	79	1343	1700	79	1343
	2	10/16/2009	0.0	0.0	0.0	20	17	12	0	840	89	748	1540	78	1201	1540	78	1201
14	1	10/19/2009	0.0	0.0	0.0	20	17	12	0	890	89	792	1910	79	1509	1910	79	1509
	2	10/21/2009	0.0	0.0	0.0	19	17	11	0	865	90	779	1940	79	1533	1940	79	1533
15	1	10/26/2009	0.0	0.0	0.0	19	17	11	0	2140	85	1819	2320	77	1786	2320	77	1786
	2	10/27/2009	0.0	0.0	0.0	19	17	11	0	1100	85	935	2140	77	1648	2140	77	1648
16	1	11/3/2009	0.0	0.0	0.0	19	17	12	0	1400	84	1176	2660	75	1995	2660	75	1995
	2	11/5/2009	0.0	0.0	0.0	19	17	10	0	1340	83	1112	2320	74	1717	2320	74	1717

7.5 Appendix E - QAC Addition Calculation

Nansemond Pl	ant						
AVG Flo	w Chem. Toilet Waste		Chemical Liquid Addition				
[gal/day]	[gal/day]		2 oz per 5 gal				
18500000	3,539		59.14 mL per 18.925 L				
			Chemical Stock 3.12 mL/L				
Dose 1							
			Assumptions:				
Peak Duration	20	days	Receive peak waste during one month of the year. Receive				
Duration/day	4	hr	the waste out of the 20 days in 4 hour periods throughout				
Reactor Size	3	L	the 8 hour workday.				
Chem Dose	11796.7	gal					
Flow Volume	370000000	gal					
Percentage	0.00319						
Reactor Dose	9.56	mL					
Dose 2			Assumptions:				
Peak Duration	40	days	Receive peak waste during two months of the summer.				
Duration/day	5	hr	Receive the waste out of the 40 days in 5 hour periods				
Reactor Size	3	L	throughout the 8 hour workday				
Chem Dose	29491.7	gal					
Flow Volume	740000000	gal					
Percentage	0.00797						
Reactor Dose	23.91	mL					
Dose 2			Assumptions:				
Peak Duration	60	days	Receive peak waste during summer months (June, July,				
Duration/day	6	hr	August) when use of portable toilets is at a peak. Receive				
Reactor Size	3	L	the waste out of the 60 days in 6 hour periods throughout				
Chem Dose	53085.0	gal	the 8 hour workday.				
Flow Volume	1110000000	gal					
Percentage	0.01435						
Reactor Dose	43.04	mL					

7.5 Appendix F – Iron Addition Calculation and Plan

NS Treatment plant Iron Addition

Iron addition rate =	3000	lbs/day Fe
Flow Rate =	19	MGD
Fe Dose =	18.93	mg/L Fe

FeCl3 Stock Solution in Lab

FeCl3 Stock =	341	g/L FeCl3
As $Fe =$	117	g/L Fe

Day 1

	NS Mixed	Fe Dose	Fe Stock	VIP SE
Reactor	Liquor (L)	(mg/L Fe)	(uLs)	(L)
A	1	0	0	2
В	1	20	511	2
C	1	35	895	2
D	1	50	1278	2

- Add 1.0 L NS mixed liquor to each reactor
- Add 2 L of VIP SE, and start pH and DO control
- Wait 10 Min
- Record pH
- Add FeCl3 stock solution to Reactors B, C, D
- Add Corresponding Alkalinity
- Record pH reduction immediately after Fe Addition
- Wait 2 hours
- -Start experiment as normal

Day 2

	VIP Mixed	Fe Dose	Fe Stock	VIP SE
Reactor	Liquor (L)	(mg/L Fe)	(uLs)	(L)
A	1	0	0.0	2
В	1	20	170.4	2
C	1	35	298.2	2
D	1	50	426.0	2

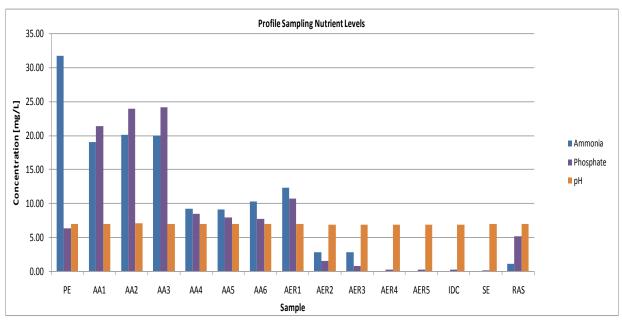
- Add 1.0 L VIP mixed liquor to each reactor
- Start aeration and wait ~10 min
- measure pH by dipping calibrated probe into top of reactor
- Add FeCl3 stock solution to Reactors B, C, D
- Wait 2 hours
- measure pH by dipping calibrated probe into top of reactor

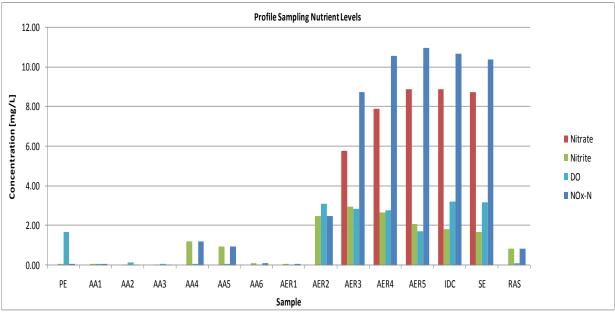
- Add 2 L of VIP SE, and start pH and DO control
- Wait for all pH and DO to stabilize at setpoint
- Wait at least 15-20 min
- -Start experiment as normal

7.6 Appendix G – Weekly Profile Sampling Data

Week 1 – 7/23/09

Date	Sample	Time	Time from Start to Finis	h	Sample Number	Reac	ctor Type		monia by CH TNT		rate by CH TNT		Nitrite by HACH PP	+	te HACH Nitrite IACH	Phos	sphate by CH TNT		DO		рН
dd-mmm-yy	h:m	ım	min					ma/	L NH3-N	ma/l	L NO3-N	m	ng/L NO2-N	mg/	L NOx-N	ma/	L PO4-P		[mg/L]		[unitless]
23-Jul-09	7:1		0		PE		-		31.80		0.00	† · ·	0.07	_ ŭ	0.07		6.30	_	1.66		6.99
23-Jul-09	7:2		9		AA1	Ana	aerobic		19.10		0.00		0.05	_	0.05		21.46	_	0.06		7.00
23-Jul-09	7:5		43		AA2		aerobic	_	20.10		0.00		0.03	_	0.03		23.98	_	0.12		7.05
23-Jul-09	8:3		77		AA3		aerobic	_	20.00		0.00		0.03	_	0.03		24.15		0.05		7.01
23-Jul-09	9:0		111		AA4		noxic	_	9.25		0.00		1.21	_	1.21		8.48		0.06		6.98
23-Jul-09	9:2		129		AA5		noxic		9.10		0.00		0.95	_	0.95		7.90		0.07		6.97
23-Jul-09	9:4	12	147		AA6		noxic	1	10.30		0.00		0.08		0.08		7.73		0.03		6.94
23-Jul-09	10:	00	165		AER1	Ae	erobic	1	12.35	-	0.00		0.06		0.06	,	10.70		0.04		7.01
23-Jul-09	10:	32	197		AER2	Ae	erobic		2.87		0.00		2.48		2.48		1.53		3.10		6.87
23-Jul-09	11:	04	229		AER3	Ae	erobic		2.84	,	5.77		2.95		8.72		0.82		2.83		6.85
23-Jul-09	11:	36	261		AER4	Ae	erobic		0.00		7.89		2.65		10.54		0.25		2.78		6.84
23-Jul-09	12:	08	293		AER5	Ae	erobic		0.00		8.86		2.08		10.94		0.25		1.71		6.89
23-Jul-09	12:	11	296		IDC		-		0.00		8.86		1.81		10.67		0.24		3.20		6.90
23-Jul-09	15:	11	476		SE		-		0.00		8.71		1.65		10.36		0.18		3.18		7.02
23-Jul-09	15:	27	492		RAS		-		1.16	-	0.00		0.81		0.81		5.16		0.10		6.98
Actual S	RT	Ae	erobic MLSS [mg/L]		Average A		00		AER pH	Al	UR (NH4				AUR MLVSS/I	_	PR (NOx [mg/L				SNPR MLVSS/hr]
12.60			3940	-	2.0		+		6.84		Įmg/	L/111	'J ['	ng/g n	1LV 33/1	"]	[IIIg/L	1111111	+	ilig/g	IVIL V OO/TII]
		DUI			2.0		:rID0				Olevice.	. ID	0.00 0	M A	NA DO		ANIA NO		ı		
Highest AER N Concentra			R (PO ₄ -P ke Slope)		SPUR	Clan	rifier IDC N	NH4	Clarifier NOx-l				C PO₄-P tration		NA PO ₄ - elease		ANA NO		A4 NC	x-N	AA6 NOx-N
[mg/L]		[m	ng/L/hr]	mg/	/g MLVSS/h) r	[mg/L]		[mg/L			[mg/			ıg/L]		[mg/L]	_	[mg/L		[mg/L]
2.95							0.0000		10.670	00		0.2	2	17	7.85		0.000		1.21		0.08
AA1 F Release [mg/	Slo			ea	cific PC se Rate _VSS/h)	Rele	ease	O ₄ -P e Slope _/hr]	е	P Re	ele	ecific F ase Ra ILVSS	ate	R	3 PC elea Slop ng/L/	se e	Р	Rele	eas	fic PO ₄ - e Rate /SS/hr]
AA4 PO ₄ -P Release Slope [mg/L/hr]	AA4 P F	Releas	ific PO ₄ - se Rate VSS/hr]	Re	5 PO ₄ -P elease Slope g/L/hr]	PR	Specifi elease g MLV	Rate	SI	PO ₄ - lease lope g/L/hr	AA(Rel	pecific Poease Rat	e U	(AA1) C ptake S [mg/L/	Slope	(AA1 COD U [mg/g		Rate	(/	A1) PO ₄ -P/ AA1) COD [mg/mg]
# Chen	n Us	ed	FeCl3	Hr	s Used	4 k	FeC Addi		В		ive olid t Flov	~	ARC	ΥF	low	CR	CY F	low	N	RC	Y Flow
[lb	s1			[hr	·1	十	[mg	/ 1		MC	3D1		[N	1GD	, †		MGD		1	[N/	IGD]
10					000	╅	4.0		1 '	18.	_			3.84			1.832		1		9.5
ANA MLTS:			A % OL		ANA ILVSS		AN MLT	1X	А	NX VO	%	N	ANX 1LVS		AE MLT	R	AI	ER ' √OL			AER ILVSS
[mg/L]	['	%]	_[mg/L]		[mg	/L]		[%]]	[mg/L]		[mg	/L]		[%]		_[mg/L]
2360	T	6	69		1628		39	40		65			2561	$\Box I$	394	40		65			2561

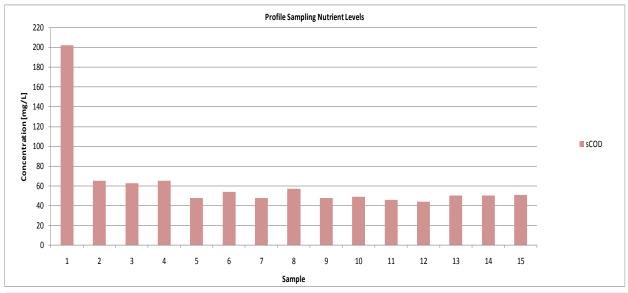


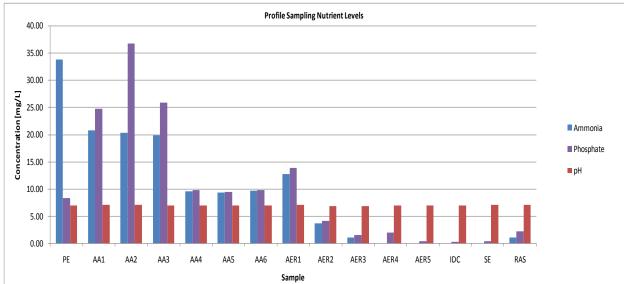


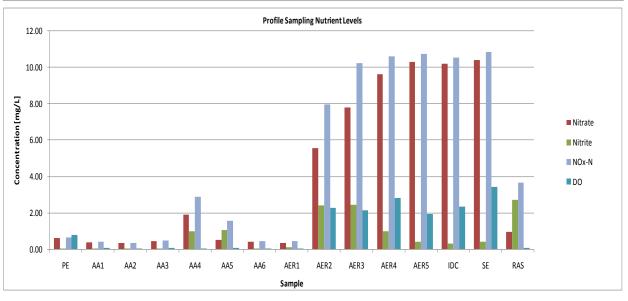
Week 2 – 7/27/09 & 7/29/09

	Train 7, Aera	tion Tank 4, Cla	rifier 4								
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Type	Ammonia by HACH TNT	Nitrate by HACH TNT	Nitrite by HACH PP	NOx-N by Nitrate HACH + Nitrite HACH	Phosphate by HACH TNT	DO	рН
dd-mmm-yy	h:mm	hr			mg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg/L NOx-N	mg/L PO4-P	[mg/L]	[unitless]
27-Jul-09	7:46	0	PE		33.80	0.61	0.02	0.64	8.35	0.79	7.01
27-Jul-09	8:13	1.15	AA1	Anaerobic	20.80	0.39	0.02	0.41	24.72	0.07	7.08
27-Jul-09	8:43	2.31	AA2	Anaerobic	20.30	0.33	0.01	0.34	36.79	0.06	7.05
27-Jul-09	9:13	3.46	AA3	Anaerobic	19.90	0.45	0.03	0.48	25.94	0.08	7.03
27-Jul-09	9:43	4.61	AA4	Anoxic	9.60	1.92	0.98	2.90	9.79	0.05	7.01
27-Jul-09	10:00	5.76	AA5	Anoxic	9.35	0.51	1.05	1.56	9.50	0.09	7.02
27-Jul-09	10:19	6.92	AA6	Anoxic	9.65	0.41	0.05	0.46	9.85	0.05	7.04
27-Jul-09	10:35	0.00	AER1	Aerobic	12.75	0.35	0.11	0.46	13.87	0.05	7.05
27-Jul-09	11:05	1.79	AER2	Aerobic	3.69	5.56	2.40	7.96	4.11	2.28	6.86
27-Jul-09	11:35	3.57	AER3	Aerobic	1.12	7.79	2.44	10.23	1.61	2.14	6.87
27-Jul-09	12:05	5.36	AER4	Aerobic	0.00	9.62	0.98	10.60	2.00	2.83	6.96
27-Jul-09	12:35	7.15	AER5	Aerobic	0.00	10.30	0.42	10.72	0.46	1.95	6.96
27-Jul-09	12:40		IDC	-	0.00	10.20	0.32	10.52	0.32	2.33	6.98
27-Jul-09	15:46		SE	-	0.00	10.40	0.42	10.82	0.42	3.41	7.11
27-Jul-09	16:13		RAS		1.16	0.95	2.72	3.67	2.22	0.07	7.07
		U. A.	NAO	•	1.10	0.30	2.12	3.07	2.22	0.07	7.07
*NOTE: These valu	•	-									
QA/QC S	ample:	SE									
EDΔ	365.1	ED	A 350.1	FI	PA 353.2	ED	A 353.2	ED	A 353.2		EPA 353.2
LIA	303.1	147	1 330.1	La	H 333.2	LA	A 333.2	1.1.2	1 333.2	+	LI A 333.2
sCOD	by CEL	C)-PO ₄		NH ₄ -N	N	O _{2,3} -N	N	O ₃ -N		NO ₂ -N
	g/L]	٠,	ng/L	+	mg/L	_	mg/L	+ -	ng/L	+	mg/L
		1	ig/L		mg/L		nig/L	1	ilg/L		nig/L
	02										
	65										
·	20	_									
	53										
6	65										
6	65 48										
6 4 5	55 48 54										
6 4 5	65 48										
6 4 5	55 48 54										
6 2 5 2	55 48 54 48										
6 4 5 4 5	55 48 54 48 57										
£ £ £ £ £ £ £	65 48 54 48 57 48										
5 4 5 4 4 4 4	55 48 54 48 57 48 49										
E 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	55 48 54 48 57 48 49 46										
5 2 5 4 4 4 4 5	55 18 54 18 57 18 19 16 14										
5 5 4 5 4 4 4 4 5 5 6 7 7 8 7 8 8 7 8 8 7 8 8 7 8 8 8 8 8 8	55 148 54 148 57 148 149 146 144 50		_		<0.20		12.3		11.9		0.43
5 5 4 5 4 4 4 4 5 5 6 7 7 8 7 8 8 7 8 8 7 8 8 7 8 8 8 8 8 8	55 18 54 18 57 18 19 16 14		-		<0.20		12.3		11.9		0.43
5 5 4 5 4 4 4 4 5 5 6 7 7 8 7 8 8 7 8 8 7 8 8 8 8 8 8 8 8 8	55 148 54 148 57 148 149 146 144 50				<0.20		12.3		11.9		0.43
5 5 4 5 4 4 4 4 5 5 6 7 7 8 7 8 8 7 8 8 7 8 8 7 8 8 8 8 8 8	55 148 54 148 57 148 149 146 144 50				<0.20	AUR			11.9		0.43
5 5 4 5 4 4 4 4 5 5 6 7 7 8 7 8 8 7 8 8 7 8 8 8 8 8 8 8 8 8	55 48 54 48 57 48 49 46 44 50 50	erobic MLSS	- Average A	ER DO	<0.20		12.3 (NH4-N ppe)	SAUR	11.9 NPR (NOx-N	I Slope)	0.43 SNPR
6 4 4 5 5 4 4 4 4 4 5 5 5 5 5 5 5 5 5 5	55 48 54 48 57 48 49 46 44 50 50	erobic MLSS [mg/L]	- Average A			Slo	(NH4-N ope)				
66	55 48 54 48 57 48 49 46 44 50 50			L]	Min AER pH	Slo [mg	(NH4-N ope)	SAUR	NPR (NOx-N	nin]	SNPR
Actual SI [days] 14.31 Highest AEI Concentr	655 48 54 48 57 48 49 46 44 50 50 51 RRT A	[mg/L] 4060 PUR (PO ₄ -P ptake Slope)	[mg/	L] 55 Clar	Min AER pH [unitless] 6.86	Slo [mg 1.	(NH4-N ppe) [m 63 C NOx-N	SAUR ng/g MLVSS/hr] 0.61 Clarifier IDC PO. Concentration	NPR (NOx-N [mg/L/n 1.83 1-P Max AN Rele	A PO ₄ -P	SNPR [mg/g MLVSS/hr] 0.68 Avg. ANA NO ₃ -N Concentration
66	655 48 54 48 57 48 49 46 44 50 51 RT A R NO ₂ -N I U U L]	[mg/L] 4060 PUR (PO ₄ -P	[mg/	L] 55 Clar	Min AER pH [unitless] 6.86	Side [mg	(NH4-N ppe) [m 63 C NOx-N //L]	SAUR ng/g MLVSS/hr] 0.61 Clarifier IDC PO.	NPR (NOx-N [mg/L/n 1.83	A PO ₄ -P	SNPR [mg/g MLVSS/hr] 0.68 Avg. ANA NO ₃ -N

AA4 NOx-N	AA6 NOx-N [mg/L]	AA1 Pon Release	Slope /hr]	AA1 Specific PO P Release Rate [mg/g MLVSS/hr	Release Slope [mg/L/hr]	÷	AA2 Specific F P Release Ra [mg/g MLVSS	ate	AA3 PC Release [mg/L	Slope /hr]	P Re	pecific PO ₄ - lease Rate MLVSS/hr]
2.90	0.46	13.5) ડ	14.43	10.50		11.21		2.16)		2.31
AA4 PO ₄ -P Release Slope [mg/L/hr] -1.07	AA4 Specific PC Release Rate [mg/g MLVSS/l	Slop	e /hr]	AA5 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr] -0.02	AA6 PO ₄ -P Release Slope [mg/L/hr] 1.03		6 Specific PO ₄ -P Release Rate ng/g MLVSS/hr] 0.39	. ,	COD Uptake Slope [mg/L/hr] 50.40	(AA1) S COD Upta [mg/g MI 53.	ake Rate _VSS/hr]	` '
# Chem Uso		Hrs Used		l3 Addition [mg/L]	Active Biosol Influent Flov [MGD]		ARCY FI		CRCY [MG	D]	[1	CY Flow MGD]
0	0	000		0.000	18.11		19.02		11.1	17		19.5

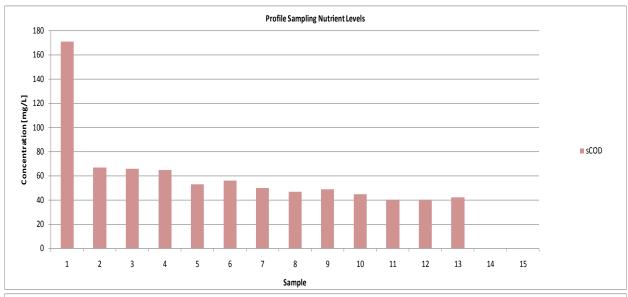


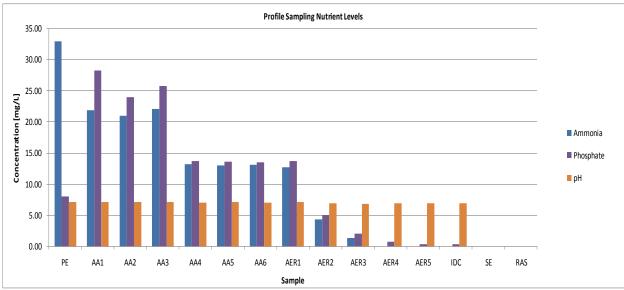


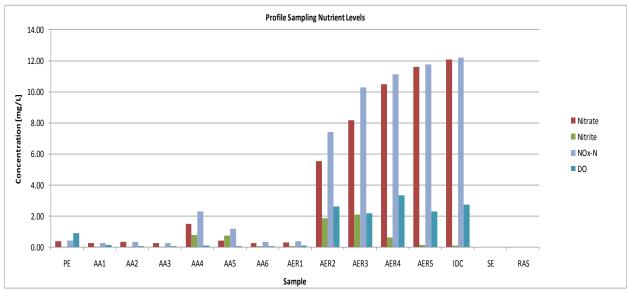


		tion Tank 5, Cla	TITICI J										
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Typ	e HACH TNT	Nitrate by HACH TNT	Nitrite by HAC	HACH	Ph F	osphate by IACH TNT	DO		рН
dd-mmm-yy	h:mm	hr	25		mg/L NH3-N	mg/L NO3-N	mg/L NO2-N		m	g/L PO4-P	[mg/L		[unitless]
29-Jul-07	7:15	0	PE		33.00	0.39	0.03	0.42		8.08	0.92		7.19
29-Jul-07	7:24	1.08	AA1	Anaerobic	21.90	0.27	0.02	0.28		28.23	0.13		7.15
29-Jul-07	7:58	2.16	AA2	Anaerobic	21.00	0.35	0.01	0.36		23.98	0.07		7.11
29-Jul-07	8:32	3.25	AA3	Anaerobic	22.10	0.26	0.01	0.26		25.78	0.06		7.11
29-Jul-07	9:06	4.33	AA4	Anoxic	13.20	1.52	0.79	2.31		13.71	0.09		7.09
29-Jul-07	9:24	5.41	AA5	Anoxic	13.00	0.44	0.73	1.17		13.64	0.07		7.10
29-Jul-07	9:42	6.49	AA6	Anoxic	13.10	0.28	0.05	0.33		13.48	0.06		7.07
29-Jul-07	10:00	0.00	AER1	Aerobic	12.70	0.30	0.08	0.38		13.71	0.10		7.10
29-Jul-07	10:32	1.79	AER2	Aerobic	4.34	5.55	1.88	7.43		5.06	2.64		6.91
29-Jul-07	11:04	3.57	AER3	Aerobic	1.37	8.17	2.10	10.27		2.03	2.20		6.89
29-Jul-07	11:36	5.36	AER4	Aerobic	0.00	10.50	0.62	11.12		0.72	3.34		6.98
29-Jul-07	12:08	7.15	AER5	Aerobic	0.00	11.60	0.16	11.76		0.41	2.31		6.96
29-Jul-07	12:11		IDC	-	0.00	12.10	0.12	12.22		0.32	2.74		6.97
SAMPLE NOT COL	LECTED		SE	-	0.00	0.00	0.00	0.00		0.00	0.00		0.00
SAMPLE NOT COL	LECTED		RAS		0.00	0.00	0.00	0.00		0.00	0.00		0.00
*NOTE: These value	es are <1 mg/L N	JH ₄ -N											
QA/QC Sa	•	NONE											
	-	NONE											
EPA :	365.1	EP A	A 350.1		EPA 353.2	EP	A 353.2	EF	PA 3	53.2		EPA 3	53.2
sCOD b	•	_	-PO ₄		NH ₄ -N		O _{2,3} -N		NO ₃			NO ₂ ·	
[mg		n	ng/L		mg/L		mg/L		mg/	L		mg/	L
17	71		-		-		-		-			-	
6	7												
6	6												
6													
5	3												
5	6												
5													
4													
4	9												
4	5												
4													
4													
4	2												
C)												
)												
Actual SR	RT A	erobic MLSS	Average A		Min AER pH	Slo	(NH4-N ope)	SAUR		PR (NOx-N			NPR
Actual SR	T A	[mg/L]	[mg/	L]	[unitless]	Slo [mg	ope) /L/hr] [mg/g MLVSS/hi		[mg/L/m		[mg/g N	MLVSS/hr]
Actual SR	ET A			L]		Slo [mg	ope)					[mg/g N	
Actual SR [days] 12.59 Highest AER Concentra	R NO ₂ -N F	[mg/L] 4220 PUR (PO ₄ -P ptake Slope)	[mg/ 2.1:	L] 2 C	[unitless] 6.89 arifier IDC NH ₄ -N	Slo [mg 1.	ppe) [166 C NOx-N	0.60 Clarifier IDC PC	r] D ₄ -P	[mg/L/m 1.96 Max ANA Relea	in] A PO ₄ -P	[mg/g M	MLVSS/hr] 0.70 .NA NO ₃ -N centration
Actual SR [days] 12.59 Highest AER	R NO ₂ -N Fation U	[mg/L] 4220 PUR (PO ₄ -P	[mg/	L] 2 C	[unitless] 6.89	Slo [mg	Depe) [[[] [] [] [] [] [] [] [] [mg/g MLVSS/hi 0.60 Clarifier IDC PC	r] D ₄ -P	[mg/L/m 1.96 Max ANA	in] PO ₄ -P ase	Avg. A	MLVSS/hr] 0.70 .NA NO ₃ -N

			AA1 PC) ₄ -P	AA1 Specific	PO ₄ -	AA2 PC	O ₄ -P		AA2 Sp	ecific PO ₄ -	A	A3 PC	D₄-P	AA3 S	Specific PO ₄ -
AA4 NOx-N	Α	A6 NOx-N	Release	Slope	P Release F	Rate	Release	Slope		P Rele	ease Rate	Rele	ease :	Slope	P Re	elease Rate
[mg/L]		[mg/L]	[mg/L/	/hr]	[mg/g MLVS	S/hr]	[mg/L/	/hr]		[mg/g N	//LVSS/hr]	[mg/L/	/hr]	[mg/g	MLVSS/hr]
2.31		0.33	16.1	3	14.94		4.30)		3	3.98		4.08	3		3.77
AA4 PO ₄ -P Release Slope		I Specific PO ₄ -F Release Rate	AA5 PO ₄ -P		AA5 Specific PC Release Rate		AA6 PO ₄ -P Rel Slope	lease i		Specific Release R) COD U Slope	ptake	(AA1) S COD Upta		(AA1) PO ₄ -P/ (AA1) COD
[mg/L/hr]	[m	g/g MLVSS/hr]	[mg/L/	hr]	[mg/g MLVSS/	hr]	[mg/L/hr]		[mo	g/g MLVS	SS/hr]	[mg/L/hr]]	[mg/g ML	VSS/hr]	[mg/mg]
-0.14		-0.05	-0.0	1	0.00		0.58			0.21		40.45		37.	45	0.40
# Chem Us	ed	FeCl3 H	rs Used	FeC	l3 Additior		Active Bio		-	ARC	CY Flow	CR	RCY	Flow	NR	CY Flow
[lbs]		[h	r]		[mg/L]		[MGE	D]		[]	ИGD]		[MG	SD]	[MGD]
0		0.0	00		0.000		19.29	9		1	9.11	1	11.8	11		19.5
ANA MLTSS		NA % VOL	ANA MLVS:	S	ANX MLTSS	-	NX % VOL	-	AN LV	IX 'SS	AEI MLTS	-		ER % VOL	N	AER MLVSS
[mg/L]		[%]	[mg/L]	[mg/L]		[%]	[n	ng	/L]	[mg/	L]		[%]		[mg/L]
1500		72	1080		4220		66	2	278	35	422	0		66		2785



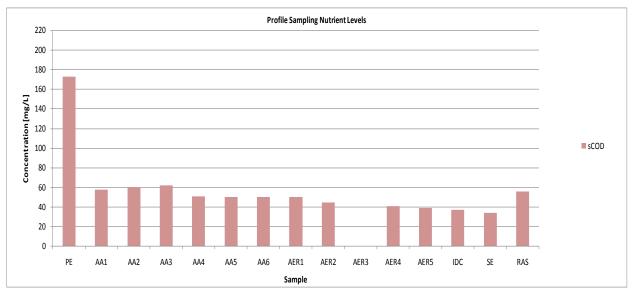


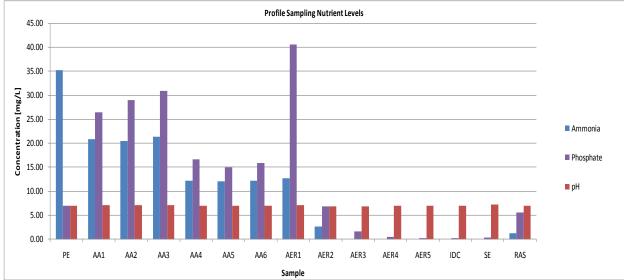


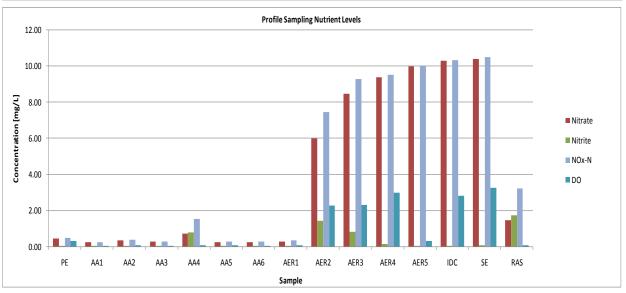
Week 3 – 8/5/09 & 8/6/09

rkutile 1:	Train 5, Aerat	ion rank 5, Gia									
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Type	Ammonia by HACH TNT	Nitrate by HACH TNT	Nitrite by HACH PP	NOx-N by Nitrate HACH + Nitrite HACH	Phosphate by HACH TNT	DO	pH
dd-mmm-yy	h:mm	hr			mg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg/L NOx-N	mg/L PO4-P	[mg/L]	[unitless]
5-Aug-09	7:46	0	PE		35.20	0.44	0.04	0.47	7.02	0.30	6.97
5-Aug-09	8:13	1.15	AA1	Anaerobic	20.90	0.24	0.01	0.25	26.43	0.06	7.06
5-Aug-09	8:43	2.30	AA2	Anaerobic	20.50	0.35	0.02	0.37	29.04	0.07	7.05
5-Aug-09	9:13	3.45	AA3	Anaerobic	21.30	0.27	0.02	0.29	30.92	0.06	7.05
5-Aug-09	9:43	4.60	AA4	Anoxic	12.15	0.73	0.80	1.53	16.64	0.07	7.03
5-Aug-09	10:00	5.75	AA5	Anoxic	12.05	0.24	0.03	0.27	14.95	0.07	7.03
5-Aug-09	10:19	6.90	AA6	Anoxic	12.20	0.25	0.02	0.27	15.89	0.06	7.01
5-Aug-09	10:35	0.00	AER1	Aerobic	12.75	0.30	0.04	0.34	40.63	0.07	7.04
5-Aug-09	11:05	1.78	AER2	Aerobic	2.66	6.00	1.44	7.44	6.79	2.28	6.89
5-Aug-09	11:35	3.56	AER3	Aerobic	0.00	8.45	0.82	9.27	1.66	2.32	6.88
5-Aug-09	12:05	5.34	AER4	Aerobic	0.00	9.36	0.16	9.52	0.44	2.97	6.92
5-Aug-09	12:35	7.13	AER5	Aerobic	0.00	9.98	0.04	10.02	0.26	0.30	6.94
5-Aug-09	12:40		IDC	-	0.00	10.30	0.03	10.33	0.26	2.80	6.96
5-Aug-09 5-Aug-09	15:46 16:13		SE RAS		0.00 1.23	10.40 1.48	0.08 1.74	10.48 3.22	0.30 5.61	3.24 0.08	7.23
	ues are <1 mg/L N	HN	KAS	•	1.23	1.40	1.74	3.22	0.01	0.00	7.05
QA/QC S	•	SE									
	365.1		A 250 1	EX) A 252 2	ED	A 252.2	ED	A 353.2		EDA 252.2
EFA	303.1	EFA	A 350.1	Er	PA 353.2	Er	A 353.2	Er.	A 333.2	1	EPA 353.2
	by CEL	O) -PO ₄		NH ₄ -N		O _{2,3} -N		IO ₃ -N		NO ₂ -N
[m	g/L]	n	ng/L		mg/L	1	mg/L	1	ng/L		mg/L
	73 58 60 62										
\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\	51 50 50 50 45 45 41 39										
<	50 50 50 45 225 41 39 37	C	0.234		<0.20		11.2		11.2		<0.01
<	50 50 50 45 45 :25 41 39	(0.234		<0.20		11.2		11.2		<0.01
<	50 50 50 45 225 41 39 37).234		<0.20		11.2		11.2		<0.01
<	50 50 50 45 45 225 41 39 37 34	Cobic MLSS	0.234 Average AER		<0.20	AUR (NH4-N S	Т	SAUR	11.2 NPR (NOx-N	Slope)	<0.01
Actual SI	50 50 50 45 45 225 41 39 37 34	robic MLSS	Average AER	DO Min	AER pH	AUR (NH4-N S	lope)	SAUR	NPR (NOx-N		
<	50 50 50 45 45 225 41 39 37 34			DO Min			lope)			nin]	SNPR
Actual SI	50 50 50 45 425 41 39 37 34 56 RT Aer	robic MLSS	Average AER	DO Min	AER pH nitless] 6.88	AUR (NH4-N S [mg/L/hr] 1.58	lope) [mg/	SAUR 'g MLVSS/hr]	NPR (NOx-N [mg/L/m 1.65	nin]	SNPR [mg/g MLVSS/hr]
Actual SI (days) 9.81	50 50 50 45 425 41 39 37 34 56 RT Aer	robic MLSS [mg/L] 3760 PUR (PO ₄ -P	Average AER [mg/L] 1.59	DO Min	AER pH	AUR (NH4-N S [mg/L/hr] 1.58	lope) [mg/	SAUR 'g MLVSS/hr] 0.59 larifier IDC PO	NPR (NOx-N [mg/L/m 1.65 4-P Max AN Relo	in]	SNPR [mg/g MLVSS/hr] 0.62 Avg. ANA NO ₃ -

AA4 NOx-N [mg/L] 1.53		AA6 NOx-N [mg/L] 0.27		AA1 PO ₄ -l Release Slo [mg/L/hr] 12.90	ре	P Re	Specific I elease R g MLVSS 6.96	ate	Releas	e Slope 'L/hr]	AA2 Specific Release [mg/g ML\]	Rate /SS/hr]	AA3 F Release [mg/l	e Slope L/hr]	P F	Specific PO ₄ - elease Rate g MLVSS/hr] 2.56
AA4 PO ₄ -P Release [mg/L/hr] -0.30	Slope	AA4 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr] -0.11		PO ₄ -P Release [mg/L/hr] -0.30	Slope	Re	Specific Poelease Rate /g MLVSS	е	AA6 PO ₄ -P Slope [mg/L/ 4.38	e hr]	AA6 Specific PO ₄ Release Rate [mg/g MLVSS/hi 1.64] [r	COD Uptake Slope mg/L/hr] 44.79	Úpt [mg/g	Specific CO take Rate MLVSS/hri 24.15	(AA1) COD
# Chem Used [lbs]	Fe	Cl3 Hrs Us [hr] 0.000	ed	Add [mg	CI3 lition g/L]	1	Bi Influ		olid Flow BD]		CY Flow [MGD] 19.38		CY Flo MGD] 1.087	ow	[N	Y Flow (GD) 9.5
ANA MLTSS [mg/L] 2440	5 A	NA % VOL [%] 76	MI [r	ANA LVSS mg/L]	AN)	X ML [mg/ 376	.TSS L]		NX % VC [%] 71		.NX MLVSS [mg/L] 2670	AER [m	MLTSS ng/L] 760	۷]	ER % 'OL %] 71	AER MLVSS [mg/L] 2670



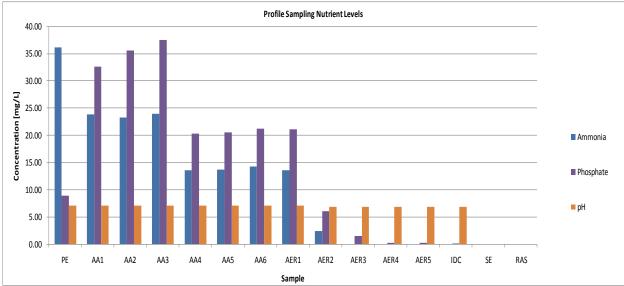


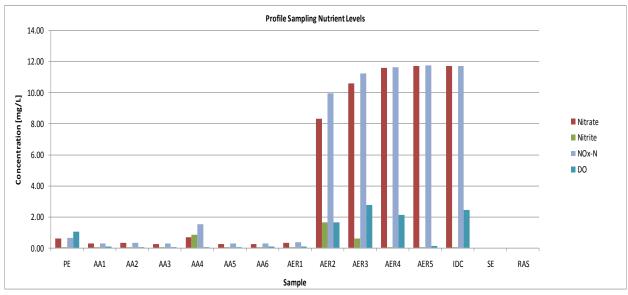


PROFILE 2:	Train 4, Ae	ration Tank 4, CI	arifier 4										
Date	Sample Tim	HRT along BNR Process (w/out e Recycles)	Sample Number	Reactor Type	Ammonia by HACH TNT	Nitrate by HACH TNT	Nitrite by HAC	NOx-N by Nitrate CH HACH + Nitrite HACH	Ph	osphate by IACH TNT	DO	pH	
dd-mmm-yy	h:mm	hr			mg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg/L NOx-N	m	g/L PO4-P	[mg/L	[unitless]	
6-Aug-09	7:15	0	PE		36.10	0.62	0.03	0.65		8.89	1.04	7.05	
6-Aug-09	7:24	0.97	AA1	Anaerobic	23.80	0.29	0.02	0.31		32.63	0.09	7.13	
6-Aug-09	7:58	1.93	AA2	Anaerobic	23.30	0.32	0.01	0.33		35.57	0.07	7.11	
6-Aug-09	8:32	2.90	AA3	Anaerobic	24.00	0.26	0.01	0.28		37.53	0.05	7.09	
6-Aug-09	9:06	3.86	AA4	Anoxic	13.60	0.68	0.87	1.55		20.31	0.06	7.06	
6-Aug-09	9:24	4.83	AA5	Anoxic	13.70	0.25	0.03	0.28		20.56	0.05	7.06	
6-Aug-09	9:42	5.79	AA6	Anoxic	14.25	0.27	0.03	0.30		21.21	0.09	7.06	
6-Aug-09	10:00	0.00	AER1	Aerobic	13.55	0.32	0.06	0.38		21.13	0.09	7.05	
6-Aug-09	10:32	1.78	AER2	Aerobic	2.49	8.31	1.66	9.97		6.10	1.64	6.86	
6-Aug-09	11:04	3.56	AER3	Aerobic	0.00	10.60	0.62	11.22		1.50	2.75		
6-Aug-09	11:36	5.34	AER4	Aerobic	0.00	11.60	0.05	11.65		0.29	2.14	6.88	
6-Aug-09	12:08	7.13	AER5	Aerobic	0.00	11.70	0.06	11.76		0.22	0.13	6.93	
6-Aug-09	12:11		IDC	-	0.00	11.70	0.02	11.72		0.16	2.46	6.93	
SAMPLE NOT CO	LLECTED		SE		0.00	0.00	0.00	0.00		0.00	0.00	0.00	
SAMPLE NOT CO	LLECTED		RAS		0.00	0.00	0.00	0.00		0.00	0.00	0.00	
*NOTE: These value		NILI NI	1010		0.00	0.00	0.00	0.00		0.00	0.00	0.00	
	- U												
QA/QC S	ampie:	NONE											
EPA	365.1	EP	A 350.1	E	PA 353.2	EF	PA 353.2	E	PA 3	53.2		EPA 353.2	
sCOD	by CEL	()-PO ₄		NH ₄ -N	N	IO _{2,3} -N]	NO ₃ .	-N		NO ₂ -N	
[m	g/L]	1	mg/L		mg/L		mg/L		mg/	L		mg/L	
2:	28		-		-		-		-			-	
7	7												
7	0												
7	7												
5	6												
3	88												
	7												
	60												
	10												
	88												
	34												
	52												
	5												
)												
)												
Actual SR	т	Aerobic MLSS	Average AEF	R DO Mi	n AER pH	AUR (NH4-N S	Slope)	SAUR	Ni	PR (NOx-N S	ilope)	SNPR	
	.1		-			,			+-"				rl
[days] 9.93		[mg/L] 4000	[mg/L] 1.35	- - 	unitless] 6.86	[mg/L/hr] 1.66	<u>l</u> [n	ng/g MLVSS/hr] 0.58	+	[mg/L/min 1.97		[mg/g MLVSS/hi 0.68	r]
Highest AEI	R NO ₂ -N	PUR (PO ₄ -P	1.33		0.00	1.00		Clarifier IDC PC	D ₄ -P	Max ANA	PO ₄ -P	Avg. ANA NO ₃	:-N
Concenti	ation	Uptake Slope)	SPUR	Cla	rifier IDC NH4	N Clarifier ID	C NOx-N	Concentration	on	Relea	se	Concentration	n
[mg/l	_]	[mg/L/hr]	[mg/g MLV	SS/hr]	[mg/L]	[mg]/L]	[mg/L]		[mg/	L]	[mg/L]	
1.66		3.77	1.31		0.0000	11.7		0.2		28.6		0.345	

AA4 NOx-N [mg/L] 1.55	AA6 NOx-N [mg/L] 0.30	N Rele	1 PO ₄ -P ase Slope ag/L/hr]	P Re [mg/g	pecific P lease Ra MLVSS/ 12.77	te Rele	2 PO ₄ - ase Slo ng/L/hr 9.32	оре	A2 Specifi Release [mg/g ML\ 6.42	Rate /SS/hr]	AA3 P Release [mg/L 6.3	Slope /hr]	AA3 Specific PO. P Release Rate [mg/g MLVSS/hr 4.38
AA4 PO ₄ -P Release Slope [mg/L/hr]	AA4 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr]	AA5 PO ₄ -P Release Slope [mg/L/hr]	AA5 Speci	ific PO ₄ -P e Rate	AA6 PC	O ₄ -P Release Slope mg/L/hr]	AA6 S Re	Specific PO lease Rate g MLVSS/h) ₄ -P (AA1)	COD Uptake Slope [mg/L/hr]	(AA1) S Upt	Specific COD ake Rate MLVSS/hr]	
-0.19	-0.06	0.05	0.0)2		1.07		0.37		67.19		46.33	0.28
# Chem Used F [lbs]	FeCl3 Hrs L [hr] 0.000	Jsed	FeCl3 Addition [mg/L 0.000	on .]	Bio Influe [N	ctive psolid ent Flow MGD]	/ A	RCY [MG 19.6	BD]		Y Flo GD] .08	ow N	IRCY Flow [MGD] 19.5
ANA MLTSS	ANA % VOL	ANA MLVS	SS AI	NX ML ⁻ [mg/L	TSS	ANX % \ [%]	OL.	ANX N	MLVSS g/L]	AER MI	TSS	AER ^c VOL [%]	% AER
1960	74	1450		4000		72			380	400		72	2880





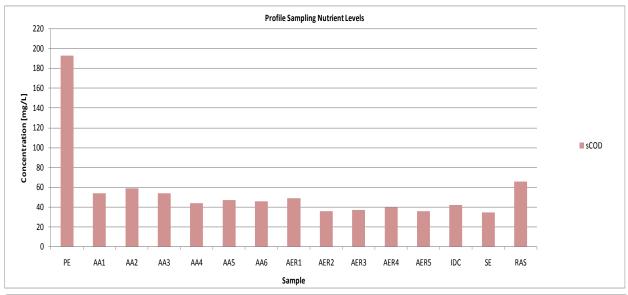


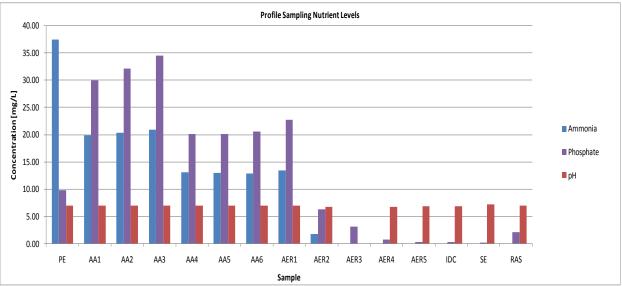
Week 4 – 8/10/09 & 8/12/09

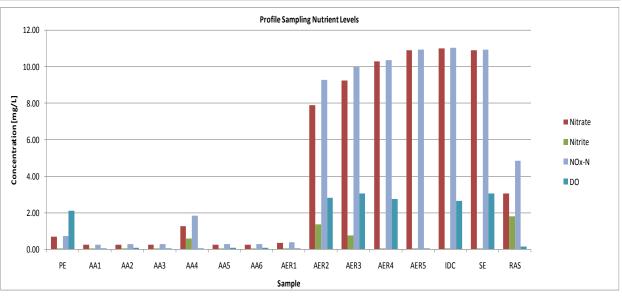
PROFILE 1:	Train 5, Aer	ation Tank 5, Cl	arifier 5									
Date	Sample Time	HRT along BNR Process (w/out		Reactor Type	Ammonia by HACH TNT	Nitrate by HACH TNT	Nitrite by HACH PP	NOx-N by Nitrate HACH + Nitrite HACH	Phosphate by HACH TNT	DO)	рН
dd-mmm-yy	h:mm	hr			mg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg/L NOx-N	mg/L PO4-P	[mg	/L]	[unitless]
10-Aug-09	6:54	0	PE		37.40	0.68	0.04	0.72	9.87	2.1	0	7.02
10-Aug-09	7:10	1.10	AA1	Anaerobic	19.90	0.24	0.01	0.25	29.94	0.0	15	7.04
10-Aug-09	7:40	2.19	AA2	Anaerobic	20.30	0.26	0.01	0.27	32.14	0.0	18	7.03
10-Aug-09	8:10	3.29	AA3	Anaerobic	20.90	0.26	0.01	0.27	34.51	0.0	15	7.02
10-Aug-09	8:40	4.38	AA4	Anoxic	13.05	1.25	0.60	1.85	20.07	0.0		7.01
10-Aug-09	8:55	5.48	AA5	Anoxic	13.00	0.26	0.02	0.28	20.15	0.0		6.99
10-Aug-09	9:10	6.57	AA6	Anoxic	12.85	0.26	0.02	0.29	20.56	0.0		6.98
10-Aug-09	9:25	0.00	AER1	Aerobic	13.45	0.33	0.05	0.38	22.68	0.0		6.97
10-Aug-09	9:55	1.70	AER2	Aerobic	1.82	7.90	1.36	9.26	6.27	2.8		6.78
10-Aug-09	10:25	3.39	AER3	Aerobic	0.00	9.24	0.76	10.00	3.18	3.0		679
10-Aug-09	10:55	5.09	AER4	Aerobic	0.00	10.30	0.05	10.35	0.77	2.7		6.78
10-Aug-09	11:25	6.79	AER5	Aerobic	0.00	10.90	0.03	10.93	0.33	0.0		6.83
10-Aug-09	11:34		IDC	-	0.00	11.00	0.03	11.03	0.30	2.6		6.87
10-Aug-09	14:34		SE	-	0.00	10.90	0.04	10.94	0.22	3.0		7.17
10-Aug-09	14:50		RAS	•	0.00	3.05	1.81	4.86	2.15	0.1	6	7.02
*NOTE: These valu	es are <1 mg/L	. NH ₄ -N		<u>-</u>								
QA/QC S	ample:	SE										
	-	ED	A 250 1	FI	24.052.0	ED		ED	252.0		EDA	252.2
EPA	365.1	EP.	A 350.1	E	PA 353.2	EP	A 353.2	EPA	A 353.2		EPA	353.2
	by CEL)-PO ₄		NH ₄ -N		O _{2,3} -N		O ₃ -N			0 ₂ -N
[mṛ	g/L]	1	mg/L		mg/L		mg/L	n	ng/L		m	g/L
19	93											
	54 59											
5	54											
4	4											
	7											
	ŀ6											
	9											
	36											
	37 37											
	10											
3	6											
	86											
4	12											
4	12 35		0.218		<0.20		11.4		11.4		0.	02
4	12		0.218		<0.20		11.4		11.4		0.	02
4	12 35		0.218		<0.20		11.4		11.4		0.	02
4 3 6	35 66			R DO M		ALIR (NIHA.N				Slone)	0.	
4 3 6 Actual SR	35 66	Aerobic MLSS	Average AEF		n AER pH	AUR (NH4-N 3	Slope)	SAUR	NPR (NOx-N			SNPR
4 3 6 Actual SR	35 66	erobic MLSS	Average AEF		n AER pH unitless]	[mg/L/hr	Slope)	SAUR g/g MLVSS/hr]	NPR (NOx-N [mg/L/mi			SNPR g MLVSS/hr]
4 3 6 Actual SR	12 135 166 T A	erobic MLSS [mg/L] 3700	Average AEF		n AER pH	,	Slope)	SAUR	NPR (NOx-N			SNPR
4 3 6 Actual SR	12 135 166 T A	erobic MLSS	Average AEF		n AER pH unitless]	[mg/L/hr	Slope)	SAUR g/g MLVSS/hr]	NPR (NOx-N [mg/L/mi			SNPR g MLVSS/hr]
Actual SR [days] 8.12	22 35 66 T A	erobic MLSS [mg/L] 3700	Average AEF [mg/L] 1.75		n AER pH unitless]	[mg/L/hr	Slope)	SAUR g/g MLVSS/hr]	NPR (NOx-N [mg/L/mi 1.81	n]	[mg/s	SNPR g MLVSS/hr]
Actual SR [days] 8.12	PR NO ₂	Lerobic MLSS [mg/L] 3700 UR (PO ₄ -P	Average AEF		n AER pH unitless]	[mg/L/hr 1.69 Clarifier IE	Slope) [my	SAUR g/g MLVSS/hr] 0.65	NPR (NOx-N [mg/L/mi 1.81 4- Max ANA	n] PO ₄ -	[mg/s	SNPR g MLVSS/hr] 0.70
Actual SR [days] 8.12	22 25 66 RT A	Img/L] 3700 UR (PO ₄ -P Uptake Slope)	Average AEF [mg/L] 1.75	Clarif	n AER pH unitless] 6.78	[mg/L/hr 1.69 Clarifier IE	Slope) J [mç DC NOx- Cl	SAUR g/g MLVSS/hr] 0.65	NPR (NOx-N [mg/L/mi 1.81 4- Max ANA	n] PO ₄ -	[mg/s	SNPR g MLVSS/hrj 0.70 ANA NO ₃ -N

AA4 NOx-I [mg/L] 1.85	N AA6 NOx-I [mg/L] 0.29	N	AA1 F Release [mg/l	e Slope L/hr]	AA1 Speci P Releas [mg/g ML1	e Rate /SS/hr]		PO ₄ -P Release Slope [mg/L/hr] 7.61	Rel	pecific PO ₄ -P ease Rate g MLVSS/hr] 9.30	AA3 PO ₄ -I Release Slo [mg/L/hr] 5.46	pe P Re	pecific PO ₄ - lease Rate MLVSS/hr] 6.67
AA4 PO ₄ -P Release Slope [mg/L/hr] 0.15	AA4 Specific PO ₄ -P Release Rate [mg/g MLVSS/hr] 0.06	Rele	.5 PO ₄ -P ase Slope ng/L/hr]	Relea: [mg/g N	cific PO ₄ -P se Rate //LVSS/hr]	AA6 F Release [mg/l	e Slope L/hr]	AA6 Specific F Release Ra [mg/g MLVS: 0.48	ate	(AA1) COD Uptake Slope [mg/L/hr] 59.83	Uptak [mg/g M	ecific COD e Rate LVSS/hr]	(AA1) PO₄- P/ (AA1) COD [mg/mg]
# Chem Used [lbs]	FeCl3 Hrs Used [hr]		Fe0 Addi [mg	CI3 tion /L]	Activ Influ		solid low		iD]	CRCY	′ Flow GD]	NRC	Y Flow GD] 9.5

		ANA					AER %	AER
ANA MLTSS	ANA % VOL	MLVSS	ANX MLTSS	ANX % VOL	ANX MLVSS	AER MLTSS	VOL	MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[mg/L]
1090	75	818	3700	70	2590	3700	70	2590

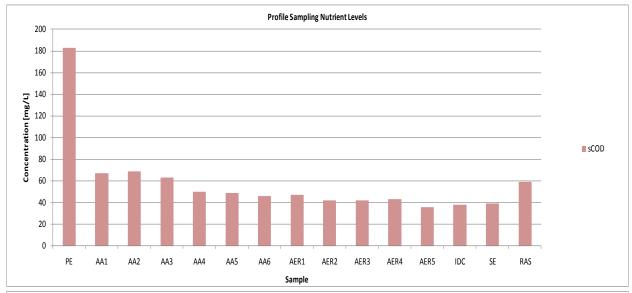


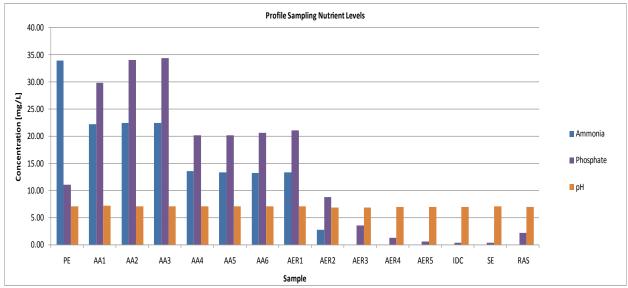


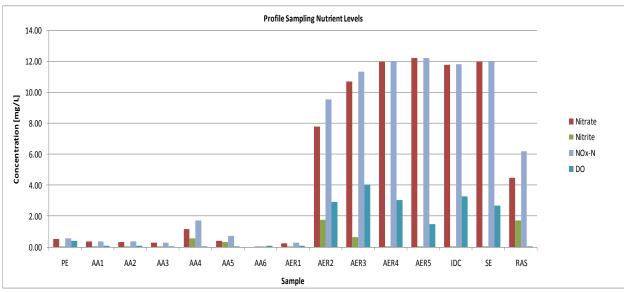


PROFILE 2:	Train 4, Aera	tion Tank 4, Cla	arifier 4									
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Type	Ammonia by HACH TNT	Nitrate by HACH TNT	Nitrite by HACH PP	NOx-N by Nitrate HACH + Nitrite HACH	Phosphate by HACH TNT	DO)	рН
dd-mmm-yy	h:mm	hr			mg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg/L NOx-N	mg/L PO4-P	[mg	/L]	[unitless]
12-Aug-09	7:52	0	PE	-	34.00	0.53	0.04	0.58	11.01	0.4		7.11
12-Aug-09	8:07	1.05	AA1	Anaerobic	22.20	0.36	0.02	0.38	29.86	0.0		7.17
12-Aug-09	8:37	2.10	AA2	Anaerobic	22.50	0.33	0.02	0.34	34.02	0.0		7.12
12-Aug-09	9:07	3.15	AA3	Anaerobic	22.40	0.28	0.03	0.30	34.43	0.0		7.11
12-Aug-09	9:37	4.21	AA4	Anoxic	13.55	1.18	0.56	1.74	20.15	0.0		7.05
12-Aug-09	9:52	5.26	AA5	Anoxic	13.35	0.42	0.31	0.73	20.15	0.0		7.04
12-Aug-09	10:07	6.31	AA6	Anoxic	13.20	0.00	0.02	0.02	20.64	0.0		7.06
12-Aug-09	10:22	0.00	AER1	Aerobic	13.30	0.25	0.02	0.27	21.05	0.1		7.05
12-Aug-09	10:58	1.70	AER2	Aerobic	2.70	7.79	1.76	9.55	8.78	2.9		6.86
12-Aug-09	11:22	3.39	AER3	Aerobic	0.00	10.70	0.64	11.34	3.59	4.0		6.88
12-Aug-09	11:52	5.09	AER4	Aerobic	0.00	12.00	0.04	12.04	1.25	3.0		6.91
12-Aug-09	12:22	6.79	AER5	Aerobic	0.00	12.20	0.03	12.23	0.63	1.4		6.91
12-Aug-09	12:27		IDC	•	0.00	11.80	0.02	11.82	0.38	3.2	6	6.92
12-Aug-09	15:37		SE	-	0.00	12.00	0.03	12.03	0.38	2.6	8	7.10
12-Aug-09	15:56		RAS		0.00	4.47	1.73	6.20	2.19	0.0	6	6.98
*NOTE: These valu	ies are <1 mg/L l	NH ₄ -N										
*NOTE: These value												
QA/QC S		IDC										
EPA	365.1	EP.	A 350.1	Е	PA 353.2	EP	A 353.2	EPA	A 353.2		EPA :	353.2
[m	by CEL g/L]		D-PO ₄	L	NH ₄ -N mg/L		O _{2,3} -N		O ₃ -N		NO ₂	
2	67 69 63 60 49 46 47 42 42											
	36											
]	38		101		0.20		10.1		10.1			2
		().484		<0.20		13.1		13.1		0.0)3
	39).484		<0.20		13.1		13.1		0.0)3
).484 I		<0.20		13.1		13.1		0.0)3
	39 59	erobic MLSS	0.484 Average AEI	R DO M	<0.20	AUR (NH4-N		SAUR	13.1 NPR (NOx-N S	slope)		03 SNPR
Actual SF	39 59		Average AEI				Slope)	SAUR	NPR (NOx-N S	' '		SNPR
Ę	39 59	erobic MLSS			in AER pH [unitless]	AUR (NH4-N	Slope)			' '		
Actual SF [days] 8.34 Highest AE N Concent	RT Ac	erobic MLSS [mg/L] 3640 JR (PO ₄ -P Uptake Slope)	Average AEI [mg/L] 2.31 SPUR	Clarif	in AER pH [unitless] 6.86	AUR (NH4-N : [mg/L/hr 1.73 Clarifier IE	Slope) [m-	SAUR g/g MLVSS/hr] 0.66 clarifier IDC PO Concentratio	NPR (NOx-N S [mg/L/min 2.19 4- Max ANA P Relea:	PO ₄ -	[mg/g Avg. A Conc	SNPR MLVSS/hr] 0.83 NA NO ₃ -N centration
Actual SF [days] 8.34	R NO ₂ -ration	erobic MLSS [mg/L] 3640 JR (PO ₄ -P Uptake Slope)	Average AEI [mg/L] 2.31	Clarif	in AER pH [unitless] 6.86	AUR (NH4-N : [mg/L/hr 1.73	Slope) [m. Coc NOx- Coc Full Code Figure 1	SAUR g/g MLVSS/hr] 0.66	NPR (NOx-N S [mg/L/min 2.19 4- Max ANA	PO ₄ -	[mg/g Avg. A Conc	SNPR MLVSS/hr] 0.83 NA NO ₃ -N

AA4 NOx-N	N AA6 NOx-N		AA1 PO ₄ -P Release Slope		A1 Specific Release R			O ₄ -P R Slope	Release			fic PO ₄ -P e Rate	AA3 P Release			pecific PO ₄ -P ease Rate
[mg/L]	[mg/L]		[mg/L/hr]	[n	mg/g MLVS	SS/hr]	1]	mg/L/h	r]	[mg/g	ML'	VSS/hr]	[mg/l	_/hr]	[mg/g	g MLVSS/hr]
1.74	0.02		13.45		9.12			8.69			5.9	0	5.2	25		3.56
AA4 PO ₄ -P Release Slope	AA4 Specific PO ₄ -P Release Rate	AA5 Po		Specific		AA6 PC			ecific Pease Rat			A1) COD ake Slope		Specifi otake R		(AA1) PO ₄ -P/ (AA1) COD
[mg/L/hr]	[mg/g MLVSS/hr]	[mg/L	/L/hr] [mg	/g MLV	/SS/hr]	[mg/L/l	hr]	[mg/g	MLVSS	/hr]	[n	ng/L/hr]	[mg/	g MLVS	SS/hr]	[mg/mg]
0.14	0.05	0.0	00	0.00)	1.03	1		0.39			46.59		31.60		0.29
# Chem Used	FeCl3 Hrs		FeCl3 Addition	n		ent F	low			Flov	v	CRC		w		CY Flow
[lbs]	[hr]		[mg/L]			MGD]		+	[MC		+		GD]		_	/IGD]
0	0.000		0.000 ANA		-1	9.86			19.0			11.	943	AEF		19.5 AER
ANA MLTS	S ANA % VO	_ M	//LVSS	ANX	MLTSS	AN.	X% ∨	/OL	ANXI	MLVS:	S	AER ML	TSS	V	OL	MLVSS
[mg/L] 1820	[%] 81	_	[mg/L] 1474		mg/L] 3640	-	[%] 72			ng/L] 621	4	[mg/l 3640			6] 2	[mg/L] 2621



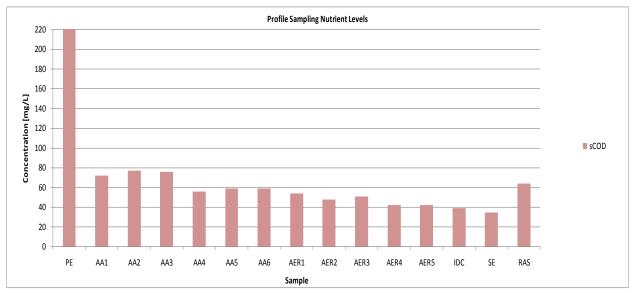


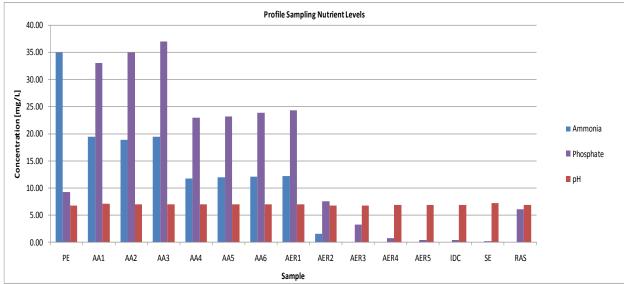


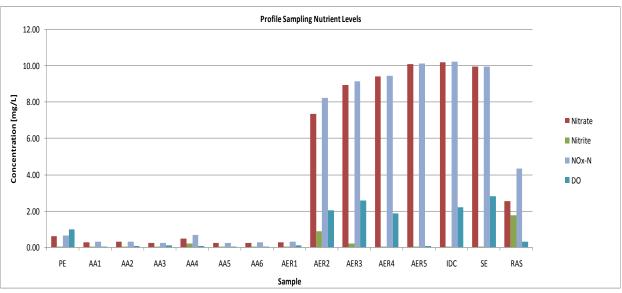
Week 5 – 8/17/09 & 8/20/09

PROFILE 1	Train 6, Ae	ration Tank 5, (Clarifier 5												
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Numbe	r Reactor Type	Ammon	iia by HACH	TNT Nitrate b	oy HACH TNT	Nitrite by HACH		N by Nitrate HA0 - Nitrite HACH	H Phosphate by	HACH TNT	DO	рН
dd-mmm-yy	h:mm	hr			m	ng/L NH3-N	mg	L NO3-N	mg/L NO2-i	N	mg/L NOx-N	mg/L F	04-P	[mg/L]	[unitless]
17-Aug-09	6:55	0	PE	-		34.90		0.63	0.03		0.66	9.3	10	0.98	6.72
17-Aug-09	7:14	1.10	AA1	Anaerobic		19.40		0.29	0.01		0.30	33.	04	0.06	7.05
17-Aug-09	7:44	2.20	AA2	Anaerobic		18.90		0.31	0.02		0.32	34.		0.07	7.03
17-Aug-09	8:14	3.31	AA3	Anaerobic		19.40		0.25	0.01		0.26	37.		0.11	7.00
17-Aug-09	8:44	4.41	AA4	Anoxic		11.70		0.47	0.21		0.68	22.		0.08	6.99
17-Aug-09	9:00	5.51	AA5	Anoxic		12.00		0.24	0.01		0.25	23.		0.06	6.98
17-Aug-09	9:15	6.61	AA6	Anoxic		12.10		0.25	0.02		0.27	23.		0.05	7.00
17-Aug-09	9:30	0.00	AER1	Aerobic		12.25		0.29	0.02		0.31	24.		0.10	6.97
17-Aug-09	10:00 10:30	0.85 1.71	AER2 AER3	Aerobic Aerobic		1.55 0.00		7.34 8.92	0.90 0.22		8.24 9.14	7.5		2.04	6.80
17-Aug-09 17-Aug-09	11:00	3.41	AER4	Aerobic		0.00		9.41	0.22		9.14	0.8		1.87	6.82
17-Aug-09	11:30	6.83	AER5	Aerobic		0.00		10.10	0.03		10.14	0.4		0.09	6.83
17-Aug-09	11:37	0.00	IDC	ACIODIC		0.00		10.20	0.04		10.14	0.3		2.20	6.92
-				-											
17-Aug-09	15:00		SE	-		0.00		9.94	0.02		9.96	0.2		2.80	7.16
17-Aug-09	15:20		RAS	-		0.00		2.55	1.78		4.33	6.1	3	0.30	6.93
	e values are <1	0													
QA/	QC S	ample	: 8	SE											
	EPA	365.1		EPA 3:	50.1	E	PA 353	3.2	EPA 35	53.2	EPA	353.2	EPA	A 35	3.2
S	[m 2 7	by CEI g/L] 26 72 77 76 56	L	O-PC			NH ₄ -N	1	NO _{2,3} ·			3-N g/L		O ₂ -l	
	£ 2	59 59 54 48 51 42													
		39													
		35		0.17	2	T	< 0.20		10.6	;	10	0.6		0.02	
		64													
Actual		robic MLSS	Average A		/lin AE		AUR (NH4			UR	,	Ox-N Slope		SNPF	
[days		[mg/L]	[mg		[unitle		[mg/		[mg/g M		<u>j [m</u>	g/L/min]	[mg/		SS/hr]
Conc	AER NO ₂ -Nentration	3700 N PUR (PO ₄ : Slo)	pe)	SPUR		Clarifier I	1.: DC NH ₄ -N	Clarifier II	DC NOx-N	Conce	DC PO ₄ -P entration	2.25 Max ANA PC Release [mg/L]		g. ANA	NO ₃ -N
	0.90	6.		2.20	2,1.11		0000		2250		9/ <u>-</u>]).4	27.74		0.30	
								•							

AA4 NOx-N	AA6 NOx-N	AA1 F	PO ₄ -P Slop	Release	AA1 Specific F		AA2 PO		AA2 Spec Releas			3 PO ₄ -P		Specific PO ₄ -P
[mg/L]	[mg/L]		[mg/L	/hrl	[mg/g MLVS	S/hr1	[mg/L/h	rl	[mg/g ML	VSS/hrl	ſ'n	ng/L/hr]	[ma/	g MLVSS/hr]
0.68	0.27		14.9	_	11.03		8.08	,	5.9		_	5.74	[9	4.23
AA4 PO ₄ -P Release Slope	AA4 Specific Release F			5 PO ₄ -P ase Slope	AA5 Specific PC Release Rate	F	AA6 PO ₄ -P Release Slope		Specific PO Release Rate	,	COD Slope	(AA1) Speci Uptake		(AA1) PO ₄ -P/ (AA1) COD
[mg/L/hr]	[mg/g MLVS	SS/hr]		g/L/hr]	[mg/g MLVSS/l	nr]	[mg/L/hr]	[m	g/g MLVSS/h	· · · ·	L/hr]	[mg/g MLV	_	
0.33	0.12		<u> </u>	0.04	0.02		1.15		0.42	64	.39	47.3	3	0.23
# Chem Used	FeCl3 H	rs Us	sed	FeCl3	3 Addition		tive Bios Ifluent Flo		ARCY	′ Flow	CR	CY Flow	NI	RCY Flow
[lbs]	[h	r]		[r	mg/L]		[MGD]		[Mo	GD]	[MGD]		[MGD]
0	0.0	000		(0.000		18.95		18.	75	1	1.602		19.5
ANA MLTS	S ANA %	VOL	ANA	MLVSS	ANX MLTS	S AN	NX % VOL	AN	XMLVSS	AER MI	_TSS	AER % V	OL /	AER MLVSS
[mg/L]	[%]		[n	ng/L]	[mg/L]		[%]		[mg/L]	[mg/	L]	[%]		[mg/L]
1720	79		1	359	3700		75		2775	370	0	75		2775

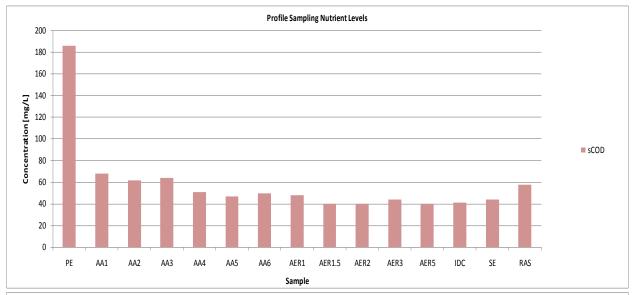


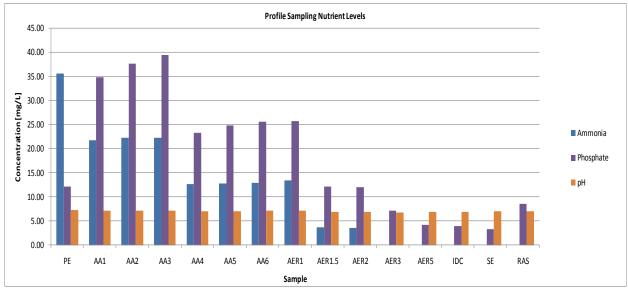


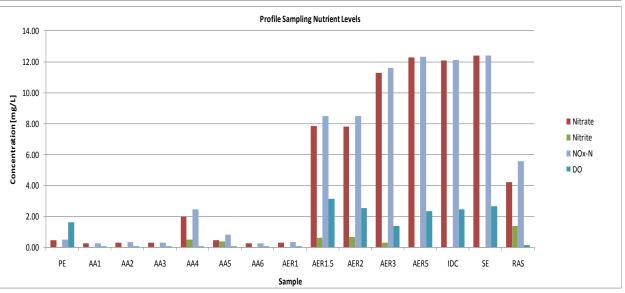


dd-mmm-yy 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	Sample Time h:mm 7:08 7:25 7:55	HRT along BNR Process (w/out Recycles) hr 0	Sample Number	Reactor Type	Ammo	nia by HACH	TNT Nitrate	LUACUTAT			N by Nitrate HAC				
dd-mmm-yy 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	h:mm 7:08 7:25	Recycles)	Sample Number	Reactor Type	Ammoi	nia hv HACH	TNT Nitrote								
dd-mmm-yy 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	h:mm 7:08 7:25	hr	Sample Number	Reactor Type	Ammo	nia hv HACH	TAIT Mitrote								
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	7:08 7:25					1110 07 111 1011	IIVI IVIIIdit	by HACH TNT	Nitrite by HAC		+ Nitrite HACH	Phosphate b	y HACH TNT	DO	pН
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	7:25	0			r	ng/L NH3-N	m	g/L NO3-N	mg/L NO2	-N	mg/L NOx-N	mg/L	PO4-P	[mg/L]	[unitless
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09		-	PE	-		35.60		0.45	0.04		0.49	12	2.07	1.62	7.22
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	7:55	1.11	AA1	Anaerobic		21.70		0.25	0.02		0.27	34	1.83	0.06	7.13
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09		2.22	AA2	Anaerobic		22.20		0.32	0.01		0.33		7.61	0.07	7.10
20-Aug-09 20-Aug-09 20-Aug-09 20-Aug-09	8:25	3.33	AA3	Anaerobic		22.30		0.30	0.01		0.32		9.40	0.06	7.09
20-Aug-09 20-Aug-09 20-Aug-09	8:55	4.44	AA4	Anoxic		12.60		1.97	0.49		2.46		3.33	0.08	7.03
20-Aug-09 20-Aug-09	9:10	5.56	AA5	Anoxic		12.80		0.47	0.37		0.84		1.80	0.08	7.06
20-Aug-09	9:25	6.67	AA6	Anoxic		12.95		0.26	0.01		0.28		5.53	0.05	7.07
	9:40	0.00	AER1	Aerobic		13.45		0.30	0.03		0.33		5.70	0.08	7.07
	9:55	0.85	AER1.5	Aerobic		3.65		7.86	0.64		8.50		2.07	3.15	6.89
20-Aug-09	10:10	1.71	AER2	Aerobic		3.60		7.80	0.68		8.48		2.04	2.53	6.88
20-Aug-09	10:40	3.41	AER3	Aerobic	100	0.00		11.30	0.32		11.62	_	.15	1.40	6.81
20-Aug-09	11:40	6.83	AER5	Aerobic		0.00		12.30	0.02		12.32		.19	2.36	6.85
20-Aug-09	11:45		IDC	-		0.00		12.10	0.02		12.12	3.	.95	2.48	6.87
20-Aug-09	14:46		SE	-		0.00		12.40	0.02		12.42	3.	.28	2.65	7.01
20-Aug-09	15:07		RAS	-		0.00		4.21	1.37		5.58	8.	.58	0.16	6.96
NOTE: These	e values are <1	ma/L NH ₄ -N													
		.23 mg/L NO ₃ -N													
		ample	. 6	E											
QA	عد عد	ampie	: 3												
	EPA	365.1]	EPA 3	50.1	E	PA 35	3.2	EPA 35	53.2	EPA	353.2	EPA	35	3.2
					_			_							
s	COD	by CE	L	O-PO) ₄]	NH ₄ -1	1	$NO_{2,3}$	-N	NO	3-N	NO	O_2 -1	7
	[m	g/L]		mg/	L		mg/L	,	mg/l	L	mg	;/L	m	g/L	,
	1	86													
		88													
		52													
		64													
		51													
		17													
		50													
		18	_												
		10 10													
		14													
		14 10													
		₽0 ₽1													
			_				0.20		10.4	<u> </u>	1.0			. 00	
		14					< 0.20	<u> </u>	13.6	,	13	.6	C	.02	
		8							_						
Actual Sl	DT Agr	obic MLSS	Average A	ER DO	Min AE	R pH	AUR (NH	4-N Slope	s/	AUR	NPR (N	Ox-N Slope	e)	SNPF	3
[days]		[mg/L]	[mg/		[unitle		•	/L/hr]		/ILVSS/hi	,	g/L/min]			SS/hr
7.67		3580	1.9		6.8			.50).57	. <u>, </u>	2.85	Įmg/s	1.08	
				<u> </u>		1									
•	AER NO ₂ -N	PUR (PO ₄	-P Uptake								DC PO ₄ -P	Max ANA P		•	NO ₃ -N
	entration	Slo	pe)	SPUR		Clarifier I	DC NH₄-N	Clarifier I	DC NOx-N	Conce	entration	Release	C	oncen	tration
Conce	ng/L]	[mg/	L/hr] [mg/g MLVS	SS/hr]	[m	ıg/L]	[m	ng/L]	[m	ıg/L]	[mg/L]		[mg/	[L]
	60	4.	68	1.77		0.0	0000	12.	1200	3	3.9	27.33		0.59	94
	.00			۵ معدمام	A1 Sn	ecific PC),-P A	 A2 P∩P	AA2 S	necific Pr)P AA	3 POP	AA3 Sn	ecific	POP
[m	1.00	Δ	A1 P∩P ₽		, UL	COMO I'C	-a F		1 / 1/74 0		-a	U 1 U4-1		JUILLO	. 04-г
[m			A1 PO ₄ -P R Slope	Cicase				-			-	•			ate
[m	x-N AA6	NOx-N A	A1 PO ₄ -P R Slope [mg/L/h		Rele	ease Rate	Re	lease Slop [mg/L/hr]	e Rel	ease Rat	e Rele	ase Slope		ase R	
[m	60	4.	•		A1 Sp	0.0 ecific PC		12. .A2 PO ₄ -P		pecific P		27.33 3 PO ₄ -P	AA3 Sp		
[m _j 0. AA4 NOx	x-N AA6	NOx-N	Slope		Rele	ase Rate	Re	lease Slop	e Rel	ease Rat	e Rele	ase Slope	Rele	ase R	

AA4 PO ₄ -P Release Slope	AA4 Specific PO ₄ -P Release Rate		5 PO ₄ -P ase Slope	AA5 Specific PC Release Rate		AA6 PO ₄ -P Release Slope		Specific PO	(, , , , ,) COD e Slope	(AA1) Specific		(AA1) PO ₄ -P/ (AA1) COD
[mg/L/hr]	[mg/g MLVSS/hr]	[m	g/L/hr]	[mg/g MLVSS/	hr]	[mg/L/hr]	[mg	/g MLVSS/h	r] [mg	/L/hr]	[mg/g MLVS	S/hr]	[mg/mg]
-0.18	-0.07	(0.26	0.10		1.17		0.44	45	5.59	31.76		0.32
# Chem Used	FeCl3 Hrs U	sed	FeCl3	3 Addition		ctive Bios Influent Flo		ARCY	′ Flow	CR	CY Flow	NR	CY Flow
[lbs]	[hr]		[r	mg/L]		[MGD]		[M0	GD]	[MGD]		[MGD]
0	0.000		(0.000		18.79		18.	43	1	1.501		19.5
ANA MLTSS	S ANA % VOL	ANA	MLVSS	ANX MLTS	S /	ANX % VOL	ANX	MLVSS	AER M	LTSS	AER % VO	DL A	ER MLVSS
[mg/L]	[%]	[n	ng/L]	[mg/L]		[%]	[r	ng/L]	[mg	/L]	[%]		[mg/L]
1840	78	1	435	3580		74	2	2649	358	30	74		2649





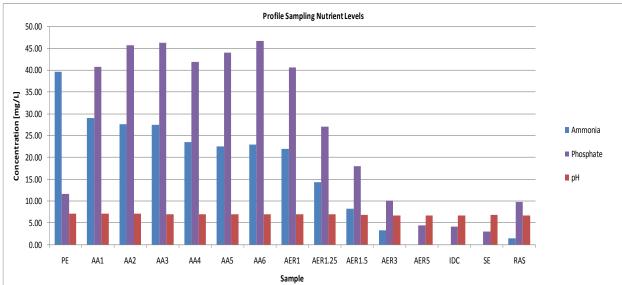


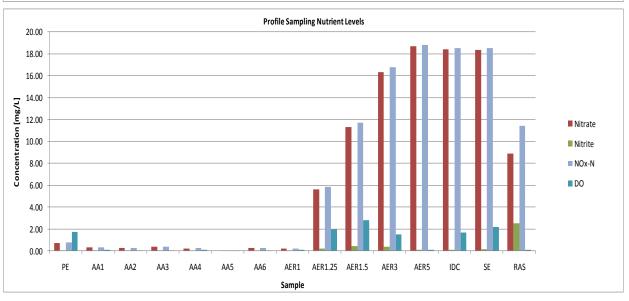
Week 6 – 8/24/09 & 8/26/09

ROFILE 1	Train 4, A	eration Tank 4	, Clarifier 4												
		HRT along BNF	3												
		Process (w/ou	t								by Nitrate HA				
Date	Sample Tim	e Recycles)	Sample Number	r Reactor Type	Ammoni	a by HACH TNT	Nitrate b	y HACH TNT	Nitrite by HAC		Nitrite HACH	Phosphate I	y HACH TNT	DO	рН
dd-mmm-yy	h:mm	hr			mç	g/L NH3-N		L NO3-N	mg/L NO2	N n	ng/L NOx-N		PO4-P	[mg/L]	[unitless]
24-Aug-09 24-Aug-09	7:18 7:33	1.10	PE AA1	- Anaerobic		39.60 29.00		0.73	0.05 0.01		0.79		.67 .79	1.72 0.09	7.09 7.06
24-Aug-09 24-Aug-09	8:03	2.20	AA1	Anaerobic		27.70		0.28	0.01		0.32		i.68	0.09	7.08
24-Aug-09	8:33	3.30	AA3	Anaerobic		27.50		0.39	0.02		0.41		.34	0.07	7.03
24-Aug-09	9:03	4.40	AA4	Anaerobic		23.60		0.23	0.02		0.25		.93	0.08	7.04
24-Aug-09	9:18	5.50	AA5 AA6	Anaerobic		22.50		0.00	0.01		0.01		.05 i.66	0.06	7.01
24-Aug-09 24-Aug-09	9:33 9:48	6.60 0.00	AA6 AER1	Anaerobic Aerobic		23.00		0.26	0.01 0.01		0.28		1.63	0.06	6.99 6.99
24-Aug-09	9:56	0.34	AER1.25	Aerobic		14.40		5.61	0.24		5.85		.08	1.94	6.99
24-Aug-09	10:04	0.68	AER1.5	Aerobic		8.25		11.30	0.44		11.74		.03	2.79	6.86
24-Aug-09	10:49	3.41	AER3	Aerobic	_	3.23		16.36	0.40		16.76		.15	1.53	6.73
24-Aug-09	11:49	6.82	AER5	Aerobic	_	0.00		18.70	0.13		18.83 18.51		47	0.11 1.67	6.72
24-Aug-09 24-Aug-09	11:54 14:54		SE	-	-	0.00		18.40 18.34	0.11 0.17		18.51		11 03	2.18	6.73
24-Aug-09	15:07		RAS		_	1.48		8.91	2.54		11.45		82	0.08	6.68
		<1 mg/L NH ₄ -N	1010			1.10		0.01	2.01		11.40	3	02	0.00	0.00
		<0.23 mg/L NO ₃ -N													
		ample		SE.											
		365.1	•	EPA 3	250.1	EDA	353	2 2	EPA 3:	52.2	EDA	353.2	EP/	25	22
	EFF	X 303.1		EPA 3	550.1	EFF	333	5.2	EPA 3.	33.2	EFA	333.2	EFF	1 33	3.2
s		by CE	EL	O-P	O ₄	N	H_4-N	1	$NO_{2,3}$		NC) ₃ -N	N	O_2 -I	N
		ng/L]	_	mg/	'L	n	ng/L		mg/	L	m	g/L	n	ng/L	,
	•	188													
		81 80	_												
		70													
		70													
		65													
		66													
		66 53													
		49													
		47													
		44													
		45				1									
		46		2.8	1	().48		19.4	ļ <u> </u>	19	9.2	(0.18	
		55													
Actual	SRT A	erobic MLSS	Average A	AER DO	Min AEI	R pH AU	R (NH4	-N Slope)	SA	AUR	NPR (N	Ox-N Slope	e)	SNPF	₹
[days	s]	[mg/L]	[mg	/L]	[unitle	ss]	[mg/l	L/hr]	[mg/g N	ILVSS/hr]	[m	ıg/L/min]	[mg/g	MLV	SS/hr]
7.24	ļ	3180	1.2	<u>.</u> 9	6.72	2	2.5	58	1	.07		3.95		1.63	
Highest	AER NO ₂	-N PUR (PC	₄ -P Uptake							Clarifier ID	OC PO ₄ -P	Max ANA P	O₄-P Av	a. ANA	NO ₃ -N
_	entration	1	ope)	SPUF	2	Clarifier IDC	NH₄-N	Clarifier I	DC NOx-N	Concer	ntration	Release		oncent	tration
ſr	ng/L]	ſmo	g/L/hr]	[mg/g MLV		[mg/L		ſm	g/L]	[mg	ı/L1	[mg/L]		[mg/	L1
	0.44	_	5.87	2.84		0.0000			5100	4.		35.00		0.24	_
					1										
		L.	AA1 PO₄-P I	Release	141 Sno	ecific PO ₄ -F	Δ /	\2 PO₄-P	ΔΔ2 Q	pecific PO)P	\3 PO₄-P	AA3 Sp	ecific	P∩D
AA4 NC) _{x-N} Δ Δ	6 NOx-N	Slope		•	ase Rate		ease Slop		ease Rate		ease Slope		ase R	
, , , , ¬ INC	~ 11 / /							Jaco Olop	- INGR	Jaco Nait			1,010	a00 11	4.0
Ima/I	1	[ma/L1	[/] /4	orl .		M //CC/L-1	г.	ma/l /hrl	[ma/a	MIVER	orl f.	ma/l/hrl	[ma/~	MI VO	C/h-l
[mg/l 0.25		[mg/L] 0.28	[mg/L/l 10.94	_		MLVSS/hr] 8.44	[r	mg/L/hr] 11.05	[mg/g	MLVSS/h 8.53	nr] [ı	mg/L/hr] 6.81	[mg/g	MLVS 5.26	S/hr]

AA4 PO ₄ -P Release Slope	AA4 Specific PO ₄ -P Release Rate		5 PO ₄ -P ase Slope	AA5 Specific PC Release Rate		AA6 PO ₄ -P Release Slope		Specific PO ₄ elease Rate	(, , , , ,) COD e Slope	(AA1) Specific		(AA1) PO ₄ -P/ (AA1) COD
[mg/L/hr]	[mg/g MLVSS/hr]	[m	ıg/L/hr]	[mg/g MLVSS/	hr]	[mg/L/hr]	[mg/	/g MLVSS/hr] [mg	/L/hr]	[mg/g MLVS	S/hr]	[mg/mg]
0.98	0.40		0.39	0.16		2.11		0.87	43	3.11	33.28		0.25
# Chem Used [lbs]	FeCl3 Hrs U: [hr]	sed	[r	3 Addition mg/L]		active Bios Influent Flo [MGD]		ARCY [MC	BD]	[1	CY Flow MGD]		CY Flow [MGD]
0	0.000		(0.000		18.98		18.	11	1	1.603		0
ANA MLTSS	S ANA % VOL	ANA	MLVSS	ANX MLTS:	5 /	ANX % VOL	ANX	MLVSS	AER M	LTSS	AER % VO	DL A	ER MLVSS
[mg/L]	[%]	[n	ng/L]	[mg/L]		[%]	[r	ng/L]	[mg	/L]	[%]		[mg/L]
1640	79	1	1296	3180		76	2	2417	318	80	76		2417

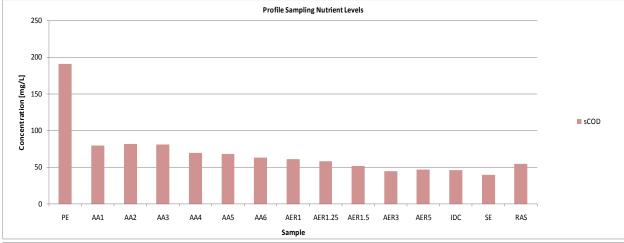


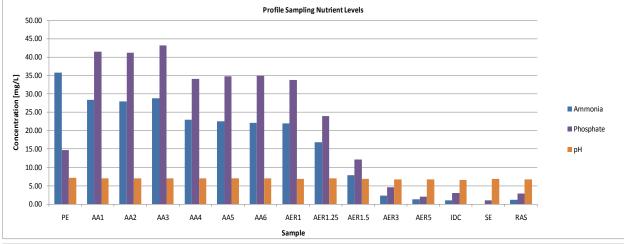


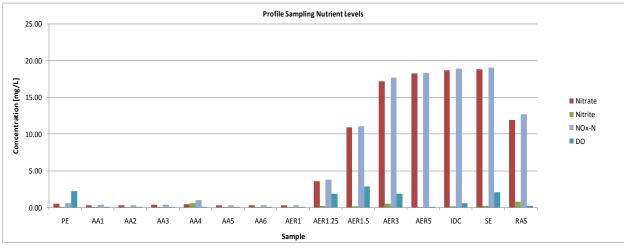


	l rain 6, Ae	ration Tank 5,	Clarifier 5												
		HRT along BNR													
Į.		Process (w/out								NOx	k-N by Nitrate H	IACH			
Date	Sample Time	Recycles)	Sample Numb	er Reactor Type	Ammonia	a by HACH TNT	Nitrate b	y HACH TNT	Nitrite by HAC	H PP	+ Nitrite HACH	l Phosphat	e by HACH TI	NT DO	pН
dd-mmm-yy	h:mm	hr			m(y/L NH3-N	mg	L NO3-N	mg/L NO2	-N	mg/L NOx-N	mç	/L PO4-P	[mg/L]	[unitless]
26-Aug-09	7:10	0	PE	-		35.70		0.51	0.06		0.57		14.77	2.23	7.20
26-Aug-09	7:16	1.13	AA1	Anaerobic		28.40		0.32	0.02		0.34		41.44	0.06	7.04
26-Aug-09	7:46	2.25	AA2	Anaerobic		28.00		0.30	0.01		0.32		41.12	0.06	7.04
26-Aug-09	8:16	3.38	AA3	Anaerobic		28.80		0.34	0.02		0.36		43.24	0.09	7.05
26-Aug-09	8:46	4.50	AA4	Anaerobic		23.00		0.44	0.60		1.04		34.10	0.10	6.97
26-Aug-09	9:01	5.63	AA5	Anaerobic		22.60		0.28	0.02		0.30		34.75	0.05	6.96
26-Aug-09	9:16	6.76	AA6	Anaerobic		22.10		0.27	0.02		0.29		34.92	0.05	6.95
26-Aug-09	9:31	0.00	AER1	Aerobic		21.90		0.27	0.02		0.29		33.77	0.10	6.94
26-Aug-09	9:39	0.34	AER1.25	Aerobic		16.90		3.59	0.20		3.79		23.98	1.84	6.95
26-Aug-09	9:47	0.68	AER1.5	Aerobic		7.89		10.90	0.18		11.08		12.16	2.86	6.88
26-Aug-09	10:32	3.41	AER3	Aerobic		2.36		17.20	0.52		17.72		4.57	1.87	6.68
26-Aug-09	11:36	6.82	AER5	Aerobic	and the same of th	1.31		18.26	0.11		18.37		2.06	0.06	6.67
26-Aug-09	11:42		IDC	-		1.04		18.70	0.18		18.88		2.97	0.62	6.65
26-Aug-09	14:30		SE	-		0.00		18.86	0.22		19.08		0.98	2.10	6.90
26-Aug-09	14:48		RAS	-		1.19		11.90	0.82		12.72		2.84	0.26	6.68
_	se values are <	1 mg/L NHN													
		-													
		0.23 mg/L NO ₃ -N				1		1			1				
QA/		ample	: ;	SE											
	EPA	365.1		EPA 3	50.1	EPA	353	3.2	EPA 3	53.2	EPA	353.2	EP	A 35	53.2
s	COD	by CE	ᆫ	O-P	\mathcal{O}_4	N	H4-N	1	$NO_{2,3}$	-N	N	O_3-N	1	VO 2-	Ν
		ıg/L]		mg/	L	n	ng/L		mg/	L	n	ng/L		mg/I	_
		91	_												
		80 82													
		81													
		70													
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	(68 63													
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	(((68 63 61 58 52													
		68 63 61 58 52 45													
	() () ()	68 63 61 58 52													
	(!	68 63 61 58 52 45					0.20		20.4	1	2	20.2		0.24	
	() () () () () () () () () ()	68 63 61 58 52 45 47					0.20		20.4	1	2	20.2		0.24	
	() () () () () () () () () ()	68 63 61 58 52 45 47 46				<	0.20		20.4	ı		20.2		0.24	
	4	68 63 61 58 52 45 47 46 40	Average	AER DO	Min AEI			N Slove							
Actual S	SRT Ae	68 63 61 58 52 45 47 46 40 55				R pH AU	R (NH4) S/	AUR	NPR	(NOx-N Slo		SNPI	R
[days	SRT Ae	68 63 61 58 52 45 47 46 40 55	[mç]/L]	[unitle	R pH AU	R (NH4	L/hr]) S/	AUR 1LVSS/h	NPR	(NOx-N Slo		SNPI g/g MLV	R /SS/hr]
	SRT Ae	68 63 61 58 52 45 47 46 40 55]/L]		R pH AU	R (NH4	L/hr]) S/	AUR	NPR	(NOx-N Slo		SNPI	R /SS/hr]
[days	4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4 4	68 63 61 58 52 45 47 46 40 55 robic MLSS	[mg]/L]	[unitle	R pH AU	R (NH4	L/hr]) S/	AUR ILVSS/h .05	NPR	(NOx-N Slo mg/L/min] 4.50	[mg	SNPI g/g MLV 1.85	R /SS/hr]
[days 7.34 Highest	SSRT Ae:	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280	[mo 1.: -P Uptake	y/L] 35	[unitle	R pH AU	R (NH4 [mg/l 2.5	L/hr] 55	S/ [mg/g M	AUR ILVSS/r .05 Clarifier	NPR [IDC PO ₄ -P	(NOx-N Slo mg/L/min] 4.50	PO ₄ -P /	SNPI g/g MLV 1.85	R /SS/hr] 5
[days 7.34 Highest A	SSRT Ae:	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280	[mo 1. -P Uptake ppe)	9/L] 35 SPUR	[unitle	R pH AU	R (NH4 [mg/l 2.5 NH ₄ -N	L/hr] 55 Clarifier II	S/ [mg/g M 1	AUR ILVSS/F .05 Clarifier Conc	NPR IDC PO ₄ -Pentration	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea	PO ₄ -P	SNPI g/g MLV 1.85 Avg. ANA Concen	R /SS/hr] 5 A NO ₃ -N tration
[days 7.34 Highest A Conce	SSRT Ae.	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Sic	[mg 1.: -P Uptake ope) [L/hr]	9/L] 35 SPUR [mg/g MLV\$	[unitle	R pH AU	R (NH4 [mg/l 2.5 NH ₄ -N	∟/hr] 55 Clarifier II [m	S/ [mg/g N 1 DC NOx-N ng/L]	AUR ILVSS/F .05 Clarifier Conc	NPR [IDC PO ₄ -Pentration mg/L]	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/	PO ₄ -P A	SNPI n/g MLV 1.85 Avg. ANA Concen [mg	R /SS/hr] 6 A NO ₃ -N tration /L]
[days 7.34 Highest A Conce	SSRT Ae:	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Sic	[mo 1. -P Uptake ppe)	9/L] 35 SPUR	[unitle	R pH AU	R (NH4 [mg/l 2.5 NH ₄ -N	∟/hr] 55 Clarifier II [m	S/ [mg/g M 1	AUR ILVSS/F .05 Clarifier Conc	NPR IDC PO ₄ -Pentration	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea	PO ₄ -P A	SNPI g/g MLV 1.85 Avg. ANA Concen	R /SS/hr] 6 A NO ₃ -N tration /L]
[days 7.34 Highest A Conce	SSRT Ae.	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slc	[mg 1.: -P Uptake ope) [L/hr]	y/L] 35 SPUR [mg/g MLV3 2.85	[unitle 6.67 SS/hr]	R pH AU	R (NH4 [mg/l 2.8 NH ₄ -N	∟/hr] 55 Clarifier II [m) S, [mg/g N 1 DC NOx-N ng/L] 8800	AUR ILVSS/F .05 Clarifier Conc	NPR IDC PO ₄ -P centration mg/L] 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/	PO ₄ -P // se	SNPI n/g MLV 1.85 Avg. ANA Concen [mg	R (SS/hr] A NO ₃ -N tration (L]
[days 7.34 Highest A Conce	SSRT Ae: SI AER NO ₂ -1 entration mg/L] 0.52	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slc	[mç 1. -P Uptake ppe) [L/hr] 92	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr]	R pH AU ss] 7 Clarifier IDC [mg/L] 1.0400	R (NH4 [mg/l 2.5 NH₄-N]	L/hr] 55 Clarifier II [m 18.) S, [mg/g M 1 DC NOx-N ng/L] 8800 AA2 S	AUR MLVSS/h .05 Clarifier Conc	NPR IDC PO ₄ -P Pentration mg/L 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4	PO ₄ -P Ase	SNPI n/g MLV 1.85 Avg. ANA Concen [mg 0.33	R //SS/hr] 6 A NO ₃ -N tration /L] 26 PO ₄ -P
[days 7.34 Highest According to 10 Missing t	SSRT Ae:	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slo	[mg. 1P Uptake ope) [L/hr] 92 A1 PO ₄ -P Slope	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr]	Clarifier IDC [mg/L 1.0400 ecific PO ₄ -Fase Rate	R (NH4 [mg/l 2.5 NH ₄ -N]) P AA Rele	Clarifier II [m 18. A2 PO ₄ -P ease Slop) S/ [mg/g M 1 DC NOx-N ng/L] 8800 AA2 S Reli	AUR MLVSS/r .05 Clarifier Conc [r	NPR IDC PO ₄ -P Reterration mg/L] 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4 AA3 PO ₄ -P	PO ₄ -P Ase L] AA3 S	SNPI 3/g MLV 1.85 Avg. ANA Concen [mg 0.3: Specific	R (SS/hr] A NO ₃ -N tration /L] 226 PO ₄ -P cate
[days 7.34 Highest A Conce [n	SSRT Ae. SI AER NO ₂ -1 entration mg/L] 0.52 Dx-N AAI	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 PUR (PO ₄ Slc	[mç 1. -P Uptake ppe) [L/hr] 92 A1 PO ₄ -P	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr] A1 Spe Relea [mg/g N	R pH AU ss] 7 Clarifier IDC [mg/L] 1.0400	R (NH4 [mg/l 2.5 NH ₄ -N]) P AA Rele	Clarifier II) S/ [mg/g M 1 DC NOx-N ng/L] 8800 AA2 S Reli	AUR MLVSS/h .05 Clarifier Conc [r	NPR IDC PO ₄ -P Reterration mg/L] 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4	PO ₄ -P Ase L] AA3 S	SNPI 3/g MLV 1.85 Avg. ANA Concen [mg 0.33	R (SS/hr] A NO ₃ -N tration /L] 226 PO ₄ -P cate
[days 7.34 Highest . Conco	SSRT Ae. SI AER NO ₂ -1 entration mg/L] 0.52 Dx-N AAI	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slo [mg/L] 6.	[mg/L/ 1P Uptake ppe) [L/hr] 92 A1 PO ₄ -P Slope [mg/L/ 14.8	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr] AA1 Spee Relea [mg/g N	Clarifier IDC [mg/L] 1.0400 ecific PO ₄ -Fase Rate MLVSS/hr]	R (NH4 [mg/l 2.8 NH ₄ -N) P AA Rele	_/hr] 55 Clarifier II) S/ [mg/g M 1 DC NOx-N ng/L] 8800 AA2 S Reli	AUR MLVSS/h .05 Clarifier Conc [r pecific Pease Ra MLVSS	NPR IDC PO ₄ -P Reterration mg/L] 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4 AA3 PO ₄ -P elease Slope [mg/L/hr]	PO ₄ -P Ase L] AA3 S	SNPI 3/g MLV 1.85 Avg. ANA Concen [mg 0.3: Specific elease R g MLVS 4.06	R (SS/hr] 6 A NO ₃ -N tration /L] 26 PO ₄ -P Rate SS/hr]
[days 7.34 Highest A Conce [n	SSRT Ae. SI AER NO ₂ -1 entration mg/L] 0.52 Dx-N AAI	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slo [mg/L] 6 NOx-N	[mg/L/ 1P Uptake ppe) [L/hr] 92 A1 PO ₄ -P Slope [mg/L/ 14.8	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr] AA1 Spee Relea [mg/g N	Clarifier IDC [mg/L 1.0400 ecific PO ₄ -F ase Rate	R (NH4 [mg/l 2.5 NH ₄ -N]) P AA Rele	_/hr] 55 Clarifier II) S/ [mg/g M 1 DC NOx-N ng/L] 8800 AA2 S Reli	AUR ILVSS/h .05 Clarifier Conc [r pecific Pease Ra MLVSS 5.45	NPR IDC PO ₄ -P Reterration mg/L] 3.0	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4 AA3 PO ₄ -P elease Slope [mg/L/hr] 6.46	PO ₄ -P Ase L] AA3 S	SNPI 3/g MLV 1.85 Avg. ANA Concen [mg 0.3: Specific elease R g MLVS 4.06	R (SS/hr] A NO ₃ -N tration /L] 226 PO ₄ -P cate
[days 7.34 Highest . Conco	SSRT Ae. SI AER NO ₂ -1 entration mg/L] 0.52 DX-N AAI L] [1]	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slo [mg/L] 6.	[mg/L/ -P Uptake ppe)	SPUR [mg/g MLV3 2.85 Release A	[unitle 6.67 SS/hr] AA1 Spee Relea [mg/g N	Clarifier IDC [mg/L] 1.0400 ecific PO ₄ -F se Rate MLVSS/hr] 0.36	R (NH4 [mg/l 2.8 NH ₄ -N) P AA Rele	Clarifier II [m] 18. A2 PO ₄ -P ease Slop mg/L/hr] 8.68) S/ [mg/g N 1 DC NOx-N ng/L] 8800 AA2 S Rele [mg/g	AUR ILVSS/r .05 Clarifier Conc [r Decific Pease Ra MLVSS 5.45 PO ₄ -P	NPR IDC PO ₄ -P entration mg/L] 3.0 PO ₄ -P Re Re S/hr]	(NOx-N Slo mg/L/min] 4.50 Max ANA Relea [mg/ 28.4 AA3 PO ₄ -P elease Slope [mg/L/hr] 6.46	PO ₄ -P // se // AA3 Se Re	SNPI 3/g MLV 1.85 Avg. ANA Concen [mg. 0.3: Specific elease R g MLVS 4.06	R (/SS/hr] (SS/hr] (NO ₃ -N) (tration (L) (26 (PO ₄ -P) (ate (SS/hr]
[days 7.34] Highest Conce	SSRT Ae: SI AER NO ₂ -1 entration mg/L] 0.52 Dx-N AAI L] II Slope	68 63 61 58 52 45 47 46 40 55 robic MLSS [mg/L] 3280 N PUR (PO ₄ Slc [mg/L] 6 NOx-N	[mg/L/ 1.4-P Uptake ppe) (L/hr] 92 A1 PO ₄ -P Slope [mg/L/ 14.8: A45 Relea	SPUR [mg/g MLV3 2.85 Release A hr] PO ₄ -P se Slope	[unitle 6.67 SS/hr] AA1 Spe Relea [mg/g N	Clarifier IDC [mg/L] 1.0400 ecific PO ₄ -P ase Rate ific PO ₄ -P e Rate	R (NH4 [mg// 2.6 NH ₄ -N]) P AA Rele [II	Clarifier II [m	DC NOx-N ng/L] 8800 AA2 Si ne Rele [mg/g	AUR MLVSS/h .05 Clarifier Conc [r pecific P ease Ras MLVSS 5.45 PO ₄ -P clate	NPR IDC PO ₄ -P entration mg/L] 3.0 PO ₄ -P Re S/hr] (AA1) COI	(NOx-N Slot mg/L/min] 4.50 Max ANA Relea [mg/L/s0] 28.4 AA3 PO ₄ -P elease Slope [mg/L/hr] 6.46 Decomposition (AA1) Specific properties of the content of	PO ₄ -P // se L] 7 AA3 Se Re [mg/	SNPI a/g MLV 1.85 Avg. ANA Concen [mg 0.32 Specific elease R g MLVS 4.06 (AA1) (AA4)	R (SS/hr] 6 A NO ₃ -N tration /L] 26 PO ₄ -P cate SS/hr]

# Chem Used	FeCl3 Hrs U	sed	FeCl3	Addition	Active Bios Influent Fl	-	ARCY	′ Flow	CRO	CY Flow	NRCY Flow
[lbs]	[hr]		[n	ng/L]	[MGD]		[M	GD]	[MGD]	[MGD]
0	0.000		0	.000	18.54		17.	92	1	1.241	0
ANA MLTS	S ANA % VOL	ANA	MLVSS	ANX MLTS	S ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[n	ng/L]	[mg/L]	[%]	[n	ng/L]	[mg/	[]	[%]	[mg/L]
2040	78	1	591	3280	74	2	427	328	0	74	2427



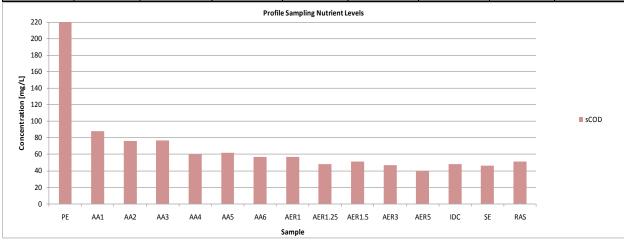


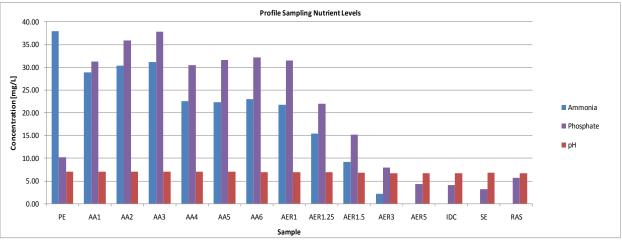


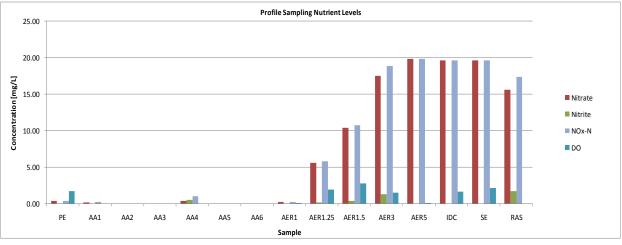
Week 7 – 9/1/09 & 9/2/09

PROFILE 1 T	Train 4, Ae	ration Tank 4	, Clarifier 4													
		HRT along BNF	,													
		Process (w/ou								NO	Ox-N by Nitrat	te HACH	1			
Date S	Sample Time	Recycles)	Sample Number	er Reactor Type	Ammon	ia by HACH TN	Nitrat	e by HACH TN	Nitrite by HAC	H PP	+ Nitrite HA	ACH	Phosphate b	y HACH TN	L DO	pН
dd-mmm-yy	h:mm	hr			m	g/L NH3-N	r	ng/L NO3-N	mg/L NO2	-N	mg/L NOx	(-N	mg/L	PO4-P	[mg/L]	[unitless]
1-Sep-09	8:02	0	PE	-		38.00		0.39	0.02		0.41			.20	1.72	7.09
1-Sep-09	8:19	0.88	AA1	Anaerobic		28.90		0.23	0.02		0.25			.33	0.09	7.06
1-Sep-09 1-Sep-09	8:49 9:19	1.77 2.65	AA2 AA3	Anaerobic Anaerobic		30.40 31.20		0.00 0.00	0.01 0.01		0.01			.89 .85	0.06	7.08 7.03
1-Sep-09	9:49	3.53	AA4	Anaerobic		22.55		0.45	0.58		1.03			.51	0.07	7.04
1-Sep-09	10:04	4.41	AA5	Anaerobic		22.35		0.00	0.02		0.02			.65	0.06	7.01
1-Sep-09	10:19	5.30	AA6	Anaerobic		23.05		0.00	0.02		0.02			.14	0.06	6.99
1-Sep-09	10:34	0.00	AER1	Aerobic		21.80		0.24	0.02		0.26		31	.49	0.10	6.99
1-Sep-09	10:42	0.34	AER1.25	Aerobic		15.40		5.65	0.22		5.87			.94	1.94	6.99
1-Sep-09	10:50	0.68	AER1.5	Aerobic		9.19		10.40	0.40		10.80			.26	2.79	6.86
1-Sep-09	11:35	3.42	AER3	Aerobic		2.21		17.56	1.32		18.88			99	1.53	6.73
1-Sep-09	12:35	6.84	AER5	Aerobic	-	0.00		19.82	0.02		19.84		4.		0.11	6.72
1-Sep-09	12:40		IDC		_	0.00		19.64	0.02		19.66			08	1.67	6.73
1-Sep-09	15:41		SE	-	_	0.00		19.64	0.02		19.66			20	2.18	6.87
1-Sep-09	15:57		RAS	-		0.00		15.60	1.78		17.38		5.	74	0.08	6.68
*NOTE: These																
		0.23 mg/L NO ₃ -N														
QA/C		ample) <u>:</u>	SE									1			
	EPA	365.1		EPA 3	350.1	EP	A 35	53.2	EPA 3	53.2	EF	PA	353.2	EP.	A 35	3.2
				O D	_			N.T.	NIO	ъ.т	Ι,		NT	_		
S		by CE	-	O-P mg			H ₄ - ng/l		NO _{2,3}		1	mg	3-N		1O ₂ -1 ng/L	
		19/ L] 313		mg.	<i>,</i> L	1	ng/i		mg/	<u>L</u>		mg	<i>/</i> L		ng/ L	,
		88														
		76 77														
		60														
		62														
		57 57														
		48														
		51														
		47 40	_													
		48														
		46				-	<0.2	0	20.	8		20	.8		0.02	
	;	51														
Actual SI	DT Ag	robic MLSS	Average /	AER DO	Min AE	R pH AL	IR (NH	4-N Slope	2) 5	AUR	NP	R (NO	Dx-N Slope	2)		
		[mg/L]	[mg		[unitle		,			-		`			SNPF	₹
[days] 5.21	J	2800	1.2		6.7		[m/		·	/II \/ Q Q	/hrl	[mo	/l /minl	·	SNPF	
5.21	I	2000		20 [g/L/hr]	[mg/g N		/hr]		/L/min]	·	g MLV	SS/hr]
			<u> </u>	29	0.7				[mg/g N	/ILVSS/ .32	/hr]		/L/min] 4.65	·		SS/hr]
1			•	29 [0.7			g/L/hr]	[mg/g N	.32			4.65	[mg.	g MLV 2.28	SS/hr]
Highest A		1	₄ -P Uptake	·				g/L/hr] 2.70	[mg/g N	.32 Clarifie	er IDC PO	4-P I	4.65 Max ANA P	[mg.	/g MLV 2.28 vg. ANA	SS/hr]
-	NER NO ₂ -l	1	•	29 SPUF			2	g/L/hr] 2.70	[mg/g N	.32 Clarifie		4-P I	4.65	[mg.	g MLV 2.28	SS/hr]
Conce		SI	₄ -P Uptake	·	₹	2	2 : NH ₄ -N	2/L/hr] 2.70 Clarifier	[mg/g N	.32 Clarifie Con	er IDC PO	4-P I	4.65 Max ANA P	[mg.	/g MLV 2.28 vg. ANA	SS/hr] NO ₃ -N
Concer [mg	entration	SI [mg	₄ -P Uptake ope)	SPUF	₹	2 Clarifier IDC	2 : NH ₄ -N -]	g/L/hr] 2.70 U Clarifier	[mg/g N	.32 Clarifie Con	er IDC PO	4-P I	4.65 Max ANA Po Release	[mg.	vg. ANA	SS/hr] NO ₃ -N ration L]
Concer [mg	entration ig/L]	SI [mg	₄ -P Uptake ope) ₁ /L/hr]	SPUF [mg/g MLV	₹	2 Clarifier IDC [mg/l	2 : NH ₄ -N -]	g/L/hr] 2.70 U Clarifier	[mg/g Notes In the second seco	.32 Clarifie Con	er IDC PO. ncentration [mg/L]	4-P I	4.65 Max ANA Po Release [mg/L]	[mg.	g MLV 2.28 vg. ANA Concent	SS/hr] NO ₃ -N ration L]
Concer [mg	entration ig/L]	SI [mg	₄ -P Uptake ope) J/L/hr] .44	SPUF [mg/g MLV 2.66	SS/hr]	Clarifier IDC	: NH ₄ -N -] 0	g/L/hr] 2.70 Clarifier [i	[mg/g N 1	.32 Clarifie Con	er IDC PO, ncentration [mg/L] 4.1	4-P I	4.65 Max ANA Port Release [mg/L] 27.66	[mg.	/g MLV 2.28 vg. ANA Concent [mg/	NO ₃ -N ration
Concer [mg	entration ig/L] .32	SI [mg	4-P Uptake ope) y/L/hr] .44 AA1 PO ₄ -P	SPUF [mg/g MLV 2.66	SS/hr]	Clarifier IDC [mg/l 0.000]	; NH ₄ -N -] 0	g/L/hr] 2.70 Clarifier 19 AA2 PO ₄ -F	[mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	.32 Clarifie Con	er IDC PO, ncentration [mg/L] 4.1 PO ₄ -P	AA:	Max ANA Pering Release [mg/L] 27.66	[mg,	vg. ANA Concent [mg/ 0.11	NO ₃ -N rration
Concer [m, 1.	entration g/L] .32 x-N AA	SI [mg	4-P Uptake ope) y/L/hr] .44 AA1 PO ₄ -P Slope	SPUF [mg/g MLV 2.66 Release	SS/hr] AA1 Spc Relea	Clarifier IDC [mg/l 0.000	; NH ₄ -N -] 0	AA2 PO ₄ -Felease Slo	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel	Clarifie Con	er IDC PO, neentration [mg/L] 4.1 PO ₄ -P ate	AA:	Max ANA Port Release [mg/L] 27.66 3 PO ₄ -P ase Slope	[mg,	vg. ANA Concent [mg/ 0.11	NO ₃ -N ration L] 3
Concer [mg/L] AA4 NOx	entration ag/L] .32 x-N AAG] [SI	4-P Uptake ope) y/L/hr] .44 AA1 PO ₄ -P Slope [mg/L/	SPUF [mg/g MLV 2.66 Release /	SS/hr] AA1 Spo Relea	Clarifier IDC [mg/l 0.000 ecific PO ₄ - ase Rate MLVSS/hr	; NH ₄ -N -] 0	7/L/hr] 2.70 Clarifier [i 15 AA2 PO ₄ -F elease Slo [mg/L/hr]	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel	Clarifie Con	er IDC PO, neentration [mg/L] 4.1 PO ₄ -P ate	AA:	Max ANA Port Release [mg/L] 27.66 3 PO ₄ -P ase Slope g/L/hr]	[mg,	g MLV 2.28 vg. ANA Concent [mg/ 0.11	NO ₃ -N ration L] 3
Concer [m, 1.	entration ag/L] .32 x-N AAG] [SI [mg	4-P Uptake ope) y/L/hr] .44 AA1 PO ₄ -P Slope	SPUF [mg/g MLV 2.66 Release /	SS/hr] AA1 Spo Relea	Clarifier IDC [mg/l 0.000	; NH ₄ -N -] 0	AA2 PO ₄ -Felease Slo	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel	Clarifie Con	er IDC PO, neentration [mg/L] 4.1 PO ₄ -P ate	AA:	Max ANA Port Release [mg/L] 27.66 3 PO ₄ -P ase Slope	[mg,	vg. ANA Concent [mg/ 0.11	NO ₃ -N ration L] 3
Concer [mg 1. AA4 NOx [mg/L] 1.03	entration g/L] .32 x-N AAG] [SI [mg/5]	AA1 PO ₄ -P Slope	SPUF [mg/g MLV 2.66 Release /	SS/hr] AA1 Spo Relea [mg/g I	Clarifier IDC [mg/l 0.000 ecific PO ₄ - ase Rate MLVSS/hr	2 NH ₄ -N -] 0 P Re	7/L/hr] 2.70 Clarifier [i 15 AA2 PO ₄ -Felease Slo [mg/L/hr] 10.96	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel [mg/g	Clarifie Con pecific lease R. MLVS 9.84	er IDC PO, ncentration [mg/L] 4.1 PO ₄ -P ate	AA:	Max ANA Pe Release [mg/L] 27.66 3 PO ₄ -P ase Slope g/L/hr] 7.16	AA3 S Rel [mg/g	vg. ANA Concent [mg/ 0.11 pecific ease R MLVS 6.43	NO ₃ -N ration L] 3 PO ₄ -P ate S/hr]
Concer [mg 1. AA4 NOx [mg/L] 1.03	entration g/L] .32 x-N AAA] [SI [mg 5 5 6 NOx-N mg/L] 0.02 1.4 Specific P	AA1 PO ₄ -P Slope [mg/L/11.8*	SPUF [mg/g MLV 2.66 Release /	SS/hr] AA1 Spo Relea [mg/g I	Clarifier IDC [mg/l 0.000 ecific PO ₄ - ase Rate MLVSS/hr 10.61	2 NH ₄ -N -] 0	7/L/hr] 2.70 Clarifier [I 15 AA2 PO ₄ -F elease Slo [mg/L/hr] 10.96	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel [mg/g	Clarifie Con pecific lease Rig MLVS 9.84 PO ₄ -P	er IDC PO, ncentration [mg/L] 4.1 PO ₄ -P ate (AA1) C	AAA:Relea	Max ANA Per Release [mg/L] 27.66 3 PO ₄ -P ase Slope g/L/hr] 7.16	AA3 S Rel	vg. ANA Concent [mgy 0.11 pecific ease R MLVS 6.43 (AA1)	NO ₃ -N ration L] 3 PO ₄ -P ate S/hr]
AA4 NOx [mg/L] 1.03 AA4 PO. Release S	entration 19/L] 32 x-N AAA [] [6 NOx-N mg/L] 0.02 4 Specific P Release Ra	AA1 PO ₄ -P Slope [mg/L/ 11.8* O ₄ -P AA5 e Relea:	SPUF [mg/g MLV 2.66 Release /	SS/hr] AA1 Spore Release Rele	Clarifier IDC [mg/l 0.000 ecific PO ₄ - ase Rate MLVSS/hr 10.61	NH ₄ -N NH ₄ -N Relea	AA2 PO ₄ -Felease Slo [mg/L/hr] 10.96	IDC NOx-N mg/L] 0.6550 P AA2 S pe Rel [mg/g AA6 Specific Release F	Clarifie Con pecific lease Rig MLVS 9.84 PO ₄ -P Rate	er IDC PO, ncentration [mg/L] 4.1 PO ₄ -P ate SS/hr] (AA1) C Uptake S	AAA Releas	Max ANA Per Release [mg/L] 27.66 3 PO ₄ -P ase Slope g/L/hr] 7.16 (AA1) Spec Uptake	AA3 S Rel [mg/c	vg. ANA Concent [mg/ 0.11 pecific ease R MLVS 6.43 (AA1) (AA1)	NO ₃ -N ration L] 3 PO ₄ -P ate S/hr]
Concer [mg 1. AA4 NOx [mg/L] 1.03	entration eg/L] .32 x-N AAA [] [SI [mg 5 5 6 NOx-N mg/L] 0.02 1.4 Specific P	AA1 PO ₄ -P Slope [mg/L/ 11.8* O ₄ -P AA5 e Relea:	SPUF [mg/g MLV 2.66 Release /	AA1 Spo Relea [mg/g I	Clarifier IDC [mg/l 0.000 ecific PO ₄ - ase Rate MLVSS/hr 10.61	22 S NH ₄ -N NH ₄ -N NH ₄ -N AA6 Relea	7/L/hr] 2.70 Clarifier [I 15 AA2 PO ₄ -F elease Slo [mg/L/hr] 10.96	IDC NOx-N mg/L] 0.6550 AA2 S pe Rel [mg/g	Clarifie Con pecific lease Rig MLVS 9.84 PO ₄ -P Rate	er IDC PO, ncentration [mg/L] 4.1 PO ₄ -P ate (AA1) C	AAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAAA	Max ANA Per Release [mg/L] 27.66 3 PO ₄ -P ase Slope g/L/hr] 7.16	AA3 S Rel [mg/g	vg. ANA Concent [mgy 0.11 pecific ease R MLVS 6.43 (AA1) (AA1)	NO ₃ -N ration L] 3 PO ₄ -P ate S/hr]

# Chem Used	FeCl3 Hrs U	sed FeCl3	3 Addition	Active Bios		ARCY	∕ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]	[r	ng/L]	[MGD]		[MGD]		[MGD]		[MGD]
0	0.000	C	0.000	18.92		18		11.482		0
ANA MLTS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]] [mg		[%]	[mg/L]
1410	79	79 1114		73	2	044	280	0	73	2044

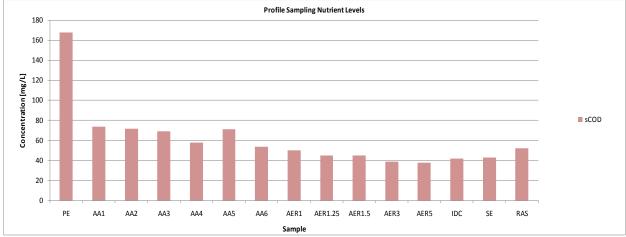


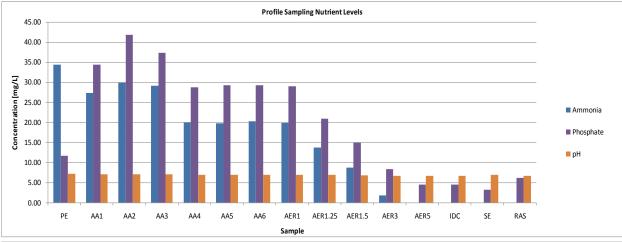


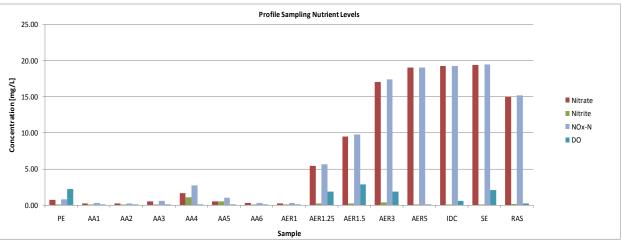


ROFILE 2	Train 6	, Aera	ition Tank 5	, Clarifie	r 5																
			HRT along BNF																		
Date	Sample 1		Process (w/out Recycles)		e Numbe	mber Reactor Type		Ammonia by HACH TNT		Nitrate by HACH TNT		T Niti	Nitrite by HACH PP		NOx-N by Nitrate HACH + Nitrite HACH			Phosphate by HACH TNT		рН	
dd-mmm-yy	h:mm	1	hr					mg/L NH3-N		mg/L NO3-N			mg/L NO2-N n		mg/L N	NOx-N mg/L P		L PO4-P	[mg/L		
2-Sep-09	7:10	_	0	-	PE -			34.40			0.74		0.07		0.8			11.67	2.23	7.20	
2-Sep-09	7:25		0.89				:	27.30		0.26			0.02		0.28		34.43		0.06	7.04	
2-Sep-09	7:55	_	1.77	_	\A2	Anaerobio Anaerobio		29.90			0.24		0.01		0.2			11.77	0.06	7.04	
2-Sep-09	8:25		2.65	_	AA3	Anaerobio	_	29.10		0.54			0.01		0.56		37.36		0.09	7.05	
2-Sep-09	8:55		3.53	А	AA4	Anaerobio	:	20.05		1.68			1.05	2.73		3	28.72		0.10	6.97	
2-Sep-09	9:10		4.41	A	AA5	Anaerobio	:	19.80	9.80		0.54		0.49		1.0			29.21	0.05	6.96	
2-Sep-09	9:25		5.30	A	AA6	Anaerobio	:	20.30	20.30		0.28		0.01		0.29		2	29.21	0.05	6.95	
2-Sep-09	9:40		0.00	Al	ER1	Aerobic		19.85			0.26		0.02	0.02		0.28		29.04	0.10	6.94	
2-Sep-09	9:48		0.34	AEI	R1.25	Aerobic		13.75	13.75		5.45		0.20		5.6	5.65 2		20.97	1.84	6.95	
2-Sep-09	9:56		0.68	AE	R1.5	Aerobic		8.77			9.51		0.24	24		.75 15		15.01	2.86	6.88	
2-Sep-09	10:41		3.42	Al	ER3	Aerobic	1.84				17.08		0.34		17.4	2		8.42	1.87	6.68	
2-Sep-09	11:41		6.84	Al	ER5	Aerobic		0.00			19.02		0.03		19.0	15		4.50	0.06	6.67	
2-Sep-09	11:47	,		I	DC	-	0.00				19.24		0.01		19.2	!5		4.50	0.62	6.65	
2-Sep-09	14:47	,			SE	-	0.00		19.44			0.01		19.45			3.20		6.90		
2-Sep-09	14:59)		R	RAS	-			0.00		15.02		0.16		15.1	8		6.20	0.26	6.68	
*NOTE: Thes		_	ma/L NH -N			0.															
			23 mg/L NO ₃ -N																		
			mple		- C	SE		i			1										
QA			_	• :				_													
	EF	PA	365.1		_	EPA	350.1]	EP <i>A</i>	353	3.2	EF	PA 35	53.2	E	PA	353.2	EF	A 35	3.2	
		_				0.1	20		NTI	T 18	т.		10	ът		NIO	N.T.	Ι,	10	. т	
- 5			by CE	<u> </u>	_	O-PO ₄				NH ₄ -N			NO _{2,3} -N		_		3-N	1	NO ₂ -N		
	[g/L]		_	mg	/L	ㅗ	n	ıg/L			mg/I	_		mg	g/L	<u> </u>	mg/I	_	
			86		-1																
			4																		
			2																		
69																					
58 71																					
					-																
	54 50																				
	45																				
			-5																		
		3	9																		
		3	8																		
		4	2																		
		4	3			2.78 <				0.20			20.5			20).5		0.02		
		5	2																		
												T									
				1																	
Actual	SRT	Aero	bic MLSS	Aver	rage A	AER DO Min AER pH AL				R (NH4-N Slope)		e)	SAUR		NPR (NC		IOx-N Slop	0x-N Slope)		SNPR	
[days			[ma	ng/L] [unitless]			[mg/L/hr]			+	[mg/g MLVSS/hr		S/hr] [mg/L/min]		g/l /minl	[mg/g MLVSS/hr		/SS/hrl			
5.38						1.35 6.67						+	1.22		,	4.28		1,,,	2.12		
0.00					1.0		0.0		1	۷.۰	i I		'i				0	-	۷. ۱۵		
Highest	AER NO	O ₂ -N	PUR (PC	4-P Upt	ake									Clarifie	er IDC P	O₄-P	Max ANA F	PO ₄ -P	Avg. AN	A NO ₃ -N	
Conc	entratio	n	,	ope)		SPU	R	Clarifi	er IDC	NH₄-N	Clarifier	IDC	NOx-N		ncentrati		Releas		Concer	-	
	. ,					0.0															
	[mg/L] [mg/L/hr]								[mg/L]			mg/L			[mg/L]			[mg/L]		[mg/L]	
	0.34		1 4	.82		2.39	1		0.0000	,	19	9.250	IU		4.5		30.10		0.5	90	
				1A1 PC)₄-P F	Release	AA1 Sr	ecific	PO F	ΔΔ	A2 PO₄-F		AA2 Sp	ecific	PO ₄ -P	Δ.	\3 PO₄-P	ДДЗ	Specific	PO ₄ -P	
												,		·		ase Slope			pecific PO ₄ -P		
				Slope	·					·						•		Release Rate			
			ng/L/ł			g/g MLVSS/hr]		[mg/L/hr]			[mg/g MLVS				ng/L/hr]	[mg	[mg/g MLVSS/hr]				
2.73	3	0	.29		15.94				14.37		13.39		12.06		06		5.84		5.26		
4445	<u> </u>		0	ا ي م	^ ^ -	₂ -	A A F O		<u>, </u>	^^^	, [0	DO 5	l ., .		[_	4) DC	
AA4 PO			Specific P			A5 PO ₄ -P AA5 Specific								(/			(AA1) Specific COD		1) PO ₄ -		
Release	Slope	F	Release Rat	ie I	Releas	ease Slope Releas		se Rate R		Release	Release Slope		Release Rate		Uptake Slope		Uptak	Uptake Rate		COD	
[mg/L	/hr]	[mo	g/g MLVSS	/hr]	[ma	ng/L/hr] [mg/g M		MLVSS/hr]		[ma/	[mg/L/hr] [[mg/g MLVSS/hr]		[mg/L/hr]		[mg/g M	[mg/g MLVSS/hr]		g/mg]	
-0.3	_		-0.18			.11		0.06		2.0			1.00	-		12		.86		0.37	
	-										1						- 00				

# Chem Used	FeCl3 Hrs U	sed FeCl3	Addition	Active Bios Influent Fl		ARCY	′ Flow	CRO	CY Flow	NRCY Flow	
[lbs]	[hr]	[r	ng/L]	[MGD]		[M	GD]	[MGD]		[MGD]	
0	0.000	C	.000	18.74		17.82		11.361		0	
ANA MLTSS	ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	TSS	AER % VC	L AER MLVSS	
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	_] [mg,		[%]	[mg/L]	
1370	81	1110	2800	72	2	016	2800		72	2016	



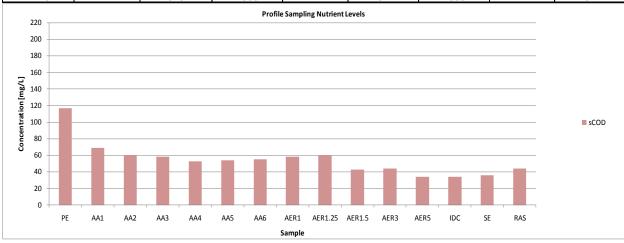


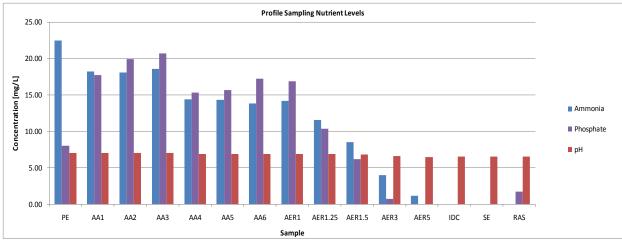


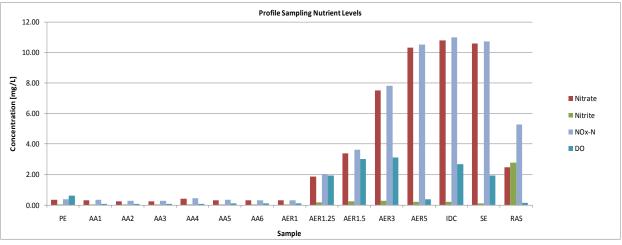
Week 8 – 9/9/09

'KOFILE 1	-	=	AL 101 -															
1	rain 6, A	eration Tank 5	, Clarifier 5															
		HRT along BNI																
		Process (w/ou			_ .							NOx-N by				te by HAC		
Date	Sample Time		Sample Nur	nber Reactor	Type Am	nmonia by HAC		Nitrate by H		Nitrite by I			rite HAC			T844	DO	pН
dd-mmm-yy	h:mm	hr				mg/L NH3-N	١	mg/L No		mg/L N			L NOx-N			P04-P	[mg/L]	[unitless]
9-Sep-09	8:10	0	PE	- Annore	hia	22.50		0.36		0.0			0.38			3.02	0.60	7.04
9-Sep-09 9-Sep-09	8:24 8:44	0.68 1.36	AA1 AA2	Anaero Anaero		18.20 18.10		0.33		0.0			0.35 0.26			7.70 9.90	0.08	7.03 7.03
9-Sep-09	9:19	2.04	AA3	Anaero		18.60		0.24		0.0			0.26			0.70	0.07	7.04
9-Sep-09	9:45	2.72	AA4	Anaero	bic	14.40		0.4		0.0			0.43			5.30	0.08	6.91
9-Sep-09	9:58	3.40	AA5	Anaero		14.35		0.32		0.0			0.34			5.70	0.11	6.89
9-Sep-09	10:15	4.08	AA6	Anaero		13.85		0.30		0.0			0.32			7.20	0.11	6.88
9-Sep-09 9-Sep-09	10:28 10:36	0.00 0.26	AER1 AER1.2	Aerob Aerob		14.20 11.55		0.29		0.0			0.31 2.03			6.90 0.40	0.11 1.95	6.90 6.90
9-Sep-09	10:45	0.53	AER1.5			8.53		3.40		0.2			3.63			5.21	3.01	6.86
9-Sep-09	11:30	2.63	AER3	Aerob	ic	4.02		7.52		0.2	29		7.81		C).73	3.10	6.61
9-Sep-09	12:30	5.27	AER5	Aerob	ic	1.21		10.3		0.2			10.51).00	0.39	6.47
9-Sep-09	12:38		IDC	-		0.00		10.7		0.2			10.99			0.00	2.69	6.52
9-Sep-09	15:26		SE	-		0.00		10.6		0.1			10.71).00	1.93	6.56
9-Sep-09	15:45		RAS	-		0.00		2.49		2.7	'9		5.28		1	.74	0.13	6.52
		<1 mg/L NH ₄ -N																
		<0.23 mg/L NO ₃ -N																
		<0.5 mg/L PO ₄ -P																
QA/		ample	=	SE					+			_						
	EPA	365.1		EPA	350.	1 EF	PA	353.2	E	PA 3	53.2	E	PA	353.	.2	EP	A 35	53.2
	COD	 05		O-I	20.		NILI	. NI	,	NIO	NI		NO	3-N			10-	NT.
s		by CE	_	mg		_	mg	4-N	-	NO _{2,3}		-	mg				1O ₂ -1 mg/L	
		17		1112	,/ L _		IIIg	<u>,, r</u>		IIIg/	_		1112	5/ L _			mg/ L	_
		69																
		60 58																
		53																
		54																
		54 55																
	:	54 55 58 60																
	:	54 55 58 60 43																
		54 55 58 60																
		54 55 58 60 43																
		54 55 58 60 43 44 34 34 34		0.0	97		<	20		11.7	7		11	.6			0.12	
		54 55 58 60 43 44 34		0.0	97		<.:	20		11.7	7	I	11	6			0.12	
		54 55 58 60 43 44 34 34 34		0.0	97		<	20		11.7	7	<u> </u>	11	6		<u> </u>	0.12	
		54 55 58 60 43 44 34 34 36								11.	7	<u> </u>	11	6		T	0.12	
Actual S		54 55 58 60 43 44 34 34 34	Average			ER pH			Tope)		7 AUR	T NI		. 6 Ох-N (Slope		0.12 SNPF	
Actual S	SRT Ae	54 55 58 60 43 44 34 34 36	 		Min A	ER pH /	AUR		ope)	S			PR (N			_		₹
	SRT Ae	54 55 58 60 43 44 34 34 34 34 36	[m	AER DO	Min A		AUR	(NH4-N S	ope)	S. [mg/g N	AUR		PR (N	Ox-N \$		_	SNPF	R 'SS/hr]
[days	SRT Ae	54 55 58 60 43 44 34 34 34 34 36 44	[m	AER DO	Min A	tless]	AUR	(NH4-N S [mg/L/hr]	ope)	S. [mg/g N	AUR //LVSS/		PR (N	Ox-N (_	SNPF g/g MLV	R 'SS/hr]
[days 4.35	SRT Ae	54 55 58 60 43 44 34 34 34 34 36 44 robic MLSS [mg/L]	[m	AER DO	Min A	tless]	AUR	(NH4-N S [mg/L/hr]	ope)	S. [mg/g N	AUR //LVSS/ .15		PR (N	Ox-N (n]	[mg	SNPF g/g MLV	R (SS/hr]
[days 4.35	GRT Ae	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600	[m	AER DO	Min A [unit	tless]	AUR	(NH4-N S [mg/L/hr] 2.22		S. [mg/g N	AUR //LVSS/ .15 Clarifie	hr]	PR (N [mṛ	Ox-N \$ g/L/mir 2.58 Max Al	n]	[mg	SNPI g/g MLV 1.34	SS/hr]
[days 4.35	SRT Ae	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600	[m 1 -P Uptake	AER DO g/L] .71	Min A [unit	.47 Clarifier I	AUR	(NH4-N S [mg/L/hr] 2.22		S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con	hr]	PR (N [mṛ	Ox-N S g/L/mir 2.58 Max Al Re	n] NA PC	[mg	SNPF g/g MLV 1.34 Avg. ANA	R (SS/hr] A NO ₃ -N tration
[days 4.35 Highest A Conce	SRT Ae J AER NO ₂ - entration	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600	[m 1 -P Uptake ope)	AER DO g/L] 71 SPL	Min A [unit	.47 Clarifier I	AUR	(NH4-N S [mg/L/hr] 2.22	ifier IDC	S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con	r IDC PC	PR (N [mṛ	Ox-N S g/L/mir 2.58 Max Al Re	n] NA PC elease	[mg	SNPF g/g MLV 1.34 Avg. ANA Concen	R (SS/hr] NO ₃ -N tration
[days 4.35 Highest A Conce	SRT Ae J AER NO ₂ -entration ng/L	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600	[m 1 -P Uptake ope) /L/hr]	AER DO g/L] 71 SPU [mg/g ML'	Min A [unit	.47 Clarifier I	AUR	(NH4-N S [mg/L/hr] 2.22	ifier IDC	S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con	r IDC PC centratio	PR (N [mṛ	Ox-N S g/L/mir 2.58 Max Al Re	n] NA PC elease mg/L]	[mg	SNPF g/g MLV 1.34 Avg. ANA Concen [mg,	R [SS/hr] INO ₃ -N tration [L]
[days 4.35 Highest A Conce	SRT Ae J AER NO ₂ -entration ng/L	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600 N PUR (PO. Sic	[m 1 -P Uptake ope) /L/hr]	AER DO g/L] .71 SPL [mg/g ML] 2.5	Min A [unit 6. JR VSS/hr]	.47 Clarifier I	AUR DC NH	(NH4-N S [mg/L/hr] 2.22	ifier IDC [mg/ 10.98	S. [mg/g M 1	AUR //LVSS/ .15 Clarifie Con	r IDC PC centration [mg/L]	PR (N [mg	Ox-N S g/L/mir 2.58 Max Al Re	NA PCelease	[mg	SNPF g/g MLV 1.34 Avg. ANA Concen [mg,	R (SS/hr] L NO ₃ -N tration (L) 09
[days 4.35 Highest A Conce	SRT Ae] AER NO ₂ -entration ng/L]	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600 N PUR (PO. Sic	[m 1 -P Uptake ope) /L/hr] 91	AER DO g/L] .71 SPU [mg/g ML] 2.5	Min A [unit 6. JR VSS/hr] 5	Clarifier IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	AUR DC Ni g/L] 0000	(NH4-N S [mg/L/hr] 2.22 	ifier IDC [mg/ 10.98	S. [mg/g M 1	AUR //LVSS/ .15 Clarifie Con	r IDC PC centratic mg/L] 0.0	PR (N [mg	Ox-N (g/L/mir 2.58 Max Al Re [r	NA PCelease mg/L]	[mg	SNPF g/g MLV 1.34 Avg. ANA Concen [mg.	R (SS/hr] A NO ₃ -N tration (L) 09
Highest A Conce	AER NO ₂ -entration ng/L]	54 55 58 60 43 44 34 34 36 44 36 MVSS [mg/L] 2600 N PUR (PO. Sides) [mg/L]	[m 1-P Uptake ppe) /L/hr] 91 A1 PO ₄ -P Slop	AER DO g/L] -71 SPL [mg/g ML] 2.5 Release	Min A [unit 6. UR VSS/hr] 5 AA1 S Rel	Clarifier IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	DC NH	(NH4-N S [mg/L/hr] 2.22 H ₄ -N Clat AA2 PC Release	ifier IDO [mg/ 10.98 D ₄ -P Slope	S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con [pecific Fease Ra	r IDC PC centration (mg/L) 0.0	PR (N [mg	Ox-N S g/L/mir 2.58 Max Al Re [r 1 3 PO ₄ ase SI	NA PCelease mg/L] 2.68	[mg	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg. 0.36 Specific	R (SS/hr] (NO ₃ -N) (tration (L) (D9 (PO ₄ -P) (ate
[days 4.35] Highest A Conce [in Conce	SRT Ae] AER NO ₂ - entration ng/L] 0.29	54 55 58 60 43 44 34 34 36 44 36 44 PUR (PO. Slo	[m] 1 1 1-P Uptake oppe) ///hr] 91 A1 PO ₄ -P Slop [mg/L	AER DO g/L] -71 SPU [mg/g ML] 2.5 Release	Min A [unit 6. UR VSS/hr] 5 AA1 S Rel	Clarifier IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	DC NH	(NH4-N S [mg/L/hr] 2.22 H ₄ -N Clat AA2 PC Release [mg/L	ifier IDO [mg/ 10.98 0 ₄ -P Slope	S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con [pecific Fease Raig MLVS:	r IDC PC centration (mg/L) 0.0	PR (N [mg	Ox-N \$ g/L/mir 2.58 Max Al Re [r 1 3 PO ₄ ase Sl	NA PCelease mg/L] 2.68	[mg	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg. 0.36 Specific slease R g MLVS	R (SS/hr] (NO ₃ -N) (tration (L) (D9 (PO ₄ -P) (ate
Idays 4.35 Highest A Conce	SRT Ae] AER NO ₂ - entration ng/L] 0.29	54 55 58 60 43 44 34 34 36 44 36 MVSS [mg/L] 2600 N PUR (PO. Sides) [mg/L]	[m 1-P Uptake ppe) /L/hr] 91 A1 PO ₄ -P Slop	AER DO g/L] -71 SPU [mg/g ML] 2.5 Release	Min A [unit 6. UR VSS/hr] 5 AA1 S Rel	Clarifier IIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIIII	DC NH	(NH4-N S [mg/L/hr] 2.22 H ₄ -N Clat AA2 PC Release	ifier IDO [mg/ 10.98 0 ₄ -P Slope	S. [mg/g N 1	AUR //LVSS/ .15 Clarifie Con [pecific Fease Ra	r IDC PC centration (mg/L) 0.0	PR (N [mg	Ox-N S g/L/mir 2.58 Max Al Re [r 1 3 PO ₄ ase SI	NA PCelease mg/L] 2.68	[mg	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg. 0.36 Specific	R (SS/hr] (NO ₃ -N) (tration (L) (D9 (PO ₄ -P) (ate
[days 4.35] Highest A Conce [n] AA4 NC [mg/L] 0.43	AER NO2-entration ng/L] 29 2x-N AA	54 55 58 60 43 44 34 36 44 36 44 PUR (PO. Sident Siden	[m] A1 PO ₄ -P Slop [mg/L 8.5]	AER DO g/L] .71 SPU [mg/g ML] .2.5 Release be	Min A [unit 6. UR VSS/hr] 5 AA1 S Rel [mg/g	Clarifier II Clarifier II (Clarifier II (Clarifi	DC Nh 19/L] 00000 000000000000000000000000000000	(NH4-N S [mg/L/hr] 2.22 AA2 P(Release [mg/L 6.7)	ifier IDC [mg/ 10.98 D ₄ -P Slope [hr]	S. [mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR //LVSS/ .15 Clarifie Con [pecific Fease Ray // MLVS: 7.70	r IDC PC centratic (mg/L) 0.0 PO ₄ -P ate S/hr]	PR (N	Ox-N S g/L/min 2.58 Max Al Re [r 1 3 PO ₄ ase SI ng/L/hr 4.27	NA PCelease mg/L] 2.68 -P lope	AA3 S Re [mg/	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg 0.30 Specific elease R g MLVS 4.87	R (SS/hr] NO ₃ -N tration (L] 09 PO ₄ -P tate SS/hr]
[days 4.35] Highest A Conce [In Conce AA4 NC [mg/L] 0.43	AER NO ₂ -entration ng/L] 0.29	54 55 58 60 43 44 34 34 36 44 36 40 PUR (PO. Slo	[m] 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AER DO g/L] .71 SPL [mg/g ML] .2.5 Release be	Min A [unit 6. JR VSS/hr] 5 AA1 S Rel [mg/g	Clarifier II Clarifier II Im O.C. Clease Rate G MLVSS/ 9.79	DC NH DC NH	(NH4-N S [mg/L/hr] 2.22 AA2 P(Release [mg/L 6.7)	ifier IDC [mg/ 10.98 D ₄ -P Slope [hr]	S. [mg/g N 1 1 2 NOX-N L] 60 AA2 S Rel [mg/g 6 Specific	AUR //LVSS/ .15 Clarifie Con [pecific Fease Ray // MLVS: 7.70 PO ₄ -P	r IDC PC centratic img/L] 0.0 PO ₄ -P ate S/hr]	PR (N [mg	Ox-N 3 g/L/min 2.58 Max Al Re [r 1 3 PO ₄ ase SI ng/L/hr 4.27 (AA1)	NA PCelease mg/L] 2.68 -P lope r]	AA3 S Re [mg/	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg. 0.36 Specific elease R g MLVS 4.87	R (SS/hr] NO ₃ -N tration (L] 09 PO ₄ -P tate SS/hr]
[days 4.35] Highest A Concording (n Concordi	AER NO ₂ -entration ng/L] 0.29 AAA AAA AAA BIODE	54 55 58 60 43 44 34 34 36 44 robic MLSS [mg/L] 2600 PUR (PO. Side of the second of	[m] 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AER DO g/L] .71 SPU [mg/g ML] .2.5 Release be h/hr] 9 55 PO ₄ -P ase Slope	Min A [unit 6. JR VSS/hr] 5 AA1 S Rel [mg/g	Clarifier II [m] 0.c Specific PC lease Rate g MLVSS/ 9.79 Decific PO ₄ -lease Rate	DC NH 19/L] 0000 0 0 1 P P Re	(NH4-N S [mg/L/hr] 2.22 AA2 PC Release [mg/L 6.70 AA6 PO ₄ -Pelease Slop	ifier IDC [mg/ 10.98 D ₄ -P Slope [hr]	S. [mg/g N 1 1 2 NOx-N L] 60 Rel [mg/g S Specific Release F	AUR ILVSS/ .15 Clarifie Coni pecific Fease Ra MLVS 7.70 PO ₄ -P Rate	r IDC PC centratic [mg/L] 0.0 PO ₄ -P ate S/hr]	PR (N my AA Rele n COD Slope	Ox-N 3 g/L/min 2.58 Max Al Re [r 1 3 PO ₄ ase \$1 ase \$1 u (AA1) U	NA PCelease mg/L] 2.68 -Pelope r]) Spec	AA3 S Re [mg/	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg 0.30 Specific slease R g MLVS 4.87 (AA1 P/	R (SS/hr] (NO ₃ -N) tration (L) (D9 (PO ₄ -P) tate (SS/hr] (PO ₄ -COD
[days 4.35 Highest / Conce [n () () () () () () () () () (AER NO ₂ -entration ng/L] 0.29 0.4-P A/4-P A/5lope hr] [54 55 58 60 43 44 34 34 36 44 36 40 PUR (PO. Slo	[m] A1 PO ₄ -P Slop [mg/L 8.5 A2-P AA Rele	AER DO g/L] .71 SPL [mg/g ML] .2.5 Release be	Min A [unit 6. JR VSS/hr] 5 AA1 S Rel [mg/g	Clarifier II Clarifier II Im O.C. Clease Rate G MLVSS/ 9.79	DC NH 19/L] 0000 0 0 1 P P Re	(NH4-N S [mg/L/hr] 2.22 AA2 P(Release [mg/L 6.7)	ifier IDC [mg/ 10.98 D ₄ -P Slope [hr]	S. [mg/g N 1 1 2 NOX-N L] 60 AA2 S Rel [mg/g 6 Specific	AUR ILVSS/ .15 Clarifie Coni pecific Fease Ra MLVS 7.70 PO ₄ -P Rate	r IDC PC centratic img/L] 0.0 PO ₄ -P ate S/hr]	PR (N [mg]	Ox-N 3 g/L/min 2.58 Max Al Re [r 1 3 PO ₄ ase \$1 ase \$1 u (AA1) U	NA PCelease mg/L] 2.68 -Pelope r]) Spec	AA3 \$ Ref [mg/	SNPF 3/g MLV 1.34 Avg. ANA Concen [mg 0.30 Specific elease R g MLVS 4.87 CAA1 P//	R (SS/hr] NO ₃ -N tration (L] 09 PO ₄ -P tate SS/hr]

# Chem Used	FeCl3 Hrs U	sed FeCl3	3 Addition	Active Bios	-	ARCY	/ Flow	CRO	CY Flow	NRCY Flow
[lbs]	[hr]	[r	mg/L]	[MGD]		[M	GD]	[1	MGD]	[MGD]
1500	22.000	5	5.100	24.58		17.	.68	14	4.032	0
ANA MLTSS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	L AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	'L]	[%]	[mg/L]
1140	77	878	2600	74	1	924	260	0	74	1924



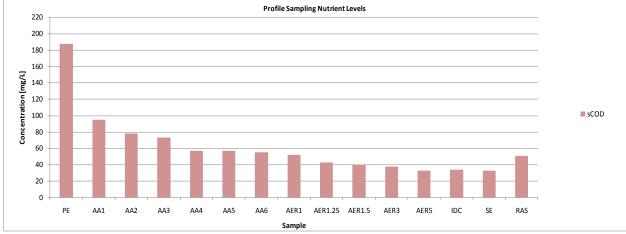


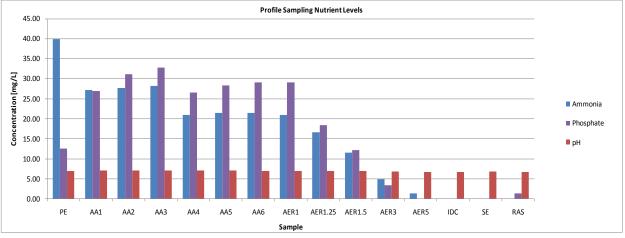


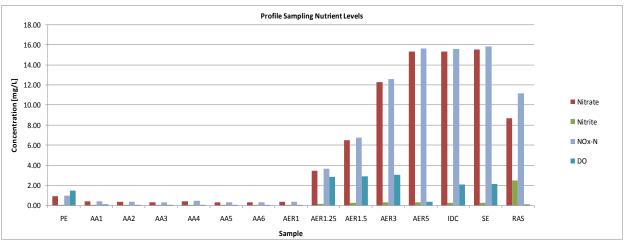
Week 9 – 9/14/09, 9/15/09, & 9/18/09

N eek ROFII F 1	1 Train 4	1. Aera	ition Tank 4	Clarifier 4														
IXOI ILL I	i iiaiii i																	
			HRT along BNF Process (w/out										NOx-N by I	Nitrate HAC	CH Phosph	nate by HA	СН	
Date	Sample		Recycles)	Sample N	mber R	Reactor Typ	e Amm	nonia by HACH T	NT Nitra	ate by HACH TN	T Nitrite by F		,	te HACH		TNT844	DO	pН
dd-mmm-yy	h:mn	m	hr					mg/L NH3-N		mg/L NO3-N	mg/L N	102-N	mg/L	NOx-N	mg	/L PO4-P	[mg/l	.] [unitle
14-Sep-09	7:00		0	PE		-		39.90		0.91	0.0			.95		12.50	1.48	_
14-Sep-09 14-Sep-09	7:15 7:45		0.84 1.68	AA1		Anaerobic Anaerobic		27.20 27.70		0.42	0.0			i.43 i.38		26.90 31.20	0.09	
14-Sep-09	8:15		2.53	AA3		Anaerobic		28.20		0.32	0.0			.33		32.80	0.08	
14-Sep-09	8:45		3.37	AA4		Anaerobic		20.90		0.42	0.0			.46		26.50	0.06	
14-Sep-09	9:00		4.21	AA5		Anaerobic		21.45		0.32	0.0			.33		28.30	0.07	_
14-Sep-09 14-Sep-09	9:15 9:30		5.05 0.00	AA6 AER		Anaerobic Aerobic		21.45 21.00		0.32	0.0			i.33 i.38		29.10 29.10	0.08	
14-Sep-09	9:38		0.33	AER1.		Aerobic		16.65		3.48	0.4			.64		18.40	2.83	
14-Sep-09	9:46		0.65	AER1		Aerobic		11.50		6.51	0.2			5.77		12.20	2.90	_
14-Sep-09	10:3		3.26	AER		Aerobic		4.92		12.26	0.3			2.59		3.35	3.05	
14-Sep-09 14-Sep-09	11:3		6.52	AER:)	Aerobic		1.34 0.00		15.30 15.30	0.3			5.62 5.58		0.00	2.08	
14-Sep-09	14:3			SE				0.00		15.54	0.2			5.81		0.00	2.16	_
14-Sep-09	14:5			RAS				0.00		8.70	2.4			1.18		1.38	0.13	_
			mg/L NH ₄ -N	1010				0.00		0.1.0	2	-	'				0.10	0.7
			23 mg/L NO ₃ -N															
			5 mg/L PO ₄ -P															
QA/	QC	Sa	mple	=	SE													
	EP	A 3	365.1		EP	PA 35	50.1	EPA	353	3.2 I	EPA 3:	53.2	EP	A 35	53.2	EF	PA 3	53.2
s	COI	Db	у СЕ	ட	(O-PC) 4	N	H4-N	1	$NO_{2,3}$	-N	1	VO3-	N	1	NO ₂ -	-N
			_			mg/I	_	n	ıg/L		mg/l			mg/I	L		mg/	L
	Li	mg	<i>/</i> L]															
	Li	18	8															
	L	18 95	8 5															
	Lı	18 95 78	5 3															
	[ı	18 95	88 5 3															
	[1	18 95 78 73 57	88 5 3 3 7			J												
	Lı	18 95 78 73 57 57	88 5 3 3 7 7															
	Lı	18 95 78 73 57	88 5 3 3 7 7 5															
	L	18 95 73 57 57 55 52 43 40	18															
	L	18 95 78 73 57 57 55 52 43 40 38	18															
	L	18 95 78 73 57 57 55 43 40 38 33	88 5 3 3 7 7 7 5 5 2 2 3 3 0 8															
	L	18 95 75 57 57 57 57 54 40 38 33 34	88 5 3 3 7 7 5 5 2 2 3 0 3 3 4),35		16.1			15.8			0.31	
		18 95 78 73 57 57 55 43 40 38 33	88 5 3 3 7 7 5 5 2 3 3 0 3 3 3 4 3 3 3 4 3 3 3 3 3 3 3 3 3						0.35		16.1			15.8	3		0.31	
		18 95 78 73 57 57 55 43 43 33 34 33	88 5 3 3 7 7 5 5 2 3 3 0 3 3 3 4 3 3 3 4 3 3 3 3 3 3 3 3 3						0.35		16.1			15.8			0.31	L
		18 95 78 73 57 57 55 43 43 33 34 33	88 5 3 3 7 7 5 5 2 3 3 0 3 3 3 4 3 3 3 4 3 3 3 3 3 3 3 3 3						0.35		16.1		<u> </u>	15.8	1		0.31	
		18 95 78 75 55 55 55 54 40 38 33 34 33 57	68 5 3 3 7 7 5 5 2 3 3 0 3 3 3 4 4	Average			Min Al			4-N Slope			NP			e)		
Actuals	SRT	188 955 765 557 557 558 440 388 33 344 355 557 Aero	68 5 3 3 7 7 7 5 2 3 3 0 3 3 4 4 3	Averag	e AER			ER pH AL	JR (NH	4-N Slope)	S	AUR		R (NOx	-N Slop		SNF	PR
[days	SRT s]	188 95 787 57 55 55 54 40 33 33 57	68 5 3 3 7 7 5 5 2 3 3 0 3 3 4 4 3 1	[1	e AER		[unitl	ER pH AL	JR (NH	/L/hr]	S [mg/g M	AUR //LVSS//		R (NOx [mg/L	(-N Slop /min]		SNF g/g ML	PR VSS/h
	SRT s]	188 95 787 57 55 55 54 40 33 33 57	68 5 3 3 7 7 7 5 2 3 3 0 3 3 4 4 3	[1	e AER			ER pH AL	JR (NH		S [mg/g M	AUR		R (NOx	(-N Slop /min]		SNF	PR VSS/h
[days	SRT s]	188 955 787 787 557 555 522 433 333 557 Aero	68 5 3 3 7 7 5 5 2 3 3 3 4 3 1 1 bic MLSS mg/L] 2760	[1	e AER ng/L]		[unitl	ER pH AL	JR (NH	/L/hr]	S [mg/g M	AUR //LVSS//	nr]	R (NOx [mg/L	r-N Slop /min] 28	[m	SNF g/g ML 1.6	PR VSS/h
[days	SRT s] 3	188 999 787 757 557 557 552 433 33 557 Aero [II	68 5 3 3 7 7 5 2 3 3 4 3 1 1 bic MLSS mg/L] 2760	-P Uptake	e AER ng/L]	R DO	[unitl	ER pH AU	JR (NH- [mg. 2.	/L/hr] .67	S [mg/g N	AUR //LVSS/h 1.32 Clarifier	IDC PO	R (NOx [mg/L 3.2 ₄ -P Ma	(-N Slop /min] 28 ax ANA F	[m	SNF g/g ML 1.6 Avg. AN	PR VSS/h 3 A NO ₃ -
[days 5.93 Highest A	SRT s] 3	188 999 787 757 557 557 552 433 33 557 Aero [II	68 5 3 3 7 7 5 2 3 3 4 3 1 1 bic MLSS mg/L] 2760	-P Uptake	e AER ng/L] 1.84	R DO I	[unitl	ER pH AUess] 67 Clarifier IDC	JR (NH- [mg. 2.	/L/hr] .67 Clarifier II	S [mg/g N	AUR //LVSS/I	IDC PO.	R (NOx [mg/L 3.2 ₄ -P Ma	x-N Slop /min] 28 ax ANA F Release	[m	SNF g/g ML 1.6 Avg. AN Conce	PR VSS/h 3 A NO ₃ - ntration
[days 5.93 Highest A Conce	SRT s] 3	188 999 787 757 557 557 552 433 33 557 Aero [II	68 5 3 3 7 7 5 5 2 3 3 3 4 3 1 1 PUR (PO. Sic [mg/L]	-P Uptake	e AER ng/L] 1.84	R DO	[unitl	ER pH AU	JR (NH- [mg. 2. : NH4-N	/L/hr] .67 Clarifier II	S [mg/g N	AUR //LVSS/I	IDC PO	R (NOx [mg/L 3.2 ₄ -P Ma	(-N Slop /min] 28 ax ANA F	[m	SNF g/g ML 1.6 Avg. AN Conce	PR VSS/h 3 A NO ₃ -
[days 5.93 Highest A Conce	SRT s] 3 AER Note that the second se	188 999 787 757 557 557 552 433 33 557 Aero [II	68 5 3 3 7 7 5 5 2 3 3 3 4 3 1 1 PUR (PO. Sic [mg/L]	P Uptake	e AER ng/L] 1.84	SPUR	[unitl	ER pH AL ess] 67 Clarifier IDC [mg/l	JR (NH- [mg. 2. : NH4-N	/L/hr] .67 Clarifier II	S [mg/g N	AUR //LVSS/I	IDC PO	R (NOx [mg/L 3.2 ₄ -P Ma	x-N Slop /min] 28 ax ANA F Releas [mg/L]	[m	SNF g/g ML 1.6 Avg. AN Conce	PR VSS/h 3 A NO ₃ - ntration g/L]
[days 5.93 Highest A Conce	SRT s] 3 AER Note that the second se	188 999 787 757 557 557 552 433 33 557 Aero [II	68 5 3 3 7 7 5 2 3 3 3 4 3 1 1 PUR (PO. Slo	-P Uptake ppe) [L/hr] 28	€ AER ng/L] 1.84	SPUR	[unitl 6.0	ER pH AU ess] 67 Clarifier IDC [mg/l 0.000	JR (NH- [mg. 2. : NH₄-N -] 0	/L/hr] .67 Clarifier II [m	S [mg/g N]	AUR MLVSS/t .32 Clarifier Conc	IDC POcentration	R (NOx [mg/L 3.2 4-P Man	x-N Slop /min] 28 ax ANA F Release [mg/L] 20.30	[m	SNF g/g ML 1.6 Avg. AN Conce [m 0.3	PR VSS/h 3 A NO ₃ - ntration g/L]
[days 5.93 Highest A Conce [n	SRT s] 3 AER Note that is a second se	188 95 78 78 78 78 78 78 78 78 78 78 78 78 78	68 5 3 3 7 7 5 2 3 3 3 4 3 1 1 PUR (PO. Slo	-P Uptake ope) 'L/hr] 28	e AERng/L] 1.84 [mg/	SPUR	[unitl 6.0 SS/hr]	ER pH AU ess] 67 Clarifier IDC [mg/l 0.000	JR (NH- [mg. 2. : NH4-N- -] 0	/L/hr] .67 Clarifier II	S [mg/g N / 2 2 2 2 2 2 2 2 2 2	AUR MLVSS/h .32 Clarifier Conc	PO ₄ -P	R (NOx mg/L 3.2	x-N Slop /min] 28 ax ANA F Release [mg/L] 20.30	PO ₄ -P	SNF g/g ML 1.6 Avg. AN Conce [m 0.:	PR VSS/h 3 A NO ₃ - ntration g/L] 360
[days 5.93 Highest A Conc	SRT s] 3 AER Note entration mg/L] 0.33 Dx-N /	18 95 78 75 55 55 55 44 40 33 33 57 Aero	68 5 3 3 7 7 5 2 2 3 3 3 4 3 1 1 PUR (PO. Slo [mg/L]	-P Uptako ppe) L/hr] 28 A1 PO ₄ -	e AERng/L] 1.84 [mg/ P Releippe	SPUR	[unitl 6.4 SS/hr] A1 Sp Rele	ER pH AU ess] 67 Clarifier IDC [mg/l 0.000 pecific PO ₄ - ease Rate	JR (NH- [mg. 2. : NH4-N -] 0	/L/hr] .67 Clarifier II [m 15. A2 PO ₄ -P lease Slop	S [mg/g N]	AUR //LVSS/H .32 Clarifier Conc [i	DC PO.centration mg/L] 0.0 PO ₄ -P	R (NOx [mg/L 3.2 4-P Man	x-N Slop /min] 28 ax ANA F Releas [mg/L] 20.30 PO ₄ -P e Slope	PO4-P e AA3	SNF g/g ML 1.6 Avg. AN Conce [m 0.:	PR VSS/h 3 A NO ₃ ntration g/L] 360 C PO ₄ Rate
[days 5.93 Highest AA4 NC [mg/L]	SRT si 3 AER No centratio mg/Lj 0.33 Dx-N /	18 95 78 75 55 55 55 44 40 33 33 57 Aero [II]	68 5 3 3 7 7 5 2 3 3 3 4 3 1 1 PUR (PO. Slo [mg/L] 6.	-P Uptake ppe) 'L/hr] 28 A1 PO ₄ - Slo [mg/	e AERng/L] 1.84 [mg/ Pelectric Release Pelectric Pelectric Release Pelectric Pelectric Release Pelect	SPUR	[unitl 6.4 SS/hr] A1 Sp Rele	ER pH AU ess] 67 Clarifier IDC [mg// 0.000 pecific PO ₄ - ease Rate MLVSS/hr	JR (NH- [mg. 2. : NH4-N -] 0	/L/hr] .67 Clarifier II	S [mg/g N]	AUR //LVSS/t .32 Clarifier Conc [i pecific F ease Ra multiple MLVSS	DC PO.centration mg/L] 0.0 PO ₄ -P	R (NOx [mg/L 3.2 4-P Man AA3 F Release [mg/	x-N Slop /min] 28 ax ANA F Releasi [mg/L] 20.30 PO ₄ -P e Slope 'L/hr]	PO4-P e AA3	SNF g/g ML 1.6 Avg. AN Conce [m 0.: Specific elease	PR VSS/h 3 A NO ₃ A NO ₃ of the state of t
[days 5.93 Highest A Conc	SRT si 3 AER No centratio mg/Lj 0.33 Dx-N /	18 95 78 75 55 55 55 44 40 33 33 57 Aero [II]	68 5 3 3 7 7 5 2 2 3 3 3 4 3 1 1 PUR (PO. Slo [mg/L]	-P Uptako ppe) L/hr] 28 A1 PO ₄ -	e AERng/L] 1.84 [mg/ Pelectric Release Pelectric Pelectric Release Pelectric Pelectric Release Pelect	SPUR	[unitl 6.4 SS/hr] A1 Sp Rele	ER pH AU ess] 67 Clarifier IDC [mg/l 0.000 pecific PO ₄ - ease Rate	JR (NH- [mg. 2. : NH4-N -] 0	/L/hr] .67 Clarifier II [m 15. A2 PO ₄ -P lease Slop	S [mg/g N]	AUR //LVSS/H .32 Clarifier Conc [i	DC PO.centration mg/L] 0.0 PO ₄ -P	R (NOx [mg/L 3.2 4-P Man AA3 F Release [mg/	x-N Slop /min] 28 ax ANA F Releas [mg/L] 20.30 PO ₄ -P e Slope	PO4-P e AA3	SNF g/g ML 1.6 Avg. AN Conce [m 0.:	PR VSS/h 3 A NO ₃ A NO ₃ of the state of t
[days 5.93 Highest Conc. [n (SRT s] 3 AER Note that it is a second secon	18 95 78 75 55 55 55 52 40 40 40 33 33 57 60 70 70 70 70 70 70 70 70 70 70 70 70 70	BB 5 5 3 3 5 7 7 7 5 2 2 3 3 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1	L/hr] 28 A1 PO ₄ -Slo	⊋ AER ng/L] 1.84 [mg 2 Rele pe L/hr] 72	SPUR 3/g MLVS 3.12 ease A	[unitl 6.4 SS/hr] A1 Sp Rele [mg/g	Clarifier IDC [mg/l 0.000 Decific PO4- pase Rate MLVSS/hr 7.00	R (NH- [mg, 2.] 2. NH4-N NH4-N Rel	/L/hr] .67 Clarifier II	S [mg/g N]	AUR ILVSS/t .32 Clarifier Conc [i	IDC PO. centration mg/L] 0.0 PO ₄ -P	R (NOx [mg/L] 3.2	x-N Slop /min] 28 ax ANA F Release [mg/L] 20.30 PO ₄ -P e Slope [L/hr]	PO ₄ -P e AA3	SNF g/g ML 1.6 Avg. AN Conce [m 0.: Specific elease /g MLV 5.74	VSS/h 3 A NO ₃ A NO ₃ ntratior g/L] 360 c PO ₄ Rate SS/hr
[days 5.93 Highest Aconc [n () () () () () () () () () (SRT s] 3 AER Note that the control of the control	18 95 78 78 78 78 78 78 78 78 78 78 78 78 78	BB 5 5 8 5 8 5 8 5 8 5 8 8 5 8 8 5 8 8 6 7 7 7 5 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9 9	-P Uptakeppe) 'L/hri] 28 A1 PO ₄ - Slo [mg/ 7.	e AER ng/L] 1.84 [mg Pelulum 72 A5 PO	SPUR 3/g MLVS 3.12 ease A	[unitl 6.6 SS/hr] A1 Sp Rele [mg/g	Clarifier IDC [mg/l 0.000 Decific PO ₄ -P Pecific PO ₄ -P	R (NH- mg, 2 NH4-N mg, 2 NH4-N mg, 2 NH4-N mg, 2 AA6	/L/hr] .67 Clarifier II Im In	S S [mg/g N / n / n / n / n / n / n / n / n / n /	AUR //LVSS/t .32 Clarifier Conc [i pecific F ease Ra MLVSS 8.97	r IDC PO. centration mg/L] 0.0 PO ₄ -P tte	R (NOx [mg/L 3.2 4-P Man AA3 F Release [mg/6.	x-N Slop /min] 28 ax ANA F Release [mg/L] 20.30 PO ₄ -P e Slope /L/hr] 33	AA3 R	SNF g/g ML 1.6 Avg. AN Conce [m 0.: Specific elease /g MLV 5.74 COD (A	PR VSS/h 3 A NO ₃ - ntratior g/L] 360 E PO ₄ - Rate SS/hr]
[days 5.93 Highest Conc. [n (SRT s] 3 AER Note that the second se	18 95 78 75 75 55 55 55 55 55 55 55 55 55 55 55	BB 5 5 3 3 5 7 7 7 5 2 2 3 3 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1 5 1	-P Uptakeppe) 'L/hr] 28 A1 PO ₄ - Slo [mg/ 7.	⊋ AER ng/L] 1.84 [mg 2 Rele pe L/hr] 72	SPUR SPUR 3/g MLVS 3.12 ease A	[unittle factors of the content of t	Clarifier IDC [mg/l 0.000 Decific PO4- pase Rate MLVSS/hr 7.00	IR (NH- Img. 2. NH4-N O Relea	/L/hr] .67 Clarifier II	S [mg/g N]	AUR //LVSS/t .32 Clarifier Conc [i pecific F ease Ra multiple m	IDC PO. entration mg/L] 0.0 PO ₄ -P te (AA1) Uptake	R (NOx [mg/L] 3.2	x-N Slope /min] 28 ax ANA F Release [mg/L] 20.30 PO ₄ -P e Slope [L/hr] 33	PO ₄ -P e AA3	SNF g/g ML 1.6 Avg. AN Conce [m 0.: Specific elease /g MLV 5.74 COD (A	PR VSS/h 3 A NO ₃ - ntration g/L] 360 C PO ₄ - Rate SS/hr]

# Chem Used	FeCl3 Hrs U	sed	FeCl3	Addition	Active Bios		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]		[n	ng/L]	[MGD]		[M	GD]	[]	MGD]	[MGD]
0	0.000		0	.000	19.84		18.	.03	1:	2.075	0
ANA MLTS	S ANA % VOL	ANA	MLVSS	ANX MLTS	S ANX % VOL	ANX	MLVSS	AER M	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[m	ng/L]	[mg/L]	[%]	[n	ng/L]	[mg/	′L]	[%]	[mg/L]
1360	81	1	102	2760	73	2	015	276	0	73	2015

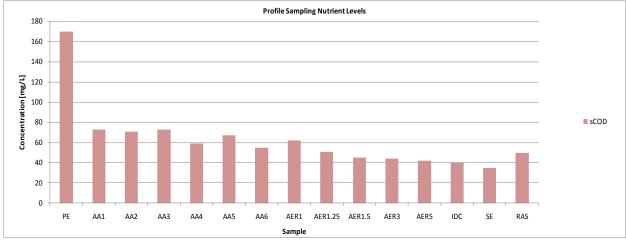


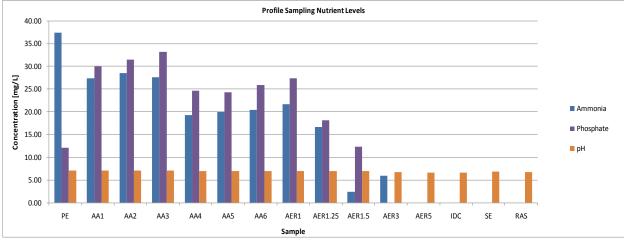


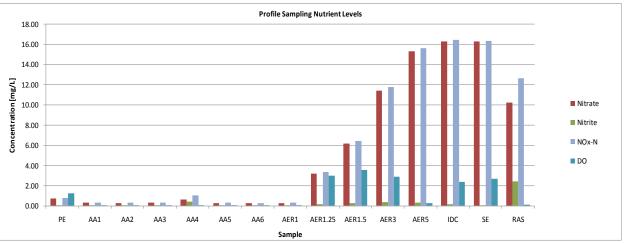


ROFILE 2	2 Train 6, Ae	ration rank 5,	Ciai iiiei J											
Date	Sample Time		Sample Number	Reactor Type		l TNT Nitra	ate by HACH TNT	Nitrite by HACH TN	+ N	oy Nitrate HAC litrite HACH	, T	ate by HACI NT844	DO	рН
dd-mmm-yy	h:mm	hr			mg/L NH3-N		mg/L NO3-N	mg/L NO2-N	mį	g/L NOx-N		L PO4-P	[mg/L]	[unitless
15-Sep-09	6:50	0	PE		37.40		0.75	0.06		0.81	_	12.10	1.26	7.15
15-Sep-09	7:08	0.84	AA1	Anaerobic	27.40		0.33	0.02		0.35		30.00	0.07	7.13
15-Sep-09 15-Sep-09	7:38 8:08	1.68 2.53	AA2 AA3	Anaerobic Anaerobic	28.50		0.30	0.01		0.31	_	31.50 33.20	0.07	7.11 7.09
15-Sep-09	8:38	3.37	AA3 AA4	Anaerobic	27.60 19.25		0.63	0.01 0.43		1.05	_	24.60	0.08	7.03
15-Sep-09	8:53	4.21	AA5	Anaerobic	19.95		0.03	0.02		0.31	_	24.30	0.00	7.02
15-Sep-09	9:08	5.05	AA6	Anaerobic	20.45		0.28	0.01		0.30	_	25.90	0.07	7.00
15-Sep-09	9:23	0.00	AER1	Aerobic	21.70		0.30	0.02		0.31	_	27.40	0.09	7.04
15-Sep-09	9:31	0.33	AER1.25	Aerobic	16.65		3.22	0.15		3.37		18.20	3.02	7.04
15-Sep-09	9:39	0.65	AER1.5	Aerobic	2.44		6.16	0.27		6.43		12.30	3.55	7.02
15-Sep-09	10:28	3.26	AER3	Aerobic	5.92		11.42	0.37		11.79		0.00	2.89	6.82
15-Sep-09	11:28	6.52	AER5	Aerobic	0.00		15.30	0.32		15.62		0.00	0.26	6.64
15-Sep-09	11:32		IDC	-	0.00		16.28	0.17		16.45		0.00	2.36	6.70
15-Sep-09	14:32		SE	-	0.00		16.30	0.02		16.32		0.00	2.67	6.93
15-Sep-09	14:45		RAS	-	0.00		10.22	2.44		12.66		0.00	0.13	6.76
_	se values are <	1 mg/L NHN												
		•												
		0.23 mg/L NO ₃ -N												
'NOTE: Thes	se values are <	0.5 mg/L PO ₄ -P												
QA/	QC S	ample	: SE	=										
	EPA	365.1	Е	PA 35	0.1 EP	A 353	3.2 E	PA 353.2	Е	PA 35	53.2	EP.	A 35	3.2
s	COD	by CE	_	O-PO	4 N	1H ₄ -N	4	$NO_{2,3}-N$		NO_3 -	N	N	O_2 -	Ν
	[m	g/L]		mg/L	, 1	mg/L	,	mg/L		mg/I	,	1	ng/I	,
	1	70												
		73												
	7													
	7	73 71	-11											
	7	73 71 73												
	7 7 7	73 71 73 59	1											
	7 7 5	73 71 73 59 57												
	7 7 5 6	73 71 73 59 57												
	77 77 55 66 67	73 71 73 59 57 55												
	7 7 5 6 6 6	73 71 73 59 57 55 52												
	7 7 5 6 6 5	73 71 73 59 57 55												
	7 7 5 6 5 6	73 71 73 59 57 55 52 51 45												
	7 7 7 6 6 6 6 2	73 71 73 59 57 55 55 52 51 45 44												
	6 6 6 5 2 2	73 71 73 59 57 55 52 51 45 44		0.160		~0.20°		17.7		17.7			. 02	
	77 77 77 77 77 6 6 6 6 6 6 7 7 7 7 7 7	73 71 73 59 57 55 52 51 45 44 42 40		0.160		<0.20		17.7		17.7			0.02	
	77 77 77 77 77 6 6 6 6 6 6 7 7 7 7 7 7	73 71 73 59 57 55 52 51 45 44		0.160		<0.20		17.7		17.7			0.02	
	77 77 77 77 77 6 6 6 6 6 6 7 7 7 7 7 7	73 71 73 59 57 55 52 51 45 44 42 40		0.160		<0.20		17.7	<u> </u>	17.7			0.02	
	77 77 77 77 8 8 8 8 8 8 2 2 2 2 2 2 2 2	73 71 73 59 57 55 52 51 45 44 42 40 85	Autrogo Al								N Slope			
	77 77 77 77 77 77 77 77 77 77 77 77 77	73 71 73 59 57 55 52 51 45 44 42 40 35 50		ER DO M	lin AER pH	AUR (NH	l4-N Slope)	SAUR		IPR (NOx		e)	SNPI	
Actual S	77 77 77 77 77 77 77 77 77 77 77 77 77	73 71 73 59 57 55 52 51 45 44 42 40 85	Average At	ER DO M		AUR (NH	H4-N Slope)					e)	SNPI	
	77 77 77 77 77 77 77 77 77 77 77 77 77	73 71 73 59 57 55 52 51 45 44 42 40 35 50		ER DO M	lin AER pH	AUR (NH	l4-N Slope)	SAUR		IPR (NOx	/min]	e)	SNPI	'SS/hr]
[days 6.38	77 77 77 77 77 77 77 77 77 77 77 77 77	73 71 73 59 57 55 52 51 45 44 42 40 35 50	[mg/L 1.96	ER DO M	lin AER pH /	AUR (NH	H4-N Slope)	SAUR [mg/g MLVS: 1.21		IPR (NOx [mg/L	/min]	e) [mg.	SNPI g MLV 1.53	'SS/hr]
[days 6.38 Highest	77 77 77 77 77 77 77 77 77 77 77 77 77	73 71 73 59 67 65 65 62 61 45 44 42 40 85 60 robic MLSS [mg/L] 2640	[mg/L 1.96 -P Uptake	ER DO M	lin AER pH // [unitless] 6.64	AUR (NH [mg. 2.	14-N Slope) g/L/hr] 2.43	SAUR [mg/g MLVS: 1.21	S/hr] er IDC P	IPR (NOx [mg/L 3.0	/min] 07 ux ANA P	(mg.	SNPI /g MLV 1.53 vg. ANA	(SS/hr]
[days 6.38 Highest A	SRT Aeess AER NO ₂ -lentration	73 71 73 59 67 65 65 62 61 45 44 42 40 35 60 robic MLSS [mg/L] 2640 PUR (PO ₄ Slo	[mg/L 1.96 -P Uptake pe)	ER DO M	lin AER pH A [unitless] 6.64 Clarifier II	AUR (NH- [mg 2. DC NH4-N	14-N Slope) g/L/hr] 2.43 U Clarifier ID	SAUR [mg/g MLVS: 1.21 Clari	S/hr] ier IDC P	IPR (NOx [mg/L 3.0	/min] 07 x ANA P Release	(mg.	SNPI /g MLV 1.53 vg. ANA Concen	SS/hr] NO ₃ -N
[days 6.38 Highest A Conce	SRT Aeess AER NO2-lentration	73 71 73 59 67 65 65 62 61 45 44 42 40 35 60 robic MLSS [mg/L] 2640 PUR (PO ₄ Sic	[mg/L 1.96 -P Uptake pe) L/hr] [r	SPUR	lin AER pH A [unitless] 6.64 Clarifier IE	AUR (NH- [mg, 2. DC NH ₄ -N	H4-N Slope) g/L/hr] 2.43 UClarifier ID	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co	S/hr] ler IDC P ncentrati	IPR (NOx [mg/L 3.0	/min] D7 DX ANA P Release [mg/L]	(mg.	SNPI /g MLV 1.53 vg. ANA Concen [mg	SS/hr] NO ₃ -N tration
[days 6.38 Highest A Conce	SRT Aeess AER NO ₂ -lentration	73 71 73 59 67 65 65 62 61 45 44 42 40 35 60 robic MLSS [mg/L] 2640 PUR (PO ₄ Sic	[mg/L 1.96 -P Uptake pe)	ER DO M	lin AER pH A [unitless] 6.64 Clarifier II	AUR (NH- [mg, 2. DC NH ₄ -N	14-N Slope) g/L/hr] 2.43 U Clarifier ID	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co	S/hr] ier IDC P	IPR (NOx [mg/L 3.0	/min] 07 x ANA P Release	(mg.	SNPI /g MLV 1.53 vg. ANA Concen	SS/hr] NO ₃ -N tration
[days 6.38 Highest A Conce	SRT Ae SI AER NO2-lentration mg/L]	73 71 73 59 67 65 65 62 61 45 44 42 40 85 60 PUR (PO4 Slo	[mg/L 1.96	SPUR ng/g MLVS: 3.53	lin AER pH // [unitless] 6.64 Clarifier II S/hr] [m; 0.0	AUR (NH- [mg. 2. DC NH4-N g/L] 000	14-N Slope) 2/L/hr] 2.43 Clarifier ID	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co yL] 540 AA2 Specific	er IDC Pencentration (mg/L) 0.0	IPR (NOx [mg/L 3.0 O ₄ -P Maion	/min] D7 EX ANA P Release [mg/L] 21.10 PO ₄ -P	O ₄ -P A	SNPI /g MLV 1.53 vg. ANA Concen [mg 0.33	NO ₃ -No ₃ -No ₄ -PO ₄ -P
[days 6.38 Highest A Conce [n	5	73 71 73 59 67 55 62 61 45 44 42 40 85 60 Tobic MLSS Tobic MLSS	[mg/L 1.96	SPUR ng/g MLVS: 3.53	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate	AUR (NH4-N [mg. 2. DC NH4-N [g/L] 0000	I4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg] 16.4 AA2 PO ₄ -P elease Slope	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co y/L] 540 AA2 Specific Release	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P	IPR (NOx [mg/L 3.0 O ₄ -P Ma ion AA3 F Releas	/min] D7 Ex ANA P Release [mg/L] 21.10 PO ₄ -P e Slope	O ₄ -P A	SNPI /g MLV 1.53 wg. ANA Concen [mg 0.33 pecific ease R	SS/hr] NO ₃ -N tration
[days 6.38 Highest Conce [n	5	73 71 73 59 67 65 65 62 61 45 44 42 40 85 60 PUR (PO4 Slo [mg/L] 7. A	[mg/L 1.96 -P Uptake pe) L/hr] [r 10 A1 PO ₄ -P Re Slope [mg/L/hr]	SPUR ng/g MLVS: 3.53	lin AER pH // [unitless] 6.64 Clarifier II S/hr] [m; 0.0	AUR (NH4-N [mg. 2. DC NH4-N [g/L] 0000	I4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg] 16.4 AA2 PO ₄ -P elease Slope [mg/L/hr]	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co yL] 540 AA2 Specific	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P	IPR (NOx [mg/L 3.0 O ₄ -P Maion	/min] D7 Ex ANA P Release [mg/L] 21.10 PO ₄ -P e Slope	O ₄ -P A	SNPI /g MLV 1.53 vg. ANA Concen [mg 0.33	SS/hr NO ₃ -N tration /L] 56 PO ₄ -P
[days 6.38 Highest A Conce [n	5	73 71 73 59 67 55 62 61 45 44 42 40 85 60 Tobic MLSS Tobic MLSS	[mg/L 1.96	SPUR ng/g MLVS: 3.53	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate	AUR (NH4-N [mg. 2. DC NH4-N [g/L] 0000	I4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg] 16.4 AA2 PO ₄ -P elease Slope	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co y/L] 540 AA2 Specific Release	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P	IPR (NOx [mg/L 3.0 O ₄ -P Ma ion AA3 F Releas	/min] D7 IX ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr]	O ₄ -P A	SNPI /g MLV 1.53 wg. ANA Concen [mg 0.33 pecific ease R	SS/hr NO ₃ -N tration /L] 56 PO ₄ -P
[days 6.38 Highest A Conce [n]	5	73 71 73 59 67 55 52 51 45 44 42 40 85 60 robic MLSS [mg/L] 2640 N PUR (POa Slo	[mg/L 1.96 -P Uptake pe) L/hr] [r 10 A1 PO ₄ -P Re Slope [mg/L/hr]	SPUR ng/g MLVS: 3.53	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate mg/g MLVSS/h	AUR (NH4-N [mg. 2. DC NH4-N [g/L] 0000	I4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg] 16.4 AA2 PO ₄ -P elease Slope [mg/L/hr]	SAUR [mg/g MLVS: 1.21 Clari C NOx-N Co y/L] 540 AA2 Specific Release [mg/g MLV	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P	IPR (NOx [mg/L 3.0 O ₄ -P Ma ion AA3 F Release	/min] D7 IX ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr]	O ₄ -P A	SNPI /g MLV 1.53 vg. ANA Concen [mg 0.33 pecific ease R MLVS 5.84	(SS/hr) NO ₃ -N tration (L] 56 PO ₄ -P tate
[days 6.38 Highest Conco	5 SRT Ae SRT Ae SRT Ae Ng/L 0.37	73 71 73 59 67 55 62 61 45 44 42 40 85 60 robic MLSS [mg/L] 2640 N PUR (POal Sic [mg/L] 7. A 6 NOx-N mg/L] 0.30	[mg/L 1.96 1.9	SPUR ng/g MLVS: 3.53	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate mg/g MLVSS/h 12.36	AUR (NH-N [mg. 2. DC NH ₄ -N [g/L] 0000 Rel nr]	14-N Slope) 3/L/hr] 2.43 Clarifier ID	SAUR [mg/g MLVS: 1.21 Claric C NOx-N Co y/L] 540 AA2 Specific Release [mg/g MLV 8.27	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P Rate SS/hr]	IPR (NOx [mg/L 3.0 O ₄ -P Maion AA3 I Release [mg/ 6.	/min] D7 IX ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr]	O ₄ -P A	SNPI (g MLV 1.53 ug. ANA Concen [mg 0.34 MLVS 5.84 (AA	NO ₃ -No ₃ -No ₄ -
[days 6.38 Highest . Conco	77 77 77 77 77 77 77 7	73 71 73 59 71 73 59 67 55 62 51 45 44 42 40 85 60 85 60 8 PUR (PO ₄ Slo [mg/L] 7. A 6 NOx-N mg/L] 0.30 A4 Specific PO A4 Specific PO	[mg/L 1.96	SPUR ng/g MLVS: 3.53 elease A/	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate mg/g MLVSS/h 12.36 A5 Specific PO ₄	AUR (NH-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-	H4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg 16.4 AA2 PO ₄ -P elease Slope [mg/L/hr] 9.02	SAUR [mg/g MLVS: 1.21 Claric C NOx-N Co y/L] 540 AA2 Specific Release [mg/g MLV 8.27	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P Rate SS/hr]	IPR (NOx [mg/L] 3.0	/min] D7 Ex ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr] 37	AA3 S Rel	SNPP 1.53 g. ANA g. ANA g. ANA g. SA g. ANA	NO ₃ -N tration /L] 566 PO ₄ -F cate SS/hr]
[days 6.38 Highest Conco	77 77 77 77 77 77 77 7	73 71 73 59 67 55 62 51 45 44 42 40 85 60 8 PUR (PO ₄ 8lc [mg/L] 2640 8 PUR (PO ₄ 8lc [mg/L] 0.30 8 A4 Specific PO Release Rat	[mg/L] 1.96 P Uptake pe) L/hr] [r] 10 A1 PO ₄ -P Re Slope [mg/L/hr] 13.48 O ₄ -P AA5 F Release	SPUR ng/g MLVS: 3.53 elease A/	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate mg/g MLVSS/h 12.36	AUR (NH-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-N-	14-N Slope) 3/L/hr] 2.43 Clarifier ID	SAUR [mg/g MLVS: 1.21 C NOx-N Co yL] 540 AA2 Specific Release [mg/g MLV 8.27 AA6 Specific POa Release Rate	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P Rate SS/hr]	IPR (NOx [mg/L 3.0 O ₄ -P Maion AA3 I Release [mg/ 6.	/min] D7 Ex ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr] 37	O ₄ -P A	SNPP 1.53 g. ANA g. ANA g. ANA g. SA g. ANA	NO ₃ -No ₃ -No ₄ -Po ₄ -
[days 6.38 Highest . Conco	77 77 77 77 77 77 77 7	73 71 73 59 71 73 59 67 55 62 51 45 44 42 40 85 60 85 60 8 PUR (PO ₄ Slo [mg/L] 7. A 6 NOx-N mg/L] 0.30 A4 Specific PO A4 Specific PO	[mg/L] 1.96 P Uptake pe) L/hr] [r] 10 A1 PO ₄ -P Re Slope [mg/L/hr] 13.48 O ₄ -P AA5 F Release	SPUR ng/g MLVS: 3.53 elease A/	lin AER pH [unitless] 6.64 Clarifier II S/hr] [mg 0.0 A1 Specific PO Release Rate mg/g MLVSS/h 12.36 A5 Specific PO ₄	AUR (NH-Ng/L) OC NH4-Ng/L) OOO Rel Rel Relea	H4-N Slope) 3/L/hr] 2.43 Clarifier ID [mg] 16.4 AA2 PO ₄ -P elease Slope [mg/L/hr] 9.02	SAUR [mg/g MLVS: 1.21 Claric C NOx-N Co y/L] 540 AA2 Specific Release [mg/g MLV 8.27	er IDC P ncentrati [mg/L] 0.0 PO ₄ -P Rate SS/hr]	IPR (NOx [mg/L] 3.0	/min] D7 xx ANA P Release [mg/L] 21.10 PO ₄ -P e Slope L/hr] 37 (AA1) Si Upta	AA3 S Rel	SNPI 1.53 g. ANA Concen [mg 0.3: pecific rease R MLVS 5.84 (AA P.	/SS/hr] A NO ₃ -N tration /L] 56 PO ₄ -P cate SS/hr] A1) PO ₄ / (AA1)

# Chem Used	FeCl3 Hrs U	sed FeCl3	3 Addition	Active Bios Influent Fl		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]	[r	ng/L]	[MGD]		[M	GD]	[MGD]	[MGD]
0	0.000	C	0.000	19.77		17.	66	1	1.995	0
ANA MLTSS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	'L]	[%]	[mg/L]
1380	79	1090	2640	74	1	954	272	0	74	2013

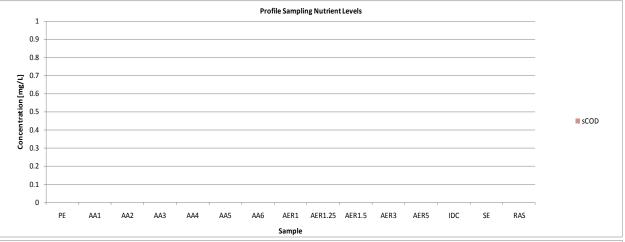


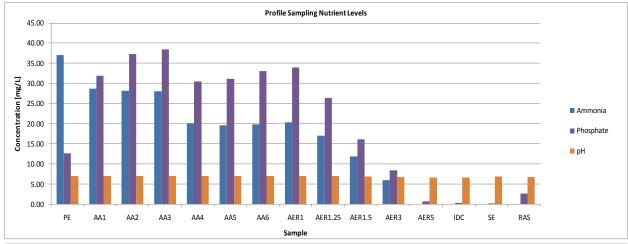


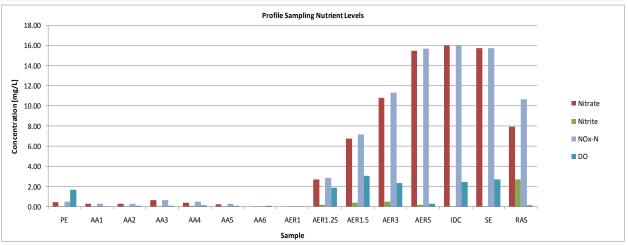


		oranom ram	5, Clarifie	er o														
1		HRT along BI	NR															
		Process (w/o											NOx-N by		CH Phos	phate by HAC		
Date	Sample Tim		Sampl	e Number	Reactor Ty	pe Amm	nonia by HA			by HACH TN				ite HACH		TNT	DO	pН
dd-mmm-yy	h:mm	hr		DE .			mg/L NH3	3-N	r	ng/L NO3-N	mg/L N			NOx-N	- 1	ng/L PO4-P	[mg/L]	[unitless]
18-Sep-09	8:20	0	_	PE	- ^		37.00			0.46	0.0			0.49		12.73	1.67	7.02
18-Sep-09 18-Sep-09	8:34 9:04	0.83 1.68	_	4A1 4A2	Anaerobi Anaerobi		28.70 28.20			0.30	0.0			0.32		31.98 37.36	0.07	7.03 7.03
18-Sep-09	9:34	2.53		4A3	Anaerobi		28.10			0.65	0.0			0.66		38.51	0.00	7.03
18-Sep-09	10:04	3.37	_	4A4	Anaerobi		20.10			0.40	0.0			0.49		30.51	0.15	7.01
18-Sep-09	10:19	4.21	_	4A5	Anaerobi		19.60			0.26	0.0			0.29		31.16	0.11	7.01
18-Sep-09	10:34	5.05	_	AA6	Anaerobi	С	19.85			0.00	0.0			0.01		33.12	0.12	6.98
18-Sep-09	10:49	0.00	A	ER1	Aerobic		20.40			0.00	0.0	01	(0.01		33.94	0.07	6.99
18-Sep-09	10:58	0.33	AE	R1.25	Aerobic		17.10			2.70	0.1	19	2	2.89		26.43	1.89	6.99
18-Sep-09	11:06	0.65		R1.5	Aerobic		11.95			6.75	0.4			7.17		16.15	3.07	6.97
18-Sep-09	11:51	3.26	_	ER3	Aerobic		6.06			10.82	0.5			1.33		8.48	2.35	6.79
18-Sep-09	12:51	6.52		ER5	Aerobic		0.00			15.48	0.2			5.69		0.71	0.31	6.65
18-Sep-09	12:59			IDC	-		0.00			16.02	0.0			6.06		0.35	2.47	6.71
18-Sep-09	15:59		_	SE	-		0.00			15.74	0.0			5.76		0.26	2.71	6.91
18-Sep-09	16:14		F	RAS	-		0.00			7.96	2.7	73	1	0.69		2.69	0.15	6.73
*NOTE: Thes	se values are	<1 mg/L NH ₄ -N																
*NOTE: Thes	se values are	<0.23 mg/L NO ₃ -	N															
*NOTE: Thes	se values are	<0.5 mg/L PO₄-P)															
QA/	QC S	ampl	e :															
		365.1		F	PA 3	50.1	E	PA	353	2 1	EPA 3:	53.2	FE	PΔ 3	53.2	EP	A 35	3.2
	1.21.2.5	303.1		1 -	1113	50.1	+ -		333		31 7 1 3.	33.2		110	33.2		11 30	.5.2
s	COD	by CE	≣L		O-PO	O_4		NH.	4-N		$NO_{2,3}$	-N	1	NO_3	-N	1	1O ₂ -	Ν
		g/L]		1	mg/	L		mg	/L		mg/l			mg/	L		mg/I	
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ActualS	SSRT AG	0 0 0 0 0 0 0 0 0	S Aver	age AE	ER DO	Min AE	ER pH	AUR	(NH4	-N Slope)	s	AUR	NF	PR (NO	x-N Sld	pe)	SNP	γ.
Actual S		0 0 0 0 0 0 0 0 0	3 Aver	age AE		Min AE			(NH4			AUR MLVSS/H		•	x-N Slo	. ,		R (SS/hr]
	s]	O O O O O O O O O O O O O O O O O O O	3 Aver		.]		less]			L/hr]	[mg/g N			[mg/		. ,		'SS/hr]
[days	3	O O O O O O O O O O O O O O O O O O O		[mg/L 1.54	.]	[unitl	less]		[mg/l	L/hr]	[mg/g N	//LVSS/h	nr]	[mg/	L/min] .98	[mg	/g ML\ 1.48	'SS/hr]
[days	s]	O O O O O O O O O O O O O O O O O O O		[mg/L 1.54	.]	[unitl	less]		[mg/l	L/hr]	[mg/g N	//LVSS/h		[mg/	L/min]	[mg	/g ML\ 1.48	'SS/hr]
[days 4.43 Highest	3	O O O O O O O O O O O O O O O O O O O		[mg/L 1.54	.]	[unitl	less] 65		[mg/l	⊥/hr] 33	[mg/g N	MLVSS/h 1.40 Clarifier	nr]	[mg/ 2 0 ₄ -P N	L/min] .98	PO ₄ -P	/g ML\ 1.48	(SS/hr]
[days 4.43 Highest A	AER NO ₂ ·	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upta Slope)	[mg/L 1.54 ake	.] SPUF	[unitl 6.6	less] 65 Clarifie	r IDC N	[mg/l	_/hr] 33 Clarifier II	[mg/g NOx-N	ALVSS/h 1.40 Clarifier Cond	IDC PC	[mg/ 2 0 ₄ -P N	L/min] .98 lax ANA Relea	PO ₄ -P	1.48 1.48 vg. ANA Concer	(SS/hr] NO ₃ -N tration
[days 4.43 Highest A Conce	AER NO ₂ · entration	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upta Blope)	[mg/L 1.54 ake	SPUF	[unitl 6.6 R [SS/hr]	less] 65 Clarifie	r IDC N	[mg/l	_/hr] 33 Clarifier II	[mg/g NOx-N g/L]	ALVSS/h 1.40 Clarifier Cond	nr] · IDC PC centration mg/L]	[mg/ 2 0 ₄ -P N	L/min] .98 lax ANA Relea [mg/	PO ₄ -P /	/g ML\ 1.48 .vg. ANA Concer [mg	'SS/hr] NO ₃ -N tration
[days 4.43 Highest A Conce	AER NO ₂ ·	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upta Slope)	[mg/L 1.54 ake	.] SPUF	[unitl 6.6 R [SS/hr]	less] 65 Clarifie	r IDC N	[mg/l	_/hr] 33 Clarifier II	[mg/g NOx-N	ALVSS/h 1.40 Clarifier Cond	IDC PC	[mg/ 2 0 ₄ -P N	L/min] .98 lax ANA Relea	PO ₄ -P /	1.48 1.48 vg. ANA Concer	'SS/hr] NO ₃ -N tration
[days 4.43 Highest A Conce	AER NO ₂ · entration	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upta Blope) g/L/hr] 6.48	[mg/L 1.54 ake	SPUF ng/g MLV 3.22	[unitl 6.6 R [SS/hr]	less] 65 Clarifie	r IDC N mg/L]	[mg/l 2.8 lH ₄ -N	_/hr] 33 Clarifier II [m 16.0	[mg/g N DC NOx-N g/L]	Clarifier Conc	IDC PC centration mg/L]	[mg/ 2 2 0 ₄ -P M	L/min] .98 lax ANA Relea [mg/ 25.7	PO ₄ -P A	/g ML\ 1.48 wg. ANA Concer [mg	(SS/hr] NO ₃ -N tration /L]
[days 4.43 Highest . Conce	AER NO ₂ : entration mg/L]	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upta Slope) g/L/hr] 6.48	[mg/L 1.54 ake [n	SPUF ng/g MLV 3.22	[unitl 6.6 R SS/hr]	Clarifie	r IDC N mg/L] 0.0000	[mg/I 2.8 IH ₄ -N	Clarifier II [m 16.42 PO ₄ -P	[mg/g N DC NOx-N g/L] 0620	Clarifier Conc	r IDC PC centration mg/L] 0.3	[mg/ 2 0 ₄ -P M n	L/min] .98 lax ANA Relea [mg/ 25.7	PO ₄ -P /	/g ML\ 1.48 avg. AN/ Concer [mg 0.3	(SS/hr] NO ₃ -N tration /L] 19 PO ₄ -P
[days 4.43 Highest According to the control of the	AER NO ₂ - entration mg/L] 0.51 Ox-N AA	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upt: Slope) g/L/hr] 6.48	[mg/L 1.54 ake [n	SPUF ng/g MLV 3.22	[unitl 6.6 R SS/hr] AA1 Sp Rele	Clarifie	r IDC N mg/L] 0.0000 PO ₄ -P	[mg/l 2.8 IH ₄ -N	Clarifier II [m 16. A2 PO ₄ -P ease Slop	DC NOx-N g/L] 0620 AA2 S e Rel	ALVSS/h 1.40 Clarifier Conc [i	mg/L] 0.3 PO ₄ -P	[mg/ 2 0 ₄ -P M n AA3	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop	PO ₄ -P // see L] 8 AA3 S	/g ML\ 1.48 vg. ANA Concer [mg 0.3	(SS/hr] NO ₃ -N tration /L] 19 PO ₄ -P tate
[days 4.43 Highest Conc [n]	AER NO ₂ - entration mg/L] 0.51 Dx-N AA	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upti Slope) g/L/hr] 6.48 AA1 PC	[mg/L 1.54 ake [n 0 ₄ -P Re Slope	SPUF ng/g MLV 3.22	[unitl 6.6 R SS/hr] AA1 Sp Rele	Clarifie Clarifie (pecific Fease Ra	r IDC N mg/L] 0.0000 PO ₄ -P	[mg/l 2.8 IH ₄ -N	Clarifier II [m 16.4 A2 PO ₄ -P ease Slop mg/L/hr]	DC NOx-N g/L] 0620 AA2 S e Rel	ALVSS/H 1.40 Clarifier Conc [i pecific F ease Ra MLVSS	mg/L] 0.3 PO ₄ -P	[mg/ 2 0 ₄ -P M n AA3 Releas	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr]	PO ₄ -P // see L] 8 AA3 S	1.48 Avg. ANA Concer [mg 0.3 Specific lease F	(SS/hr] NO ₃ -N tration /L] 19 PO ₄ -P tate
[days 4.43 Highest According to the control of the	AER NO ₂ - entration mg/L] 0.51 Dx-N AA	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upti Slope) g/L/hr] 6.48 AA1 PC	[mg/L 1.54 ake [n	SPUF ng/g MLV 3.22	[unitl 6.6 R SS/hr] AA1 Sp Rele	Clarifie	r IDC N mg/L] 0.0000 PO ₄ -P	[mg/l 2.8 IH ₄ -N	Clarifier II [m 16. A2 PO ₄ -P ease Slop	DC NOx-N g/L] 0620 AA2 S e Rel	ALVSS/h 1.40 Clarifier Conc [i	mg/L] 0.3 PO ₄ -P	[mg/ 2 0 ₄ -P M n AA3 Releas	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop	PO ₄ -P // see L] 8 AA3 S	yg ML\ 1.48 vg. ANA Concer [mg 0.3 Specific lease F g MLVS 7.09	/SS/hr] NO ₃ -N tration /L] 19 PO ₄ -P tate SS/hr]
[days 4.43 Highest Conco	AER NO ₂ : entration mg/L] 0.51 Dx-N AA	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upts Slope) g/L/hr] 6.48 AA1 PC	[mg/L 1.54 ake [n 0 ₄ -P Re Slope ng/L/hr]	SPUF	[unitl 6.6 R SS/hr] AA1 Sp Rele [mg/g	Clarifie Clarifie Clarifie MLVS: 11.40	r IDC N mg/L] 0.0000 PO ₄ -P ate S/hr]	[mg/l 2.8 lH ₄ -N	Clarifier II [m] 16. A2 PO ₄ -P ease Slop mg/L/hr] 11.88	[mg/g N 2 2 2 2 2 2 2 2 2	Clarifier Conc pecific F ease Ra MLVSS 11.63	r IDC PC centration mg/L] 0.3 PO ₄ -P	[mg/ 2 0 ₄ -P M n AA3 Releas	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr]	PO ₄ -P // see L] 8 AA3 S	yg ML\ 1.48 vg. ANA Concer [mg 0.3 Specific lease F g MLVS 7.09	(SS/hr] NO ₃ -N tration /L] 19 PO ₄ -P tate
[days 4.43 Highest Conc [n]	AER NO ₂ : entration mg/L] 0.51 Dx-N AA	O O O O O O O O O O O O O O O O O O O	O ₄ -P Upts Slope) g/L/hr] 6.48 AA1 PC	[mg/L 1.54 ake [n 0 ₄ -P Re Slope	SPUF	[unitl 6.6 R SS/hr] AA1 Sp Rele [mg/g	Clarifie Clarifie (pecific Fease Ra	r IDC N mg/L] 0.0000 PO ₄ -P ate S/hr]	[mg/l 2.8 lH ₄ -N	Clarifier II [m] 16. A2 PO ₄ -P ease Slop mg/L/hr] 11.88	DC NOx-N g/L] 0620 AA2 S e Rel	Clarifier Conc pecific F ease Ra MLVSS 11.63	r IDC PO centration mg/L] 0.3 PO ₄ -P tite S/hr]	[mg/ 2 0 ₄ -P M n AA3 Releas [mg	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr] 7.24 (AA1)	PO ₄ -P // see L] 8 AA3 S	Jeg MLV 1.48 Avg. ANA Concer [mg 0.3 Specific lease F g MLV5 7.09 (AA) (AB)	/SS/hr] A NO ₃ -N tration /L] 19 PO ₄ -P tate SS/hr] A1) PO ₄ - / (AA1)
[days 4.43 Highest Conco	AER NO ₂ · entration mg/L] 0.51 0.51 0.4-P A	O O O O O O O O O O O O O O O O O O O	D ₄ -P Upts slope) g/L/hr] 6.48 AA1 PC	[mg/L 1.54 ake [n 0 ₄ -P Re Slope ng/L/hr] 11.64	SPUF	[unitl 6.6 R SS/hr] AA1 Sp Rele [mg/g	Clarifie Clarifie Clarifie MLVS: 11.40	r IDC N mg/L] 0.0000 PO ₄ -P ate S/hr]	[mg/l 2.8 IH ₄ -N Rele [r	Clarifier II [m] 16. A2 PO ₄ -P ease Slop mg/L/hr] 11.88	[mg/g N 2 2 2 2 2 2 2 2 2	Clarifier Conc [t] Pecific F ease Ra g MLVSS 11.63	r IDC PO centration mg/L] 0.3 PO ₄ -P tite S/hr]	[mg/ 2 0 ₄ -P M n AA3 Releas	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr] 7.24 (AA1)	PO ₄ -P // sse L] 8 AA3 \$ Re [mg/	Jeg MLV 1.48 Avg. ANA Concer [mg 0.3 Specific lease F g MLV5 7.09 (AA) (AB)	(SS/hr] NO ₃ -N tration (L] 19 PO ₄ -P tate SS/hr]
Idays 4.43 Highest Aconc In In In In AA4 NC Img/L 0.49 AA4 PC Release	AER NO ₂ - entration ng/L] 0.51 Dx-N AA L] 0 O4-P A	O O O O O O O O O O O O O O O O O O O	D ₄ -P Upticlope) g/L/hr] 6.48 AA1 PC gr	[mg/L 1.54 ake [n 0 ₄ -P Re Slope ng/L/hr] 11.64 AA5 F Release	SPUF ng/g MLV 3.22 elease PO ₄ -P e Slope	[unitl 6.6 SS/hr] AA1 Sp Rele [mg/g	Clarifie Clarif	r IDC N mg/L] 0.0000 PO ₄ -P ate S/hr]	AA6 Release	Clarifier II [m 16. 22 PO ₄ -P 22 PO ₄ -P 23 PO ₄ -P 24 PO ₄ -P 25 PO ₄ -P 26 Slope	DC NOx-N g/L 0620 AA2 S e Rel [mg/g N	Clarifier Conc [i pecific F ease Ra g MLVSS 11.63 fic PO ₄ -P e Rate	PO ₄ -P tte (AA1 Uptak)	Img/ 2 2 04-P M n AA3 Releas [mg 7	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr] 7.24 (AA1) Ur	PO ₄ -P // sse L] 8 AA3 Se Re [mg/	1.48 Avg. ANA Concer [mg 0.3 Epecific lease F g MLVs 7.09 (A) OD P	(SS/hr] A NO ₃ -N tration (L] 19 PO ₄ -P tate SS/hr] A1) PO ₄ - ((AA1)
[days 4.43 Highest . Conco	AER NO ₂ : entration ng/L] 0.51 Dx-N AA L] 0 O ₄ -P A Slope	O O O O O O O O O O O O O O O O O O O	D ₄ -P Upticlope) g/L/hr] 6.48 AA1 PC gr	[mg/L 1.54 ake [n 0 ₄ -P Re Slope ng/L/hr] 11.64	SPUF ng/g MLV 3.22 elease PO ₄ -P e Slope L/hr]	[unitl 6.6 SS/hr] AA1 Sp Rele [mg/g	Clarifie Clarifie Clarifie MLVS: 11.40	r IDC N mg/L] 0.0000 PO ₄ -P ate S/hr]	Emg/II 2.€ AAA Release Emg/II AAA6 Release Emg/II Release	Clarifier II [m 16. A2 PO ₄ -P ease Slop mg/L/hr] 11.88 PO ₄ -P	[mg/g N OC NOx-N g/L] 0620 AA2 S e Rel [mg/g	Clarifier Conc [I pecific F ease Ra 11.63 fic PO ₄ -P Rate VSS/hr]	IDC PC centration mg/L] 0.3 PO ₄ -P tite S/hr] (AA1 Uptak	[mg/ 2 0 ₄ -P M n AA3 Releas [mg	L/min] .98 lax ANA Relea [mg/ 25.7 PO ₄ -P se Slop g/L/hr] 7.24 (AA1) Ur	PO ₄ -P // se L] 8 AA3 \$ e Re [mg/	1.48 Avg. ANA Concer [mg 0.3 Epecific lease F g MLVs 7.09 (A) OD P	/SS/hr] A NO ₃ -N tration /L] 19 PO ₄ -P tate SS/hr] A1) PO ₄ - / (AA1)

# Chem Used	FeCl3 Hrs U	sed	FeCl3	Addition	Primary Cla Influent Flo		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]		[n	ng/L]	[MGD]		[M0	GD]	[1	MGD]	[MGD]
0	0.000		0	.000	20.19		18.	.03	1:	2.075	0
ANA MLTS	S ANA % VOL	ANA	MLVSS	ANX MLTSS	S ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[m	ng/L]	[mg/L]	[%]	[n	ng/L]	[mg/	[L]	[%]	[mg/L]
1380	74	1	021	2720	74	2	013	272	0	74	2013



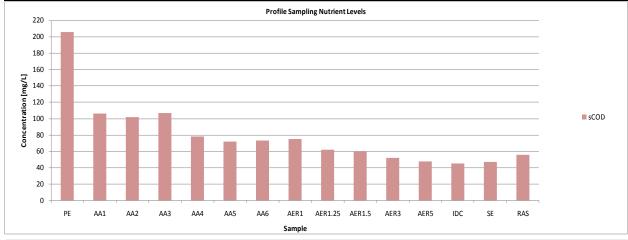


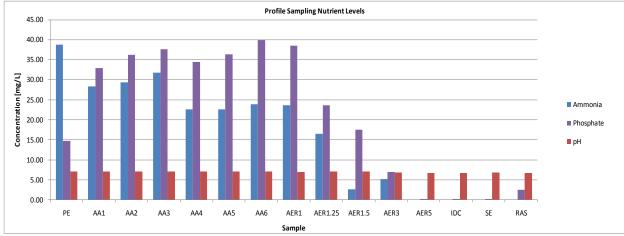


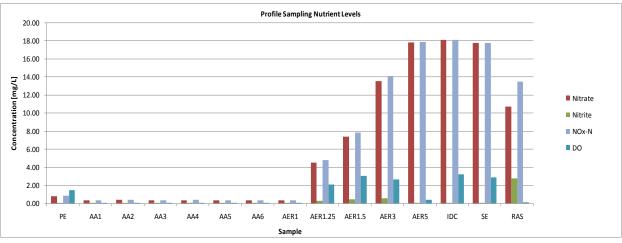
Week 10 - 9/22/09 & 9/24/09

dd-mmm-yy h:mm hr mg/L NH3-N mg/L NO3-N mg/L NO2-N mg/L NOx-N 22-Sep-09 7:03 0 PE - 38.80 0.81 0.03 0.84			
22-Sep-09 7:03 0 PE - 38.80 0.81 0.03 0.84	nosphate by HAC	CH TNT DO	pН
	mg/L PO4-P	[mg/L]	[unitless]
22 Son 00 7:40 0.97 AA4 Apparatio 29.20 0.25 0.00 0.27	14.68	1.48	7.09
22-Sep-09 7:19 0.87 AA1 Anaerobic 28.30 0.36 0.02 0.37	32.96	0.08	7.08
22-Sep-09 7:49 1.73 AA2 Anaerobic 29.40 0.38 0.01 0.39	36.22	0.08	7.09
22-Sep-09 8:19 2.60 AA3 Anaerobic 31.80 0.36 0.01 0.38	37.69	0.07	7.09
22-Sep-09 8:49 3.46 AA4 Anaerobic 22.60 0.36 0.03 0.39	34.43	0.08	7.04
22-Sep-09 9:04 4.33 AA5 Anaerobic 22.65 0.34 0.02 0.35	36.38	0.08	7.04
22-Sep-09 9:19 5.19 AA6 Anaerobic 23.85 0.34 0.02 0.36	39.97	0.07	7.03
22-Sep-09 9:34 0.00 AER1 Aerobic 23.65 0.34 0.02 0.36	38.51	0.06	7.01
22-Sep-09 (;42 0.34 AER1.25 Aerobic 16.50 4.51 0.31 4.82	23.58	2.07	7.05
22-Sep-09 9:50 0.67 AER1.5 Aerobic 2.58 7.38 0.46 7.84	17.54	3.05	7.03
22-Sep-09 10:35 3.35 AER3 Aerobic 5.18 13.52 0.59 14.11	6.98	2.68	6.83
22-Sep-09 11:42 6.71 AER5 Aerobic 0.00 17.82 0.05 17.87	0.23	0.40	6.7
22-Sep-09 11:47 IDC - 0.00 18.08 0.02 18.10	0.18	3.25	6.74
22-Sep-09 14:47 SE - 0.00 17.74 0.02 17.76	0.19	2.87	6.89
22-Sep-09 15:05 RAS - 0.00 10.74 2.76 13.50	2.45	0.12	6.73
*NOTE: These values are <1 mg/L NH ₄ -N			
QA/QC Sample: SE			
EPA 365.1 EPA 350.1 EPA 353.2 EPA 353.2 EPA 353	3.2 E	EPA 35	53.2
sCOD by CEL O-PO ₄ NH ₄ -N NO _{2,3} -N NO ₃ -N	4	NO_2 -	Z
$ [mg/L] \hspace{1cm} mg/L \hspace{1cm} mg/L \hspace{1cm} mg/L \hspace{1cm} mg/L $,	mg/I	,
206			
106			
102			
107			
78			
72			
73			
75			
62			
60			
52			
48			
45	<u> </u>	0.02	
45 47 0.142 <0.20 18.4 18.4	工	0.02	
45		0.02	
45 47 0.142 <0.20 18.4 18.4		0.02	
45 47 0.142 <0.20 18.4 18.4	T	0.02	
45 47 0.142 <0.20 18.4 18.4 56	J Slone)		
45 47 0.142 <0.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N State of the state		SNPF	₹
45 47 O.142 <o.20 (nh4-n="" (nox-n="" 18.4="" 56="" [days]="" [mg="" [unitless]="" actual="" aer="" aerobic="" aur="" average="" do="" g="" hr]="" l="" l]="" min="" min<="" mlss="" mlvss="" npr="" ph="" s="" saur="" slope)="" srt="" td=""><td>nin] [</td><td>SNPF mg/g MLV</td><td>R (SS/hr]</td></o.20>	nin] [SNPF mg/g MLV	R (SS/hr]
45 47 0.142 <0.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N State of the state	nin] [SNPF	R 'SS/hr]
45 47 O.142 <o.20 (nh4-n="" (nox-n="" 18.4="" 56="" [days]="" [mg="" [unitless]="" actual="" aer="" aerobic="" aur="" average="" do="" g="" hr]="" l="" l]="" min="" min<="" mlss="" mlvss="" npr="" ph="" s="" saur="" slope)="" srt="" td=""><td>nin] [</td><td>SNPF mg/g MLV</td><td>R 'SS/hr]</td></o.20>	nin] [SNPF mg/g MLV	R 'SS/hr]
45 47 O.142 <-O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Sidays) [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min 5.95 3780 1.65 6.70 2.52 0.89 3.49	nin] [SNPF mg/g MLV 1.23	R [SS/hr]
45 47 O.142 <-O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Sidays) [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min 5.95 3780 1.65 6.70 2.52 0.89 3.49 Highest AER NO ₂ -N PUR (PO ₄ -P Uptake Clarifier IDC PO ₄ -P Max All Clarifier IDC	ANA PO ₄ -P	SNPF mg/g MLV 1.23 Avg. ANA	R [SS/hr]
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min 5.95 3780 1.65 6.70 2.52 0.89 3.49 Highest AER NO ₂ -N PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Re	ANA PO ₄ -P	SNPF mg/g MLV 1.23 Avg. ANA Concen	SS/hr] NO ₃ -N
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min 5.95 3780 1.65 6.70 2.52 0.89 3.49 Highest AER NO ₂ -N PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Re [mg/L] [mg/L/hr] [mg/g MLVSS/hr] [mg/L]	ANA PO ₄ -P Release [mg/L]	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg.	R [SS/hr] NO ₃ -N tration [L]
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SPUR Clarifier IDC NOx-N Concentration [mg/L] [ANA PO ₄ -P	SNPF mg/g MLV 1.23 Avg. ANA Concen	SS/hr] NO ₃ -N tration
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SPUR Clarifier IDC NOx-N Concentration [mg/L] [ANA PO ₄ -P Release [mg/L]	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg.	SS/hr] NO ₃ -N tration
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SPUR Clarifier IDC NOx-N Concentration [mg/L] [ANA PO ₄ -P Release [mg/L]	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg.	R [SS/hr] L NO ₃ -N tration /L]
45 47 O.142 O.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Sidays) [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min 5.95 3780 1.65 6.70 2.52 0.89 3.49 Highest AER NO ₂ -N PUR (PO ₄ -P Uptake SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration [mg/L] [mg/L/hr] [mg/g MLVSS/hr] [mg/L] [mg/L	ANA PO ₄ -P Release [mg/L] 25.29	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg. 0.33	SS/hr] NO ₃ -N tration L[L] 66
45 47 O.142 <0.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) [days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min Slope) 5.95 3780 1.65 6.70 2.52 0.89 3.49 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Clarifier IDC PO ₄ -P Max All Clarifier IDC NOx-N Concentration Reference Img/L] [mg/L]	ANA PO ₄ -P Release [mg/L] 25.29	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg. 0.33 A3 Specif	SS/hr] NO ₃ -N tration /L] fic PO ₄ - Rate
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45 47 O.142 <0.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N States and States	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P A Slope [r/hr] [r	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg. 0.33 A3 Specif	SS/hr] NO ₃ -N tration /L] 66 fic PO ₄ : Rate SS/hr]
45 47 O.142 <0.20 18.4 18.4 56 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N SIDER NPR (NOx-N SIDER) [days] [mg/L] [mg/	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P A Slope [r/hr] [r	SNPF mg/g MLV 1.23 Avg. ANA Concen [mg. 0.33 A3 Specif P Release mg/g MLV	SS/hr] NO ₃ -N tration /L] 66 fic PO ₄ : Rate SS/hr]
45 47 O. 142 <0.20 18.4 18.4 56	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P A Slope F /hr] [r	SNPf mg/g MLV 1.23 Avg. ANA Concen [mg/g 0.33 A3 Specif P Release mg/g MLV 7.86	R SS/hr] A NO ₃ -N tration /L] 56 fic PO ₄ - A Rate SS/hr]
45 47 O.142 <0.20 18.4 18.4 56 56 SAUR NPR (NOx-N Section of the content of the c	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P Al) Specific	SNPF Img/g MLV 1.23 Avg. ANA Concen [mg/g 0.33 A3 Specif P Release mg/g MLV 7.86 (AA1)	R SS/hr] A NO ₃ -N tration /L] 56 Sic PO ₄ - PO ₄ -
Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slop	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P All Slope F All Specific Uptake Rate	SNPF Img/g MLV 1.23 Avg. ANA Concen [mg/g 0.38 A3 Specifi P Release mg/g MLV 7.86 (AA1) P/C0	R (SS/hr] (NO ₃ -N) tration (L) (56 (Rate SS/hr) (PO ₄ - DD
Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) NPR (NOx-N Slope) SAUR NPR (NOx-N Slope) NPR (NOx-N S	ANA PO ₄ -P Release [mg/L] 25.29 O ₄ -P Al) Specific	SNPF Img/g MLV 1.23 Avg. ANA Concen [mg/g 0.38 A3 Specifi P Release mg/g MLV 7.86 (AA1) P/C0	R (SS/hr] (NO ₃ -N) tration (L) (56 (Rate (SS/hr) (PO ₄ - DD) (mg]

# Chem Used	FeCl3 Hrs U	sed F	FeCl3	Addition	Active Bios Influent Fl	-	ARCY	′ Flow	CRO	CY Flow	NRCY Flow
[lbs]	[hr]		[m	ng/L]	[MGD]		[M0	GD]	[1	MGD]	[MGD]
0	0.000		0.	.000	19.30		17.	.81	1	1.191	0
	S ANA % VOL			ANX MLTSS			MLVSS				DL AER MLVSS
[mg/L]	[%]	[mg/	/L]	[mg/L]	[%]	[m	ng/L]	[mg/	'L]	[%]	[mg/L]
1200	75	900	0	3780	75	2	835	378	0	75	2835



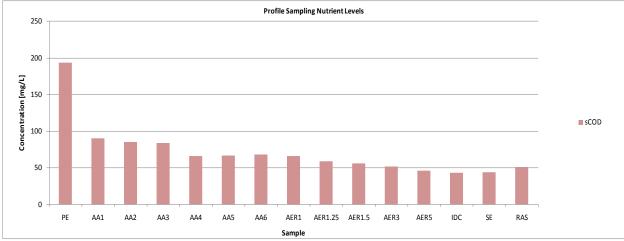


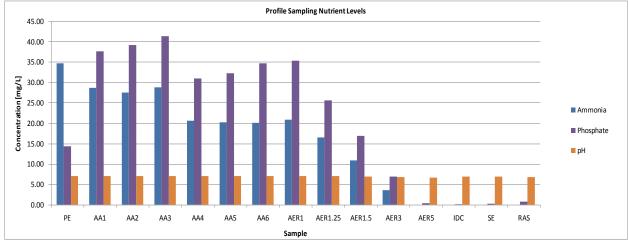


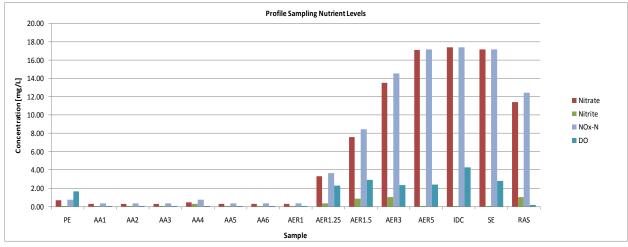
Date	Sample Time	HRT along BNR Process (w/out e Recycles)	Sample Number	Reactor Type	Ammor	nia by HACH T	NT Nitrate b	y HACH TNT	Nitrite by HACI		l by Nitrate HA Nitrite HACH	CH Phosphate	•	H DO	рН
dd-mmm-yy	h:mm	hr			m	ng/L NH3-N	mg/	L NO3-N	mg/L NO2-	N r	ng/L NOx-N		P04-P	[mg/L]	[unitless]
24-Sep-09	7:07	0	PE	-		34.70		0.69	0.03		0.73		.36	1.68	7.06
24-Sep-09	7:24	0.85	AA1	Anaerobic		28.70		0.30	0.02		0.32		.69	0.06	7.07
24-Sep-09	7:54	1.73	AA2	Anaerobic		27.50		0.31	0.02		0.33		.16	0.07	7.08
24-Sep-09	8:24	2.60	AA3	Anaerobic		28.90		0.31	0.02		0.33		.44	0.06	7.06
24-Sep-09	8:54	3.46	AA4	Anaerobic		20.60		0.48	0.27		0.75		.00	0.08	7.02
24-Sep-09	9:09	4.33	AA5	Anaerobic		20.25		0.30	0.02		0.32	32		0.06	7.01
24-Sep-09 24-Sep-09	9:24 9:39	5.19 0.00	AA6 AER1	Anaerobic		20.10		0.32	0.02		0.33	35	.75	0.06	7.00
24-Sep-09 24-Sep-09	9:48	0.00	AER1.25	Aerobic Aerobic		16.55		3.30	0.02		3.66		.70	2.28	7.02 7.02
24-Sep-09	9:56	0.67	AER1.5	Aerobic		10.85		7.58	0.84		8.42		.89	2.90	6.97
24-Sep-09	10:41	3.35	AER3	Aerobic		3.56		13.52	1.03		14.55		95	2.35	6.75
24-Sep-09	11:41	6.71	AER5	Aerobic		0.00		17.14	0.03		17.17		38	2.40	6.72
24-Sep-09	11:52	0	IDC	-	-	0.00		17.38	0.02		17.40	0.1		4.26	6.88
24-Sep-09	14:52		SE			0.00		17.16	0.02		17.18	0.1		2.77	6.98
				-	-										
24-Sep-09	15:18		RAS	-		0.00		11.42	1.03		12.45	0.	73	0.20	6.79
		<1 mg/L NH ₄ -N													
QA/	QC S	ample		SE											
	EP	A 365.1		EPA	350.	1 E	PA 35	3.2	EPA 3	53.2	EPA	353.2	E	PA 35	3.2
:	sCO	by CE	L	O-I	PO_4		NH ₄ -I	Ν	$NO_{2,3}$	3-N	NO	D_3 -N]	NO_2 -I	N
	[r	ng/L]		mg	z/L		mg/L	_	mg/	L	m	g/L		mg/L	_
		194													
		90													
		\circ													
		85													
		84													
		84 66													
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		84 66 67 68 66 59													
		84 66 67 68 66 59 56													
		84 66 67 68 66 59													
		84 66 67 68 66 59 56 52 46 43					<0.20) T	17	7	1	7.7		0.02	
		84 66 67 68 66 59 56 52 46 43					<0.20)	17.	7	1	7.7		0.02	
	1	84 66 67 68 66 59 56 52 46 43					<0.20)	17.	7	1	7.7		0.02	
		84 66 67 68 66 59 56 52 46 43					<0.20)	17.	7	1	7.7		0.02	
Actual	SRT A	84 66 67 68 66 59 56 52 46 43 44 51	Average A	AER DO	Min AE	ER pH	<0.20	•		7		7.7	e)	O.O2	2
Actual		84 66 67 68 66 59 56 52 46 43 44 51	Ŭ				AUR (NH4	1-N Slope	e) S/	\UR	NPR (N	NOx-N Slope		SNPF	
[days	s]	84 66 67 68 66 59 56 52 46 43 44 51	[mg/	/L]	[unitle	ess]	AUR (NH4	1-N Slope L/hr]	e) S/	AUR ILVSS/hr]	NPR (N	NOx-N Slope		SNPF g/g MLV	SS/hr]
	s]	84 66 67 68 66 59 56 52 46 43 44 51	Ŭ	/L]		ess]	AUR (NH4	1-N Slope L/hr]	e) S/	\UR	NPR (N	NOx-N Slope		SNPF	SS/hr]
[days	s]	84 66 67 68 66 59 56 52 46 43 44 51	[mg/	/L]	[unitle	ess]	AUR (NH4	1-N Slope L/hr]	e) S/	AUR ILVSS/hr]	NPR (I	NOx-N Slope	[m	SNPF g/g MLV	SS/hr]
[days	s])	84 66 67 68 66 59 56 52 46 43 44 51	[mg/	/L]	[unitle	ess]	AUR (NH4 [mg/ 2.	I-N Slope L/hr] 79	e) S/	AUR ILVSS/hr] .46	NPR (N [m	NOx-N Slope ng/L/min] 3.73	[m	SNPF g/g MLV 1.95	SS/hr] NO ₃ -N
[days 4.79 Highest Cond	AER NO ₂	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480	[mg, 2.0 -P Uptake pe)	(L] 0 SPUF	[unitl 6.7	ess] 72 Clarifier II	AUR (NH4 [mg/ 2.* DC NH4-N	I-N Slope L/hr] 79 Clarifier	(mg/g N IDC NOx-N	AUR ILVSS/hr] .46 Clarifier IE Concer	NPR (N	NOx-N Slope ig/L/min] 3.73 Max ANA PO Release	[m	SNPF g/g MLV 1.95 Avg. ANA Concent	SS/hr] NO ₃ -N
[days 4.79 Highest Conc	AER NO ₂ entration	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 -N PUR (PO, Sid	[mg, 2.0 -P Uptake pe) L/hr]	(L] 0 SPUF [mg/g MLV	[unitl 6.7	ess] 72 Clarifier II	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L]	I-N Slope L/hr] 79 Clarifier	(mg/g N (mg/g N 1 IDC NOx-N mg/L]	AUR ILVSS/hr] .46 Clarifier IE Concer	NPR (N [m	NOx-N Slope ng/L/min] 3.73 Max ANA PO Release [mg/L]	[m	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/	SS/hr] NO ₃ -N tration
[days 4.79 Highest Conc	AER NO ₂	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 -N PUR (PO, Sid	[mg, 2.0 -P Uptake pe)	(L] 0 SPUF	[unitl 6.7	ess] 72 Clarifier II	AUR (NH4 [mg/ 2.* DC NH4-N	I-N Slope L/hr] 79 Clarifier	(mg/g N IDC NOx-N	AUR ILVSS/hr] .46 Clarifier IE Concer	NPR (N	NOx-N Slope ig/L/min] 3.73 Max ANA PO Release	[m	SNPF g/g MLV 1.95 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 4.79 Highest Conc	AER NO ₂ entration	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 -N PUR (PO, Sid	[mg, 2.0 -P Uptake pe) L/hr]	SPUF [mg/g MLV 3.66	[unitle 6.7 R SS/hr]	ess] 72 Clarifier II	AUR (NH4 [mg/ 2.* DC NH ₄ -N g/L] 000	I-N Slope L/hr] 79 Clarifier	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR ILVSS/hr] .46 Clarifier IE Concer	NPR (NPR (NPR (NPR (NPR (NPR (NPR (NPR (NOx-N Slope ng/L/min] 3.73 Max ANA PO Release [mg/L]	[m	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/	SS/hr] NO ₃ -N tration L]
[days 4.79 Highest Conc	AER NO ₂ entration mg/L]	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 -N PUR (PO, Sid	[mg, 2.0 2.0 -P Uptake pe) L/hr] 99	SPUF [mg/g MLV 3.66	[unitle 6.7 R SS/hr]	ess] 72 Clarifier II [mg 0.0	AUR (NH4 [mg/ 2.* DC NH ₄ -N g/L] 000 P AA2	I-N Slope L/hr] 79 Clarifier [r 17	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0.	NPR (N [m]	NOx-N Slope ng/L/min] 3.73 Max ANA PO Release [mg/L] 27.08	[m]	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33	NO ₃ -N tration L]
[days 4.79 Highest Conc	AER NO ₂ entration mg/L] 1.03	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 	[mg. 2.0 2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI	SPUF [mg/g MLV 3.66 -P AA ope	[unitlender of the content of the co	Clarifier II [my 0.0 cific PO ₄ -late Rate	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L] 000 P AA2 Relea	I-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P	(a) SA [mg/g N 1 IDC NOx-N mg/L] 7.3960 AA2 Sp Rele	AUR ILVSS/hr] .46 Clarifier IC Concer [mg 0. ecific PO. ase Rate	NPR (N [m]	MOx-N Slope g/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope	[m D ₄ -P	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 3 Specific	NO ₃ -N tration L] B7 C PO ₄ - Rate
[day: 4.79 Highest Conc	AER NO ₂ entration mg/L] 1.03 Dx-N A/	84 66 67 68 66 59 56 52 46 43 44 51 2480 -N PUR (PO. Sic [mg/L] 2480 -R PUR (PO. Sic	[mg, 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0 2.0	SPUF [mg/g MLV 3.66 -P AA ope	[unitle 6.7 R SS/hr] 1 Spec Release	Clarifier II [m 0.0 clific PO ₄ -le Rate LVSS/hr]	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L] 000 P AA2 Relea [m	I-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P ase Slope	(a) SA [mg/g N 1 IDC NOx-N mg/L] 7.3960 AA2 Sp Rele [mg/g	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0. ecific PO. ase Rate	NPR (N [m]	MOx-N Slope g/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope mg/L/hr]	[m D ₄ -P	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 Specific Release	NO ₃ -N tration L] B7 C PO ₄ - Rate
[days 4.79 Highest Conc	AER NO ₂ entration mg/L] 1.03 Dx-N A/	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 	[mg. 2.0 2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI	SPUF [mg/g MLV 3.66 -P AA ope	[unitle 6.7 R SS/hr] 1 Spec Release	Clarifier II [my 0.0 cific PO ₄ -late Rate	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L] 000 P AA2 Relea [m	I-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P	(a) SA [mg/g N 1 IDC NOx-N mg/L] 7.3960 AA2 Sp Rele [mg/g	AUR ILVSS/hr] .46 Clarifier IC Concer [mg 0. ecific PO. ase Rate	NPR (N [m]	MOx-N Slope g/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope	[m D ₄ -P	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 3 Specific	NO ₃ -N tration L] B7 C PO ₄ - Rate
[day: 4.79 Highest Conc	AER NO ₂ entration mg/L] 1.03	84 66 67 68 66 59 56 52 46 43 44 51 2480 -N PUR (PO. Sic [mg/L] 2480 -R PUR (PO. Sic	[mg/ 2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI [mg/L/hr]	SPUF [mg/g MLV 3.66 -P AA ope] [n	[unitle 6.7 SS/hr] 1 Spec Releas ng/g Ml	Clarifier II [m 0.0 clific PO ₄ -le Rate LVSS/hr]	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L] 000 P AA2 Relea [m	1-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P ase Slope g/L/hr] 10.85	(a) SA [mg/g N 1 IDC NOx-N mg/L] 7.3960 AA2 Sp Rele [mg/g	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0. ecific PO. ase Rate MLVSS/h 11.46	NPR (N [m]	MOx-N Slope g/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope mg/L/hr]	AA3	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 Specific Release	NO ₃ -N tration L] 87 C PO ₄ - Rate SS/hr]
[day: 4.79] Highest Conc [t	AER NO ₂ entration mg/L] 1.03 Dx-N A/L] 5	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 PUR (PO, Slotant Solution of the state of the s	[mg/ 2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI [mg/L/hr] 16.19	SPUF [mg/g MLV 3.66 -P AA ope -P [ng/g MLV -P AA	[unitle 6.7 SS/hr] 1 Spec Releaseng/g Mil 17	Clarifier II [m] 0.0 clarifier PO ₄ -lee Rate LVSS/hr]	AUR (NH4 [mg/ 2.* DC NH ₄ -N g/L] 000 P AA2 Relea [m	1-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P ase Slope g/L/hr] 10.85	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0. ecific PO. ase Rate MLVSS/h 11.46	NPR (N [m] C PO ₄ -P ntration yL] 2 A-P AA Rele	MOx-N Slope ig/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope mg/L/hr] 7.82	Im D ₄ -P AA3 P Im g	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 3 Specific Release g/g MLV 8.25	NO ₃ -N tration L] 87 C PO ₄ - Rate SS/hr]
[day: 4.79 Highest Conc [t	AER NO ₂ entration mg/L] 1.03 Dx-N A/ L] 5 O ₄ -P / Slope	84 66 67 68 66 59 56 52 46 43 44 51 2480 PUR (PO. Slo [mg/L] 2480 A6 NOx-N [mg/L] 0.33	[mg/2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI [mg/L/hr] 16.19 AA5 P Release	SPUF [mg/g MLV] 3.66 -P AA ope r] [n O ₄ -P A	[unitle 6.7 R SS/hr] 1 Spec Releas ng/g Ml 17 A5 Spec Releas	Clarifier II [m, 0.0 iffic PO ₄ -lee Rate LVSS/hr] .09 iffic PO ₄ -P e Rate	AUR (NH4 [mg/ 2.* DC NH ₄ -N g/L] 000 P AA2 Release [m 1 AA6 PC Release	I-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P ase Slope g/L/hr] 10.85 D ₄ -P All	ing/g Nox-N mg/L] Zayana AA2 Spee Rele [mg/g A6 Specific Prelease Rate	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0. ecific PO. ase Rate MLVSS/h 11.46 D ₄ -P (A e Upt	NPR (NPR (NPR (NPR (NPR (NPR (NPR (NPR (MOx-N Slope g/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P ease Slope mg/L/hr] 7.82 (AA1) Spe COD Uptake	AA3 P [mg	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 8 Specific Release g/g MLV3 8.25 (AA1) P/CC	NO ₃ -N tration L] S7 c PO ₄ - Rate SS/hr] PO ₄ - OD
Iday: 4.79 Highest Conc It AA4 NO Img/ 0.79 AA4 P	AER NO ₂ entration mg/L] 1.03 Dx-N A/L] 5 O ₄ -P // Slope //hr]	84 66 67 68 66 59 56 52 46 43 44 51 erobic MLSS [mg/L] 2480 PUR (PO, Slotant Science) [mg/L] 0.33	[mg/2.0 -P Uptake pe) L/hr] 99 AA1 PO ₄ Release SI [mg/L/hr] 16.19	SPUF [mg/g MLV 3.66 -P AA ope In In	[unitle 6.7 SS/hr] 1 Spec Releas ng/g Ml 17 A5 Spec Releas mg/g Ml	Clarifier II [m] 0.0 clific PO ₄ -le Rate LVSS/hr] .09	AUR (NH4 [mg/ 2.* OC NH ₄ -N g/L] 000 P AA2 Relea [m 1	I-N Slope L/hr] 79 Clarifier [r 17 2 PO ₄ -P ase Slope g/L/hr] 10.85 D ₄ -P Slope A	ing/g N IDC NOx-N mg/L 3960 AA2 Sp Rele [mg/g A6 Specific P	AUR ILVSS/hr] .46 Clarifier IE Concer [mg 0. ecific PO. ase Rate MLVSS/h 11.46 D ₄ -P (A e Upt	NPR (N [m]	MOx-N Slope mg/L/min] 3.73 Max ANA PO Release [mg/L] 27.08 A3 PO ₄ -P pase Slope mg/L/hr] 7.82 (A41) Spe	[m] AA3 P [mg] ecific e Rate SS/hr]	SNPF g/g MLV 1.95 Avg. ANA Concent [mg/ 0.33 3 Specific Release g/g MLV3 8.25 (AA1)	NO ₃ -N tration L] 87 Rate SS/hr] PO ₄ - DD mg]

PROFILE 2 Train 6, Aeration Tank 5, Clarifier 5

# Chem Used	FeCl3 Hrs U	sed	FeCl3	Addition	Active Bios Influent Fl		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]		[n	ng/L]	[MGD]		[M¢	GD]	[1	MGD]	[MGD]
0	0.000		0	.000	19.73		17.	64	1	1.432	0
ANA MLTS	S ANA % VOL	ANA	MLVSS	ANX MLTS	S ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[n	ng/L]	[mg/L]	[%]	[n	ng/L]	[mg/	[L]	[%]	[mg/L]
1230	77	*	947	2480	77	1	910	248	0	77	1910



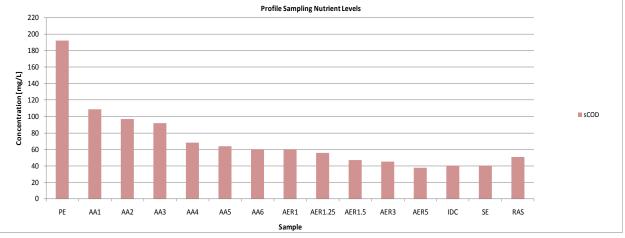


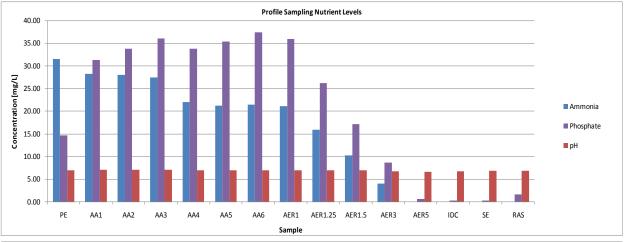


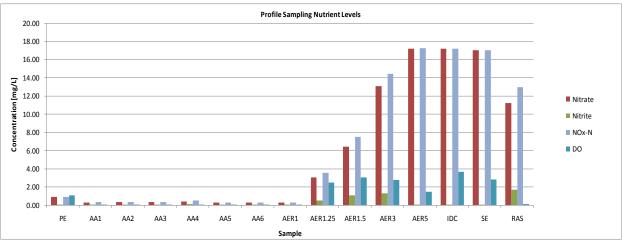
Week 11 – 9/28/09 & 9/30/09

AVIILE I	1 Train 4 Ao	ration Tank 4,	Clarifier 4													
'	i i i alli 4, AC	iadon rank 4,	olarinel 4													
		HRT along BNR														
		Process (w/out							A			y Nitrate HA				
Date	Sample Time	Recycles)	Sample Number	Reactor Type		nia by HACH TI	NI Nitrate b	y HACH TNT	Nitrite by	HACH TNT		trite HACH	Phosphate b	y HACH I	NT DO	pН
dd-mmm-yy	h:mm	hr			n	ng/L NH3-N	- v	L NO3-N		. NO2-N	mg	/L NOx-N		P04-P	[mg/L]	[unitless]
28-Sep-09	7:18	0	PE	-		31.50		0.90		0.04		0.94		.68	1.06	6.98
28-Sep-09	7:34	0.81	AA1	Anaerobic		28.30		0.32		0.02		0.34		.33	0.07	7.04
28-Sep-09	8:04	1.61	AA2	Anaerobic		28.00		0.35		0.02		0.37		.77	0.07	7.06
28-Sep-09	8:34	2.42	AA3	Anaerobic Anaerobic		27.50 22.00		0.33		0.02		0.34		.06	0.07	7.04 7.01
28-Sep-09 28-Sep-09	9:04 9:19	3.23 4.03	AA4 AA5	Anaerobic		21.20		0.39 0.27		0.13 0.03		0.53		i.77 i.41	0.07	7.00
28-Sep-09	9:34	4.84	AA6	Anaerobic		21.50		0.27		0.02		0.30		.36	0.07	7.01
28-Sep-09	9:49	0.00	AER1	Aerobic		21.15		0.27		0.03		0.29		.89	0.06	7.01
28-Sep-09	9:58	0.31	AER1.25	Aerobic		15.95		3.05		0.52		3.57		.19	2.48	7.00
28-Sep-09	10:06	0.63	AER1.5	Aerobic		10.25		6.41	1	1.08		7.49	17	.21	3.08	6.95
28-Sep-09	10:51	3.13	AER3	Aerobic		4.05		13.10	1	1.33		14.43	8.	71	2.77	6.70
28-Sep-09	11:51	6.25	AER5	Aerobic		0.00		17.22	0	0.06		17.28	0.	58	1.45	6.67
28-Sep-09	12:00		IDC	-		0.00		17.18	0	0.02		17.20	0.	33	3.67	6.71
28-Sep-09	15:00		SE	-		0.00		17.02	0	0.02		17.04	0.	29	2.80	6.89
28-Sep-09	15:25		RAS	-		0.00		11.24	1	1.72		12.96	1.	70	0.13	6.84
	se values are <	1 mg/L NH₄-N														
		ample	<u> </u>	SE												
		365.1		EPA	350	ı Fi	PA 35	3.2	FPA	353.	2	FPA	353.2	FF	PA 35	32
		303.1			330.		A 33	3.2		x 333.	_	LIA	333.2		A 33	5.2
İ																
İ				0.1	200		NITT N	т.	NIC		.	NIC	N. N.T	١,	NO 1	. T
5		by CE	<u> </u>	O-I	-		NH ₄ -1			$0_{2,3}$ -N			D ₃ -N	1	NO ₂ -1	
		ng/L] 192		mg	;/L		mg/L	<u> </u>	m	ng/L		m	g/L	<u> </u>	mg/I	_
		109														
		97														
		92														
		68 64														
		64 60														
		60														
		56														
		47														
		45 38														
		36 40														
		4 0														
		51		0.3	32		<0.20	<u> </u>	1	17.6		1'	7.6		0.02	
				0.3	32		<0.20	<u> </u>	1	17.6		1	7.6		0.02	
				0.3	32		<0.20	<u> </u>	1	17.6		17	7.6		0.02	
				0.3	32	<u></u>	<0.20) <u> </u>	1	17.6		1′	7.6	T	0.02	
ı			A						T							
Actual		robic MLSS	Average A	AER DO	Min Al		NUR (NH4	-N Slope	e)	SAUR		NPR (N	NOx-N Slope		SNPF	
[days	s]	[mg/L]	[mg	AER DO	Min AE	ess]	NUR (NH4	-N Slope ∟/hr]	e)	SAUR /g MLVS	SS/hr]	NPR (N	NOx-N Slope		SNPF g/g MLV	SS/hr]
	s]			AER DO	Min Al	ess]	NUR (NH4	-N Slope ∟/hr]	e)	SAUR	SS/hr]	NPR (N	NOx-N Slope		SNPF	SS/hr]
[days 4.13	s] 3	[mg/L] 2300	[mg.	AER DO	Min AE	ess]	NUR (NH4	-N Slope ∟/hr]	e)	SAUR /g MLVS 1.65		NPR (N	NOx-N Slope g/L/min] 4.03	[mg	SNPF g/g MLV 2.28	SS/hr]
[days 4.13	S] 3 AER NO ₂ -	[mg/L] 2300 N PUR (PO ₄	[mg, 1.9 -P Uptake	AER DO /L] 7	Min AE [unitl 6.6	ess]	NUR (NH4	-N Slope L/hr] 92	e) [mg/	SAUR /g MLVS 1.65 Clar	ifier IDC	NPR (N [m	NOx-N Slope g/L/min] 4.03 Max ANA PO	[mg	SNPF g/g MLV 2.28 Avg. ANA	SS/hr]
[days 4.13 Highest A	AER NO ₂ -centration	[mg/L] 2300	[mg, 1.9 -P Uptake	AER DO	Min AE [unitl 6.6	ess]	NUR (NH4 [mg/ 2.9	-N Slope L/hr] 92	e)	SAUR /g MLVS 1.65 Clar		NPR (N [m	NOx-N Slope g/L/min] 4.03	[mg	SNPF g/g MLV 2.28	SS/hr]
[days 4.13 Highest A Conce	AER NO ₂ -centration	[mg/L] 2300 N PUR (PO ₄ Slo	[mg, 1.9 -P Uptake pe) L/hr]	AER DO (L] 7 SPUI	Min AE [unitl 6.6	ess] 67 Clarifier IE	NUR (NH4 [mg/ 2.9 DC NH4-N 1/L]	-N Slope L/hr] 32 Clarifier [n	[mg/	SAUR /g MLVS 1.65 Clar	ifier IDC Concentr [mg/L	NPR (N [m	MOx-N Slope g/L/min] 4.03 Max ANA PO Release [mg/L]	[mg	SNPF g/g MLV 2.28 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 4.13 Highest A Conce	AER NO ₂ -centration	[mg/L] 2300 N PUR (PO ₄ Slo	[mg. 1.9 -P Uptake pe)	AER DO /L] 7 SPU	Min AE [unitl 6.6	ess] 67 Clarifier IE	NUR (NH4 [mg/ 2.9 DC NH4-N 1/L]	-N Slope L/hr] 32 Clarifier [n	[mg/	SAUR /g MLVS 1.65 Clar	ifier IDC	NPR (N [m	NOx-N Slope g/L/min] 4.03 Max ANA PO Release	[mg	SNPF g/g MLV 2.28 Avg. ANA Concen	SS/hr] NO ₃ -N tration
[days 4.13 Highest A Conce	AER NO ₂ -centration	[mg/L] 2300 N PUR (PO ₄ Slo	[mg, 1.9 -P Uptake pe) L/hr]	AER DO (L] 7 SPUI	Min AE [unitl 6.6	ess] 67 Clarifier IE	NUR (NH4 [mg/ 2.9 DC NH4-N 1/L]	-N Slope L/hr] 32 Clarifier [n	[mg/	SAUR /g MLVS 1.65 Clar	ifier IDC Concentr [mg/L	NPR (N [m	MOx-N Slope g/L/min] 4.03 Max ANA PO Release [mg/L]	[mg	SNPF g/g MLV 2.28 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 4.13 Highest A Conce	AER NO ₂ -centration	[mg/L] 2300 N PUR (PO ₄ Slo	[mg, 1.9 -P Uptake pe) L/hr]	SPUI [mg/g MLV 3.98	Min AE [unitl] 6.6	ess] 67 Clarifier IE	OC NH4-N b/L]	-N Slope L/hr] 32 Clarifier [n	IDC NOx	SAUR /g MLVS 1.65 Clar	ifier IDC concentr [mg/L 0.3	NPR (N [m	MOx-N Slope g/L/min] 4.03 Max ANA PO Release [mg/L]	[mç	SNPF g/g MLV 2.28 Avg. ANA Concent	NO ₃ -N tration
[days 4.13 Highest A Conce	AER NO ₂ -centration mg/L]	[mg/L] 2300 N PUR (PO ₄ Slo	[mg 1.9 -P Uptake pe) L/hr]	SPUF [mg/g MLV 3.98	Min Al [unitl 6.6 SS/hr]	ess] 67 Clarifier IE [mg 0.00	MUR (NH4 [mg/] 2.9 DC NH4-N J/L] D000	-N Slope L/hr] 32 Clarifier [n 17	IDC NOxng/L]	SAUR /g MLVS 1.65 Clar c-N C	ifier IDC	NPR (N [m	MOx-N Slope [g/L/min] 4.03 Max ANA P(Release [mg/L] 22.68	O ₄ -P /	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32	NO ₃ -N tration L] 22
[days 4.13 Highest Conco	AER NO ₂ -centration mg/L] 1.33	mg/L 2300 PUR (PO ₄ Sloo mg/L 7.	[mg 1.9 1.9 -P Uptake pe) L/hr] 05 AA1 PC Release \$	SPUI SPUI [mg/g MLV 3.98	Min AE [unitl 6.6 SS/hr] A1 Sp	Clarifier IE [mg 0.00 pecific PC pase Rate	MUR (NH4- [mg/] 2.9 OC NH4-N [VL] OOO AAA Rele	-N Slope L/hr] 32 Clarifier [n 17 2 PO ₄ -P ase Slope	IDC NOxng/L]	SAUR /g MLVS 1.65 Clar (-N C	ifier IDC concentr [mg/L 0.3	NPR (N [m	MOx-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P ease Slope	O ₄ -P /	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32 3 Specif Release	NO ₃ -N tration L] ic PO ₄ Rate
[days 4.13 Highest . Conce	AER NO ₂ -lentration mg/L] 1.33 Ox-N A/L]	[mg/L] 2300	[mg 1.9 1.9 -P Uptake pe) L/hr] 05 AA1 PC	SPUF [mg/g MLV 3.98 04-P ASlope hr] [Min AE [unitl 6.6 SS/hr] AA1 Sp P Rele mg/g M	Clarifier IE [mg 0.00	MUR (NH4 [mg/] 2.9 DC NH4-N [/L] DO00 AA Rele	-N Slope L/hr] 32 Clarifier [n] 17	IDC NOxng/L]	SAUR //g MLVS 1.65 Clar c-N C	ifier IDC concentr [mg/L 0.3 ific PC e Rate	NPR (N [m	Mox-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P	O ₄ -P /	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32	NO ₃ -N tration L] ic PO ₄ Rate
[days 4.13 Highest Conco	AER NO ₂ -lentration mg/L] 1.33 Ox-N A/L]	[mg/L] 2300 N PUR (PO ₄ Slo [mg/ 7.	[mg, 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1,9 1,	SPUF [mg/g MLV 3.98 04-P ASlope hr] [Min AE [unitl 6.6 SS/hr] AA1 Sp P Rele mg/g M	ess] Clarifier IE [mg 0.00 escific PC ease Rate	MUR (NH4 [mg/] 2.9 DC NH4-N [/L] DO00 AA Rele	-N Slope L/hr] 32 Clarifier [n] 17 2 PO ₄ -P ase Slop	IDC NOxng/L]	SAUR /g MLVS 1.65 Clar C-N C	ifier IDC concentr [mg/L 0.3 ific PC e Rate	NPR (N [m	Mox-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P ease Slope mg/L/hr]	O ₄ -P /	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32 3 Specif Release	NO ₃ -N tration L] ic PO ₄ Rate
[days 4.13 Highest Conce	AER NO ₂ -1 centration mg/L] 1.33 Ox-N AA /L] 33	[mg/L] 2300 PUR (PO ₄ Slo	[mg 1.9 1.9 -P Uptake pe) L/hr] 05 AA1 PC Release \$\frac{1}{2} \text{[mg/L/ 7.82]}	SPUI SPUI [mg/g MLV 3.98 04-P A Slope hr] [Min AE [unitl 6.6 SS/hr] AA1 Sp P Rele mg/g M	Clarifier IE mg 0.00 escific PC ease Rate MLVSS/hi 0.06	OC NH4-N (yL) 000 AA Rele	-N Slope L/hr] 32 Clarifier [n 17 2 PO ₄ -P ase Slop ng/L/hr] 10.08	IDC NOx ng/L] 2040	SAUR /g MLVS 1.65 Clar C-N C A2 Spec P Releas ng/g MLV 11.6	ifier IDC concentr [mg/L 0.3 iffic PC e Rate VSS/hr	NPR (N [m	Mox-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P ease Slope mg/L/hr] 7.11	AA3	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32 3 Specif Release a/g MLV 8.25	NO ₃ -N tration L] 22 ic PO ₄ Rate SS/hr]
[days 4.13 Highest Conco	AER NO ₂ -1 centration mg/L] 1.33 Ox-N A//L] 3	[mg/L] 2300 N PUR (PO ₄ Slo [mg/ 7.	[mg 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9	SPUI SPUI [mg/g MLV 3.98 04-P A Slope hr] [Min AR [unitl] 6.6 SSS/hr] A1 Sp P Relea mg/g N SA3 Speak	ess] Clarifier IE [mg 0.00 escific PC ease Rate	MUR (NH4 [mg/] 2.9 DC NH4-N [/L] DO00 AA Rele	-N Slope L/hr] 32 Clarifier [n 17 2 PO ₄ -P ase Slop ng/L/hr] 10.08	IDC NOxng/L]	SAUR /g MLVS 1.65 Clar CA2 Spec P Releas ng/g MLV 11.6	ifier IDC concentr [mg/L 0.3 ific PC e Rate /SS/hi	NPR (N [m	Mox-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P ease Slope mg/L/hr]	AAC P I [mg	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32 3 Specif Release	NO ₃ -N tration L] ic PO ₄ Rate SS/hr]
Idays 4.13 Highest Conce	AER NO ₂ -1 centration mg/L] 1.33 Ox-N A//L] 3	[mg/L] 2300	[mg 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9 1.9	SPUF [mg/g MLV 3.98 04-P AD4-P FO4-P E Slope	Min AE [unitl] 6.6 SS/hr] A1 Sp P Relea My G N SAA5 Spee	Clarifier IE [mg 0.00 pecific PC pase Rate MLVSS/hi 0.06	DC NH4-N (yL) (yL) (AA6 PC	-N Slope L/hr] 22 Clarifier [n 17 2 PO ₄ -P ase Slop ng/L/hr] 10.08 0 ₄ -P A Slope F	IDC NOxing/L] 2040 AA Special	SAUR /g MLVS 1.65 Clar C A2 Spec P Releas ng/g MLV 11.6	ifier IDC concentr [mg/L 0.3 ific PC e Rate /SS/hr i9	NPR (N [m	Mox-N Slope [g/L/min] 4.03 Max ANA PO Release [mg/L] 22.68 A3 PO ₄ -P ease Slope mg/L/hr] 7.11 (AA1) Spe	AA3 P	SNPF g/g MLV 2.28 Avg. ANA Concent [mg/ 0.32 3 Specif Release g/g MLV 8.25	NO ₃ -N ration L] 22 ic PO ₄ Rate SS/hr]

# Chem Used [lbs]	FeCl3 Hrs U		3 Addition	Active Bios Influent FI		_	∕ Flow GD1		CY Flow	NRCY Flow
0	0.000		mg/L] 0.000	20.70			.33	_	2.17	0
ANA MLTSS	S ANA % VOL	ANA MLVS	ANX MLTSS	S ANX % VOL	ANX	MLVSS	AER M	LTSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	'L]	[%]	[mg/L]
1120	77	862	2300	77	1	771	230	0	77	1771

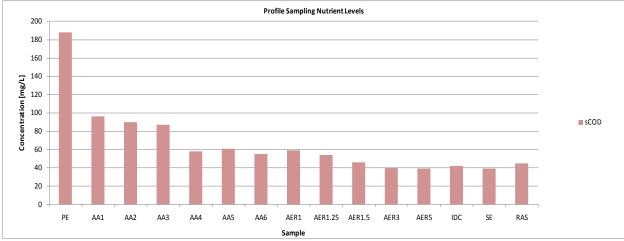


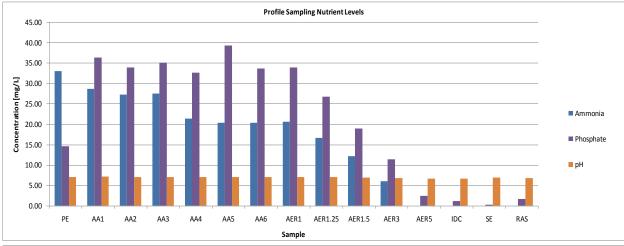


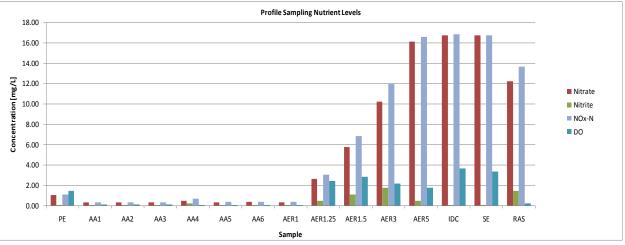


	2 Train 5, Ae	ration Tank 5,	Clarifier 5												
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Typ	e Ammo	onia by HACH T	'NT Nitrate b	by HACH TNT	Nitrite by HACI		N by Nitrate HAC Nitrite HACH		te by HAC IT844	H DO	рН
dd-mmm-yy	h:mm	hr				mg/L NH3-N	mg/	L NO3-N	mg/L NO2-	N	mg/L NOx-N	ma/L	PO4-P	[mg/L]	[unitless]
30-Sep-09	7:15	0	PE	-		33.10	_ · ·	1.03	0.03		1.06		4.68	1.44	7.04
30-Sep-09	7:30	0.84	AA1	Anaerobic		28.70		0.31	0.01		0.32		6.38	0.11	7.13
30-Sep-09	8:00	1.61	AA2	Anaerobic		27.30		0.32	0.01		0.33	3	3.94	0.12	7.12
30-Sep-09	8:30	2.42	AA3	Anaerobic		27.50		0.32	0.02		0.34	3	5.08	0.12	7.10
30-Sep-09	9:00	3.23	AA4	Anaerobic		21.40		0.45	0.22		0.67		2.63	0.08	7.06
30-Sep-09	9:15	4.03	AA5	Anaerobic		20.35		0.32	0.03		0.35		9.32	0.08	7.04
30-Sep-09	9:30	4.84	AA6	Anaerobic		20.45		0.34	0.02		0.36		3.77	0.07	7.05
30-Sep-09 30-Sep-09	9:45 9:53	0.00	AER1 AER1.25	Aerobic Aerobic		20.65 16.70		0.33 2.60	0.01		0.34 3.04		3.94 6.76	0.05 2.40	7.03 7.01
30-Sep-09	10:01	0.63	AER1.5	Aerobic		12.15		5.77	1.07		6.84		9.01	2.40	6.97
30-Sep-09	10:46	3.13	AER3	Aerobic		6.06		10.24	1.77		12.01		1.39	2.18	6.79
30-Sep-09	11:46	6.25	AER5	Aerobic		0.00		16.14	0.47		16.61		2.47	1.74	6.64
30-Sep-09	11:59	0.20	IDC	-		0.00		16.76	0.07		16.83		1.13	3.67	6.71
30-Sep-09	14:59		SE	_	-	0.00		16.74	0.02		16.76).30	3.33	6.93
				-											
30-Sep-09	15:15 se values are <	1 ma/l NIII NI	RAS	-		0.00		12.24	1.44		13.68		1.66	0.20	6.75
		•	- 14	<u> </u>											
QA		ample		SE											
	EPA	365.1		EPA	350.	1 E	PA 35	3.2	EPA 3	53.2	EPA	353.2	E	PA 35	3.2
	sCOD	by CE	.	O-1	PO ₄		NH ₄ -1	7	$NO_{2.3}$:-N	NC	D 3-N		NO ₂ -1	N
`		ng/L]			g/L	-	mg/L		mg/			g/L	1	mg/L	
		188		1112	5/10		IIIg/ L	_	IIIg/		1112	5/12		mg/ I	
		96													
		90													
		87													
		58													
		61													
		55													
		55 59													
		55 59 54													
		55 59 54 46													
		55 59 54 46 40													
		55 59 54 46 40 39													
		55 59 54 46 40 39					<0.20	, T	17.	6	12	7.6		0.02	
		55 59 54 46 40 39 42					<0.20)	17.	6	17	7.6		0.02	
	T	55 59 54 46 40 39					<0.20)	17.	6	17	7.6		0.02	
		55 59 54 46 40 39 42													
Actual		55 59 54 46 40 39 42	Average A	LER DO	Min A	ER pH /	<0.20			6 AUR		7.6 IOx-N Slop	e)	0.02 SNPF	
Actual [days	SRT Ae	55 59 54 46 40 39 42 39 45	Average A			ER pH /		1-N Slope	e) S/		NPR (N				₹
	SRT Ae	55 59 54 46 40 39 42 39 45		'L]	[unit		AUR (NH4	I-N Slope L/hr]	s) S/	AUR	NPR (N	IOx-N Slop		SNPF	R [SS/hr]
[days	SRT Aes]	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120	[mg/	'L]	[unit	less]	AUR (NH4 [mg/	I-N Slope L/hr]	s) S/	AUR 1LVSS/hr .83	NPR (N	IOx-N Slope g/L/min] 3.29	[m	SNPF ig/g MLV 2.04	R [SS/hr]
[days 3.63 Highest	SRT Aes] 3 AER NO ₂ -	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120	[mg/	′L] 4	[unit 6.	less]	AUR (NH4 [mg/	I-N Slope L/hr]	s) S/	AUR ILVSS/hr .83 Clarifier II	NPR (N [m	IOx-N Slop g/L/min]	[m	SNPF ig/g MLV	R [SS/hr]
[days 3.63 Highest	SRT Aes]	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120	[mg/	'L]	[unit 6.	less] 64	AUR (NH4 [mg/	1-N Slope L/hr] 95	s) S/	AUR ILVSS/hr .83 Clarifier II	NPR (N	IOx-N Slope g/L/min] 3.29	[m	SNPF ig/g MLV 2.04	R [SS/hr]
[days 3.63 Highest Cond	SRT Aess] 3 AER NO ₂ -centration	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO	[mg/ 1.8 -P Uptake ope)	'L] 4 SPU	[unit 6.	less] 64 Clarifier II	AUR (NH4 [mg/ 2.: DC NH4-N	I-N Slope L/hr] 95 Clarifier	(mg/g N IDC NOx-N	AUR ILVSS/hr .83 Clarifier II Conce	NPR (N [m	IOx-N Slop g/L/min] 3.29 Max ANA P Release	[m PO ₄ -P	SNPF g/g MLV 2.04 Avg. ANA Concent	R [SS/hr] NO ₃ -N tration
[days 3.63 Highest Cond	SRT Aess] AER NO ₂ -centration	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO	[mg/ 1.8 1-P Uptake ope)	'L] 4 SPU [mg/g ML\	[unit 6. R (SS/hr]	less] 64 Clarifier II	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L]	I-N Slope L/hr] 95 Clarifier	(mg/g N [mg/g N 1 IDC NOx-N	AUR ILVSS/hr .83 Clarifier II Conce	NPR (N [m DC PO ₄ -P ntration	IOx-N Slop g/L/min] 3.29 Max ANA P Release [mg/L]	PO ₄ -P	SNPF ig/g MLV 2.04 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 3.63 Highest Cond	SRT Aess] 3 AER NO ₂ -centration	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO	[mg/ 1.8 g-P Uptake ope) /L/hr]	SPU [mg/g ML\	[unit 6. R (SS/hr]	less] 64 Clarifier II	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L]	I-N Slope L/hr] 95 Clarifier [n	(mg/g N IDC NOx-N	AUR ILVSS/hr .83 Clarifier II Conce	NPR (N [m	JOx-N Slope g/L/min] 3.29 Max ANA P Release [mg/L] 24.64	PO ₄ -P	SNPF g/g MLV 2.04 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 3.63 Highest Cond	SRT Aess] AER NO ₂ -centration	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO, Slo	[mg/ 1.8 -P Uptake ppe) /L/hr] 98	SPU [mg/g ML\ 3.71	[unit 6.	Clarifier II [m: 0.0	AUR (NH4 [mg/ 2.1 DC NH ₄ -N g/L] 000 AA2 F	I-N Slope L/hr] 95 Clarifier [r] 16	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR 1LVSS/hr .83 Clarifier II Conce [m] 1	NPR (N [m	JOx-N Slope g/L/min] 3.29 Max ANA P Release [mg/L] 24.64	PO ₄ -P e	SNPF g/g MLV 2.04 Avg. ANA Conceni [mg/0.34	R SS/hr] NO ₃ -N tration /L] 42
[days 3.63 Highest Cond	SRT Aess] AER NO ₂ -centration mg/L] 1.77	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO, Slo	[mg/ 1.8 g-P Uptake ope) /L/hr]	SPU [mg/g ML\ 3.71	[unit 6.	Clarifier II	AUR (NH4 [mg/ 2.: DC NH ₄ -N g/L]	I-N Slope L/hr] 95 Clarifier [r] 16	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR 1LVSS/hr .83 Clarifier II Conce [m] 1	NPR (N [m	JOx-N Slope g/L/min] 3.29 Max ANA P Release [mg/L] 24.64	PO ₄ -P	SNPF g/g MLV 2.04 Avg. ANA Concent [mg/ 0.34	R SS/hr] NO ₃ -N tration /L] 42
[days 3.63 Highest Cond	SRT Aees] AER NO ₂ -centration mg/L] 1.77	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO, Slo	[mg/ 1.8 -P Uptake ppe) /L/hr] 98	SPU [mg/g ML\] 3.71 P AA ppe P	[unit 6.	Clarifier II [m 0.0 cific PO ₄ - se Rate	AUR (NH4 [mg/ 2.1 DC NH ₄ -N g/L] 000 AA2 F	I-N Slope L/hr] 95 Clarifier [r] 16 PO ₄ -P	(mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR **ILVSS/hr** .83 Clarifier II Conce [m] 1 fific PO ₄ -Fe Rate	NPR (N m DC PO ₄ -P ntration g/L]	JOx-N Slope g/L/min] 3.29 Max ANA P Release [mg/L] 24.64	PO ₄ -P e	SNPF g/g MLV 2.04 Avg. ANA Conceni [mg/0.34	SS/hr] NO ₃ -N tration L] 42 PO ₄ -P ate
[day: 3.63 Highest Cond	SRT Aees] AER NO ₂ -centration mg/L] 1.77 Dx-N AA	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO. Sic	[mg/ 1.8 -P Uptake ppe) /L/hr] 98 AA1 PO ₄ - Release Slo	SPU [mg/g ML\] 3.71 P AA ppe P	[unit 6.	Clarifier II [m	AUR (NH4 [mg/ 2.* DC NH ₄ -N g/L] 000 AA2 F Release	I-N Slope L/hr] 95 Clarifier [r] 16 PO ₄ -P 9 Slope L/hr]	[mg/g N 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	AUR **ILVSS/hr** .83 Clarifier II Conce [m] 1 iffic PO ₄ -Fe Rate _VSS/hr]	NPR (N [m DC PO ₄ -P ntration g/L] .1 AA3 R Release	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope	PO ₄ -P e	SNPF 1g/g MLV 2.04 Avg. ANA Concent [mg/0.34	SS/hr] NO ₃ -N tration L] 42 PO ₄ -P ate
[day: 3.63] Highest Conc. [Interpretation of the control of the co	SRT Ae s s AER NO ₂ - centration mg/L 1.77 Dx-N AA L 7	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO, Slo Slo [mg/L] 0.36	[mg/L/hr] 98 AA1 PO ₄ - Release Slo	SPU (mg/g ML\) 3.71 P AAA Ope P [[m	[unit 6. R //SS/hr] 1 Spec Release g/g ML 20.	Clarifier II [m] 0.0 clific PO ₄ - se Rate VSS/hr]	AUR (NH4 [mg/ 2.1 DC NH ₄ -N g/L] 0000 AA2 P Release [mg/ 8.7	I-N Slope L/hr] 95 Clarifier [r] 16 PO ₄ -P 9 Slope L/hr]	[mg/g M	AUR **ILVSS/hr** .83 Clarifier II Conce [m] 1 iffic PO ₄ -Fe Rate _VSS/hr]	NPR (N [m DC PO ₄ -P ntration g/L] .1 AA3 R Release	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope	PO ₄ -P e	SNPF 19/9 MLV 2.04 Avg. ANA Concent [mg/0.34 Specific Felease Raig MLVS:	SS/hr] NO ₃ -N tration L] 42 PO ₄ -P ate
[days 3.63 Highest Conc [r	SRT Ae s s AER NO ₂ - centration mg/L 1.77 Dx-N AA L 7	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO. Slo [mg/L] 5.	[mg/L/hr] 98 AA1 PO ₄ - Release Slo	SPU (mg/g ML\) 3.71 P AAA Ope P [[m	[unit 6. R //SS/hr] 1 Spec Release g/g ML 20.	Clarifier II [m	AUR (NH4 [mg/ 2.1 DC NH ₄ -N g/L] 0000 AA2 P Release [mg/ 8.7	I-N Slope L/hr] 95 Clarifier [r 16 PO ₄ -P Slope L/hr] 73	[mg/g M	AUR ILVSS/hr .83 Clarifier II Conce [m. 1 ific PO ₄ -F e Rate _VSS/hr] 96	NPR (N [m DC PO ₄ -P ntration g/L] .1 AA3 R Release	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope	PO ₄ -P e AA3 S Re	SNPF g/g MLV 2.04 Avg. ANA Concent [mg/ 0.32 Specific Felease Re (g MLVS) 9.36	SS/hr] NO ₃ -N tration /L] 42 PO ₄ -P ate S/hr]
[day: 3.63] Highest Conc. [in AA4 NC. [mg/0.66]	SRT Aees] AER NO ₂ - Dentration mg/L] 1.77 Dx-N AA L] 7 PO ₄ -P A	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO, Slo Slo [mg/L] 0.36	[mg/ 1.8 1.8	SPU (mg/g ML\) 3.71 P AAA Ope P [[m	[unit 6. R / SS/hr] 1 Spec Release g/g ML 20. AA5 Spec	Clarifier II [m] 0.0 clific PO ₄ - se Rate VSS/hr]	AUR (NH4 [mg/ 2.1 DC NH ₄ -N g/L] 0000 AA2 P Release [mg/ 8.7	I-N Slope L/hr] 95 Clarifier [r 16 PO ₄ -P Slope L/hr] 73	[mg/g N IDC NOx-N mg/L] .8300 AA2 Spec Releas [mg/g MI 11.	AUR ILVSS/hr .83 Clarifier II Conce [m. 1 ific PO ₄ -F e Rate VSS/hr] 96	NPR (N [m DC PO ₄ -P ntration g/L .1 AA3 F Release [mg/	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope [/L/hr] 83	PO ₄ -P e AA3 S Re	SNPF g/g MLV 2.04 Avg. ANA Concent [mg/ 0.32 Specific Felease Re (g MLVS) 9.36	SS/hr] NO ₃ -N tration /L] 42 PO ₄ -P ate S/hr]
[day: 3.65] Highest Conco	SRT Aees] AER NO ₂ -centration mg/L] 1.77 Ox-N AA L] 7 Oq-P A	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO. Slo [mg/L] 0.36 A4 Specific PC Release Rate	[mg/L/hr] AA1 PO ₄ - Release Slo [mg/L/hr] 15.16 Q ₄ -P AA5 I Release	SPU SPU SPU May a MLV 3.71 P AA AA PP AA PP AA PP AA PP AB PO4-P P B Slope	[unit 6. R //SS/hr] 1 Spec Release g/g ML 20. AA5 Spec Release Releas	Clarifier II [m 0.0 ciffic PO ₄ - se Rate VSS/hr] 77 ecific PO ₄ -P ase Rate	AUR (NH4 [mg/L] DC NH ₄ -N g/L] 000 AA2 P Release [mg/l 8.7	I-N Slope L/hr] 95 Clarifier [r] 16 PO ₄ -P 9 Slope L/hr] 73 O ₄ -P	[mg/g M] AA6 Specific I P Release R	AUR AUR AUR Clarifier II Conce [m] 1 fific PO ₄ -F e Rate VSS/hr] 96 PO ₄ - (A	NPR (N [m OC PO ₄ -P ontration of L] .1 AA3 F Release [mg/6. AA1) COD cake Slope	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope (L/hr] 83 (AA1) Spec Uptake	PO ₄ -P e AA3 S Ref [mg/	SNPF 19/9 MLV 2.04 Avg. ANA Concent [mg/ 0.34 Specific Felease Ref 19 MLVS: 9.36 (AA1) F	R SS/hr] A NO ₃ -N tration /L] 42 PO ₄ -P ate S/hr]
[day: 3.63] Highest Conc [I	SRT Aes SRT Ae	55 59 54 46 40 39 42 39 45 robic MLSS [mg/L] 2120 N PUR (PO. Slc	[mg/L/hr] AA1 PO4- Release Ski [mg/L/hr] 15.16 Q4-P AA5 Release hr] [mg	SPU (mg/g MLV 3.71 P AA OPP	[unit 6. R //SS/hr] 1 Spec Release g/g ML 20. AA5 Spec Release [mg/g N]	Clarifier II [m] 0.0 cific PO ₄ - se Rate VSS/hr] 77	AUR (NH4-Ng/L] OC NH4-Ng/L] OOO AA2 F Release [mg/] 8.7	I-N Slope L/hr] 95 Clarifier [r] 16 PO ₄ -P 9 Slope L/hr] 73 O ₄ -P 8 Slope	[mg/g M] IDC NOx-N mg/L] 8300 AA2 Spec Releas [mg/g MI] 11. AA6 Specific I	AUR AUR AUR Clarifier II Conce [m] 1 fific PO ₄ -F e Rate VSS/hr] 96 PO ₄ - (A	NPR (N [m DC PO ₄ -P ntration g/L] .1 AA3 F Release [mg/ 6.	Max ANA P Release [mg/L] 24.64 PO ₄ -P e Slope (L/hr] 83	PO ₄ -P e AA3 S Re [mg/	SNPF g/g MLV 2.04 Avg. ANA Concent [mg/ 0.32 Specific Felease Re (g MLVS: 9.36 (AA1) F	R SS/hr] NO ₃ -N tration L] 42 PO ₄ -P ate S/hr] PO ₄ -P/ DD

# Chem Used	FeCl3 Hrs U	sed FeCl3	3 Addition	Active Bios Influent Fl		ARCY	ſ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]	[r	ng/L]	[MGD]		[M	GD]	[1	MGD]	[MGD]
0	0.000	C	0.000	19.81		17.	.38	1	1.67	0
ANA MLTSS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	_TSS	AER % VC	L AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	L]	[%]	[mg/L]
960	76	730	2120	76	1	611	212	0	76	1611



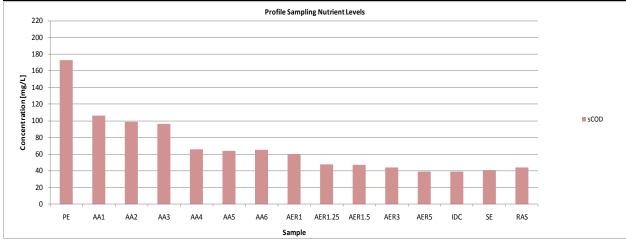


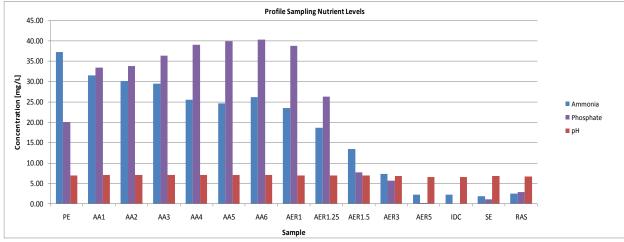


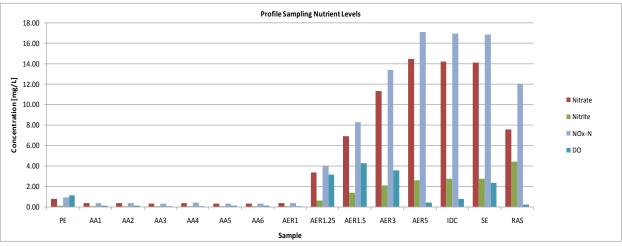
Week 12 – 10/6/09 & 10/8/09

vv eek			AL ''' 4											
'KOFILE 1	i rain 4, A	eration Tank 4,	Clarifier 4											
Date	Sample Tim	<u> </u>	Sample Number	Reactor Type		ia by HACH TNT	Nitrate by HACH TNT	Nitrite by HACH TN	T + Ni	y Nitrate HAC trite HACH	Phosphate b			рН
dd-mmm-yy	h:mm	hr			m	ıg/L NH3-N	mg/L NO3-N	mg/L NO2-N	mg	/L NOx-N	Ť	P04-P	[mg/L]	[unitless]
6-Oct-09	7:03	0	PE	-		37.20	0.79	0.14		0.93		0.07	1.11	6.95
6-Oct-09	7:20	0.89	AA1	Anaerobic		31.50	0.36	0.02		0.37		3.45	0.11	7.04
6-Oct-09	7:50	1.79	AA2	Anaerobic		30.10	0.37	0.01		0.38		3.77	0.11	7.03
6-Oct-09 6-Oct-09	8:20 8:50	2.68 3.57	AA3 AA4	Anaerobic Anaerobic		29.50 25.50	0.31 0.36	0.01		0.33		6.38 3.99	0.08	7.03
6-Oct-09	9:05	4.46	AA4 AA5	Anaerobic		24.60	0.30	0.00		0.44		9.97	0.00	7.02
6-Oct-09	9:20	5.36	AA6	Anaerobic		26.15	0.31	0.02		0.33		0.30	0.03	7.02
6-Oct-09	9:35	0.00	AER1	Aerobic		23.50	0.35	0.02		0.38		3.83	0.07	6.98
6-Oct-09	9:43	0.35	AER1.25	Aerobic		18.65	3.37	0.64		4.01		6.35	3.16	7.00
6-Oct-09	9:51	0.69	AER1.5	Aerobic		13.45	6.92	1.38		8.30	7.	.75	4.29	6.98
6-Oct-09	10:36	3.46	AER3	Aerobic		7.35	11.30	2.08		13.38	5.	.68	3.54	6.78
6-Oct-09	11:36	6.92	AER5	Aerobic		2.21	14.48	2.60		17.08	0.	.14	0.42	6.62
6-Oct-09	11:45		IDC	-		2.27	14.20	2.74		16.94	0.	.13	0.75	6.63
6-Oct-09	14:45		SE	-		1.81	14.10	2.74		16.84	1.	.06	2.32	6.82
6-Oct-09	15:03		RAS	-		2.53	7.58	4.42		12.00	2.	.82	0.23	6.71
NOTE: Thes	se values are	<1 mg/L NH ₄ -N												
		Sample	: 5	SE				1						
		A 365.1		EPA 3	350.1	EP/	A 353.2	EPA 353	3.2	EPA	353.2	E	PA 35	3.2
		by CE	L	O-P	O_4	N	H ₄ -N	NO _{2,3} -1	4	NO	3- N		NO ₂ -1	Ν
s					/T		/T	/T			/T		/T	
s	[n	ng/L]		mg/	/L	n	ng/L	mg/L		mg	g/L		mg/I	_
s	[n	ng/L] 173		mg/	/L	n	ng/L	mg/L		mg	z/L		mg/L	,
S	[n	ng/L]		mg,	/L	n	ng/L	mg/L		mg	g/L		mg/I	-
S	[n	ng/L] 173 106 99 96		mg/	/L	n	ng/L	mg/L		mg	;/L		mg/I	
S	[n	ng/L] 173 106 99 96 66		mg/	/L	n	ng/L	mg/L		mg	;/L		mg/I	_
S	[n	ng/L] 173 106 99 96 66 64		mg/	L	n	ng/L	mg/L		mg	g/L		mg/I	
S	[n	ng/L] 173 106 99 96 66 64		mg	L_	n	ng/L	mg/L		mg	<u>z/L</u>		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48		mg	<u>/L</u>	n	ng/L	mg/L		mg	z/L		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48		mg	L_	n	ng/L	mg/L		mg	;/L		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44		mg/	/L	n	ng/L	mg/L		mg	;/L		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39		mg	/L	n	ng/L	mg/L		mg	;/L		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39		mg.										
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39					ng/L	mg/L			6		mg/I	
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39												
S	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39												
	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41		0.30	04		1.68	17.4		14	6		2.8	
	[n	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44	Average A	0.30 NER DO	D4 Min AE	R pH AU	1.68 R (NH4-N Slope	17.4 e) SAU		14 NPR (N	6 Ox-N Slope	_	2.8 SNPF	₹
Actual S	[r	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44	Average A	O.30	O-4 Min AE [unitle	ER pH AU	I.68 R (NH4-N Slope [mg/L/hr]	17.4 s) SAU [mg/g ML\	/SS/hr]	NPR (N	6 Ox-N Slope g/L/min]	_	2.8 SNPF ng/g MLV	R SS/hr]
Actual S	[r	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44	Average A	O.30	D4 Min AE	ER pH AU	1.68 R (NH4-N Slope	17.4 e) SAU	/SS/hr]	NPR (N	6 Ox-N Slope	_	2.8 SNPF	R SS/hr]
Actual S [days 5.01	SRT A	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760	Average A	O.30	O-4 Min AE [unitle	ER pH AU	I.68 R (NH4-N Slope [mg/L/hr]	17.4 s) SAU [mg/g ML\ 1.91	/SS/hr]	NPR (N	Ox-N Slope g/L/min] 3.21	[m	2.8 SNPP ng/g MLV 2.28	R SS/hr]
Actual S [days 5.01	SRT A	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760	Average A [mg 2.3	O.30	Min AE [unitle	ER pH AU	R (NH4-N Slope [mg/L/hr] 2.69	17.4 s) SAU [mg/g ML\ 1.91	'SS/hr] arifier IDC	NPR (N	Ox-N Slope g/L/min] 3.21	[m	SNPF ng/g MLV 2.28	R SS/hr]
Actual S [days 5.01 Highest Conc	SRT A S] AER NO ₂ entration	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO	Average A [mg 2.3 4-P Uptake	O.30 NER DO /L] 0	Min AE [unitle 6.6	ER pH AU ess] 62 Clarifier IDC	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier	17.4 s) SAU [mg/g ML\] 1.91 IDC NOx-N	/SS/hr] arifier IDC	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release	[m	SNPF ng/g MLV 2.28 Avg. ANA Concen	R SS/hr] NO ₃ -N
Actual S [days 5.01 Highest Conc	SRT A A AER NO ₂ entration mg/L	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO SI [mg/L]	Average A [mg. 2.3 4-P Uptake ppe)	O.30 NER DO /L] 0 SPUF	Min AE [unitle 6.6	ER pH AU ess] 22 Clarifier IDC [mg/L	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier	17.4 s) SAU [mg/g ML\] 1.91 CI IDC NOx-N mg/L]	arifier IDC Concenti	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA PR Release [mg/L]	[m	SNPF ng/g MLV 2.28 Avg. ANA Concen	SS/hr] NO ₃ -N
Actual S [days 5.01 Highest Conc	SRT A S] AER NO ₂ entration	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO SI [mg/L]	Average A [mg 2.3 4-P Uptake	O.30 NER DO /L] 0	Min AE [unitle 6.6	ER pH AU ess] 62 Clarifier IDC	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier	17.4 s) SAU [mg/g ML\] 1.91 IDC NOx-N	/SS/hr] arifier IDC	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release	[m	SNPF ng/g MLV 2.28 Avg. ANA Concen	SS/hr] NO ₃ -N
Actual S [days 5.01 Highest Conc	SRT A A AER NO ₂ entration mg/L	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO SI [mg/L]	Average A [mg. 2.3 4-P Uptake ppe)	O.30 NER DO /L] 0 SPUF	Min AE [unitle 6.6	ER pH AU ess] 22 Clarifier IDC [mg/L	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier	17.4 s) SAU [mg/g ML\] 1.91 CI IDC NOx-N mg/L]	arifier IDC Concenti	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA PR Release [mg/L]	[m	SNPF ng/g MLV 2.28 Avg. ANA Concen	SS/hr] NO ₃ -N
Actual S [days 5.01 Highest Conc	SRT A A AER NO ₂ entration mg/L	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO SI [mg/L]	Average A [mg. 2.3 4-P Uptake ppe)	O.30 AER DO /LI 0 SPUF [mg/g MLV 5.13	Min AE [unitle 6.6	ER pH AU ess] 22 Clarifier IDC [mg/L	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier	17.4 s) SAU [mg/g ML\] 1.91 CI IDC NOx-N mg/L]	arifier IDC Concenti [mg/l	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA PR Release [mg/L]	O ₄ -P	SNPF ng/g MLV 2.28 Avg. ANA Concen	NO ₃ -N viration
Actual S [days 5.01 Highest Conce	SRT A S AER NO ₂ entration mg/L] 2.60	ng/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 N PUR (PO SI [mg/L]	Average A [mg, 2.3 4-P Uptake ope) ////hr]	O.30 AER DO /L 0 SPUF [mg/g MLV 5.13	Min AE [unitle 6.6 SS/hr]	ER pH AU ess] i2 Clarifier IDC [mg/L 2.2700	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifler] [i	17.4 e) SAU [mg/g MLV 1.91 IDC NOx-N mg/L] 6.9400 AA2 Spec	arifier IDC Concentr [mg/l 0.1	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA Pe Release [mg/L] 20.23	O ₄ -P	SNPF ng/g MLV 2.28 Avg. ANA Concen [mg. 0.33	NO ₃ -N ration
Actual S [days 5.01 Highest Conc	SRT A S AER NO ₂ entration mg/L] 2.60 Dx-N A	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 PUR (PO SI [mg/A] 7 A6 NOX-N	Average A [mg, 2.3 4-P Uptake ppe) //L/hr] 23 AA1 PO ₄ Release SI	O.30 LER DO /L] 0 SPUF [mg/g MLV 5.13 -P AAF	Min AE [unitle 6.6 SS/hr]	Clarifier IDC [mg/L 2.270] cific PO ₄ - se Rate	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier 0 16 16 AA2 PO ₄ -P Release Slop	17.4 (mg/g MLV 1.91 IDC NOx-N mg/L] 3.9400 AA2 Spec	arifier IDC Concents [mg/l 0.1	NPR (N [mg PO ₄ -P ration]	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg 0.33 3 Specifi Release	NO ₃ -Nr Intration ILJ 34 Cc PO ₄ *
Actual S [days 5.01 Highest Conco	SRT A S AER NO ₂ entration mg/L] 2.60 Dx-N A/L]	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 PUR (PO SI [mg/L] 7 A6 NOx-N [mg/L]	Average A [mg/ 2.3 4-P Uptake ppe) /L/hr] 23 AA1 PO ₄ Release SI [mg/L/hr	O.30 LER DO /L] 0 SPUF [mg/g MLV 5.13 -P AAF	Min AE [unitle 6.6 SSS/hr] A1 Spec	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr]	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier] [t] 0 16 AA2 PO ₄ -P Release Slop [mg/L/hr]	1.74 (mg/g MLV	arifier IDC Concentr [mg/l 0.1 cific PO ₄ se Rate	NPR (N [mg PO ₄ -P ration]	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope ng/L/hr]	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg 0.33 3 Specifi Release	Rate SS/hrj
Actual S [days 5.01 Highest Conc	SRT A S AER NO ₂ entration mg/L] 2.60 Dx-N A/L]	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 PUR (PO SI [mg/A] 7 A6 NOX-N	Average A [mg, 2.3 4-P Uptake ppe) //L/hr] 23 AA1 PO ₄ Release SI	O.30 LER DO /L] 0 SPUF [mg/g MLV 5.13 -P AA lope F	Min AE [unitle 6.6 SS/hr]	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr]	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier 0 16 16 AA2 PO ₄ -P Release Slop	17.4 (mg/g MLV 1.91 IDC NOx-N mg/L] 3.9400 AA2 Spec	arifier IDC Concentr [mg/l 0.1 cific PO ₄ se Rate	NPR (N [mg PO ₄ -P ration]	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg 0.33 3 Specifi Release	Rate SS/hrj
Actual S [days 5.01 Highest Conc. [n 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	SRT A S] AER NO ₂ entration ng/L] 2.60 Dx-N A/L] 4	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 PUR (PO Si [mg/L] 0.33	Average A [mg/L/hr] 23 AA1 PO ₄ Release SI [mg/L/hr] 3.98	O.30 AER DO (L] 0 SPUF [mg/g MLV 5.13 -P AA- ope F] [n	Min AE [unitle 6.6 SS/hr] A1 Spece Releasing/g ML 4.6	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr] 62	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier 0 16 16 16 16 16 16 16	17.4 SAU [mg/g ML\ 1.91 LDC NOx-N mg/L] 5.9400 AA2 Spec P Relea: [mg/g ML 10.	arifier IDC Concentr [mg/l 0.1 cific PO ₄ se Rate -VSS/hr]	NPR (N [mg	Ox-N Slope g/L/min] 3.21 Max ANA Po Release [mg/L] 20.23 3 PO ₄ -P ase Slope ng/L/hr] 6.92	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg 0.33 3 Specifi Release	Rate SS/hr]
Actual S [days 5.01 Conc. [n 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	SRT A S AER NO2 entration mg/L] 2.60 Dx-N A/ L] 4	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 -N PUR (PO SI [mg/L] 0.33	Average A [mg/ 2.3 AA1 PO ₄ Release SI [mg/L/hr 3.98	O.30 AER DO /L] 0 SPUF [mg/g MLV 5.13 -P AA ope F] [n	Min AE [unitle 6.6 SSS/hr] A1 Spece Releasing/g ML 4.6 AA5 Spece	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr] 62	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier] [i [i] [i] [i] [i] [i] [i] [i	174 SAU [mg/g MLV 1.91 LDC NOx-N mg/L] 3.9400 AA2 Spece P Relea: [mg/g ML 10. A6 Specific PO ₄ -	arifier IDC Concentr [mg/l 0.1 siffic PO ₄ se Rate VSS/hr] 45	NPR (N [mg PO4-P ration]	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope ng/L/hr] 6.92 (AA1) Spe	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg- 0.33 Specifi Release ng/g MLV 8.02	SS/hr] NO ₃ -N rration L] 34 c PO ₄ Rate SS/hr]
Actual S [days 5.01 Highest Conc. [r] 2 AA4 NC [mg/I 0.44 AA4 F Release	SRT A A AER NO ₂ entration mg/L] 2.60 Dx-N A/ L] 4	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 PUR (PO Sing) [mg/L] 0.33 AA4 Specific F P Release Re	Average A [mg, 2.3 4-P Uptake ope) /L/hr] .23 AA1 PO ₄ Release SI [mg/L/hr] 3.98 O ₄ - AA5 in Release	O.30 AER DO (L] 0 SPUF [mg/g MLV 5.13 -P A/- ope F] [n] PO ₄ -P # e Slope	Min AE [unitle 6.6 SS/hr] A1 Spec P Release ng/g ML 4.6 AA5 Spec P Release	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr] 62 cific PO ₄ - se Rate	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier [i [i] [174 B) SAU [mg/g ML\] 1.91 LDC NOx-N mg/L] 3.9400 AA2 Spec P Releas [mg/g ML] 10. A6 Specific PO ₄ - P Release Rate	arifier IDC Concentron (mg/l 0.1 ciffic PO ₄ se Rate VSS/hr] 45	NPR (N [mg PO4-P ration -] - AA Rele [n	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope ng/L/hr] 6.92 (AA1) Spe COD Uptake	O ₄ -P	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg. 0.33 3 Specifi Release ng/g MLV 8.02	SS/hr] NO ₃ -N rration IL] 34 c c PO ₄ - Rate SS/hr]
Actual S [days 5.01 Highest Conc. [n 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2 2	SRT A S] AER NO2 entration mg/L] 2.60 Dx-N A/ L] 4 PO4-P e Slope L/hr]	mg/L] 173 106 99 96 66 64 65 60 48 47 44 39 39 41 44 erobic MLSS [mg/L] 1760 -N PUR (PO SI [mg/L] 0.33	Average A [mg, 2.3 4-P Uptake ope) /L/hr] .23 AA1 PO ₄ Release SI [mg/L/hr 3.98 O _{4*} AA5 I Releas /hr] [mg/	O.30 AER DO (L] 0 SPUF [mg/g MLV 5.13 -P AA ope F r] [n] PO ₄ -P A	Min AE [unitle 6.6 SSS/hr] A1 Spece Releasing/g ML 4.6 AA5 Spece	Clarifier IDC [mg/L 2.270 cific PO ₄ - se Rate LVSS/hr] 62	R (NH4-N Slope [mg/L/hr] 2.69 NH ₄ -N Clarifier [1	174 SAU [mg/g MLV 1.91 LDC NOx-N mg/L] 3.9400 AA2 Spece P Relea: [mg/g ML 10. A6 Specific PO ₄ -	arifier IDC Concentr [mg/l 0.1 siffic PO ₄ se Rate VSS/hr] 45 (AA1) CC SI	NPR (N [mg PO4-P ration]	Ox-N Slope g/L/min] 3.21 Max ANA Pr Release [mg/L] 20.23 3 PO ₄ -P ase Slope ng/L/hr] 6.92 (AA1) Spe	AA P [m	2.8 SNPF ng/g MLV 2.28 Avg. ANA Concen [mg- 0.33 Specifi Release ng/g MLV 8.02	SS/hr] NO ₃ -N tration IL 34 c PO ₄ - Rate SS/hr]

# Chem Used	FeCl3 Hrs U	sed FeCl3	3 Addition	Active Bios		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]	[r	ng/L]	[MGD]		[M	GD]	[]	MGD]	[MGD]
0	0.000		0.000	18.71		17.	67	1	1.223	0
ANA MLTS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	LTSS	AER % VC	L AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	'L]	[%]	[mg/L]
980	88	862	1760	80	1	408	176	0	80	1408

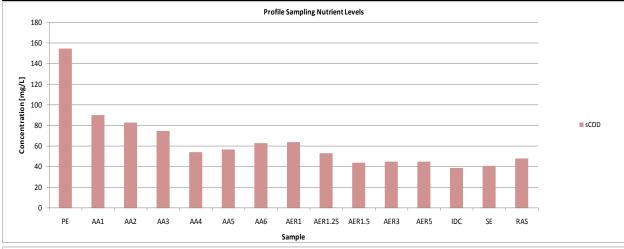


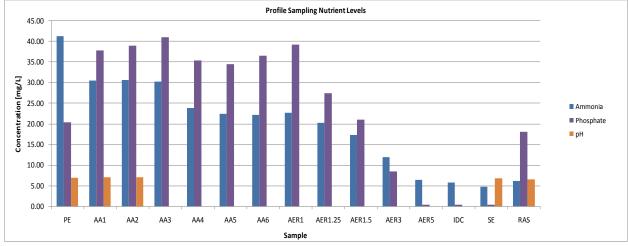


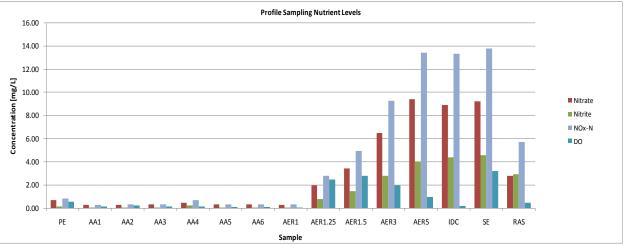


POFILE 2	Train 6 Ao	ration Tank 5,	Clarifier 5												
VOI ILL 2	main o, Ac	HRT along BNR	Siai iliei J												
Date	Sample Time	Process (w/out Recycles)	Sample Number	Reactor Type	Ammor	nia by HACH TNT	Nitrate by HACH	TNT N	Nitrite by HACH		by Nitrate HA Nitrite HACH		e by HAC T844	CH DO	pН
ld-mmm-yy	h:mm	hr			n	ng/L NH3-N	mg/L NO3-N		mg/L NO2-N	m	ng/L NOx-N	mg/L	P04-P	[mg/L]	[unitless]
8-Oct-09	7:06	0	PE	-		41.30	0.69		0.15		0.84	20).39	0.53	6.95
8-Oct-09	7:22	0.89	AA1	Anaerobic		30.50	0.28		0.01		0.30	37	.85	0.14	7.04
8-Oct-09	7:52	1.79	AA2	Anaerobic		30.70	0.29		0.01		0.30	38	3.99	0.23	7.10
8-Oct-09	8:22	2.68	AA3	Anaerobic		30.30	0.31		0.01		0.32	40).95	0.12	-
8-Oct-09	8:52	3.57	AA4	Anaerobic		23.80	0.44		0.25		0.69		5.41	0.15	-
8-Oct-09	9:07	4.46	AA5	Anaerobic		22.40	0.30		0.01		0.31		1.43	0.10	
8-Oct-09	9:22	5.36	AA6	Anaerobic		22.20	0.30		0.02		0.32		5.55	0.11	-
8-Oct-09	9:37	0.00	AER1	Aerobic		22.65	0.28		0.02		0.30).16	0.06	•
8-Oct-09	9:45	0.35	AER1.25	Aerobic		20.20	1.98		0.80		2.78		7.41	2.46	-
8-Oct-09 8-Oct-09	9:53	0.69 3.46	AER1.5 AER3	Aerobic		17.30	3.44 6.48		1.48 2.78		4.92 9.26		1.97	2.80	
8-Oct-09	10:38 11:38	6.92	AER5	Aerobic Aerobic		11.90 6.44	9.40		4.02		13.42		.48	1.97 0.97	
		0.92		- Aelobic											
8-Oct-09	11:51		IDC			5.84	8.92		4.40		13.32		.44	0.18	
8-Oct-09	14:58		SE	-		4.72	9.22		4.58		13.80		.43	3.18	6.75
8-Oct-09	15:22	_	RAS	-		6.14	2.78		2.92		5.70	18	3.03	0.46	6.52
QA/		ample		SE											
	EPA	365.1		EPA 3	350.1	EP/	A 353.2	E	EPA 35	53.2	EPA	353.2	E	PA 35	3.2
<	:COD	by CE	.	О-Р	\mathbf{O}_4	N	H4-N		NO _{2,3} -	. N	NC)3-N		NO ₂ -1	7
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		33 75 54 57 63 64 53 44 45				-	5.06		13.7		9.	05		4.65	
		33 75 54 57 63 64 53 44 45 45					5.06		13.7		9.	05		4.65	
		33 75 54 53 64 53 44 45 45 45 45		50.00	Min Af			I							
Actual		33 75 54 57 63 64 53 44 45 45	Average A	ER DO	Min AE		5.06 R (NH4-N SI	Dpe)	SA			O5 IOx-N Slope	€)	4.65 SNPF	₹
[days	SSRT Ae	83 75 54 57 63 64 53 44 45 45 45 45 45 48	[mg/	L]	[unitl	ER pH AU	R (NH4-N SI [mg/L/hr]	ppe)	SA [mg/g Ml	LVSS/hr]	NPR (N	IOx-N Slope g/L/min]		SNPF ng/g MLV	
	SSRT Ae	83 75 54 57 63 64 53 44 45 45 45 45 45 45 48		L]		ER pH AU	R (NH4-N SI	ppe)	SA	LVSS/hr]	NPR (N	IOx-N Slope		SNPF	
[days	SRT Ae	33 75 54 57 63 64 53 44 45 45 45 45 41 48 mobic MLSS [mg/L]	[mg/	L]	[unitl	ER pH AU	R (NH4-N SI [mg/L/hr]	ppe)	SA [mg/g Ml	LVSS/hr] 60	NPR (N	IOx-N Slope g/L/min] 2.26	[m	SNPF ng/g MLV 1.67	SS/hr]
[days 3.96 Highest	SSRT Ae. SSRT Ae. SSRT AAE.	33 75 54 57 63 64 53 44 45 45 39 41 48 mobic MLSS [mg/L] 1780	[mg/ 1.6 -P Uptake	[L] 5	[unitl 0.0	ER pH AU ess]	R (NH4-N SI [mg/L/hr] 2.17		SA [mg/g MI	LVSS/hr] 60 Clarifier ID	NPR (N [m	JOx-N Slope g/L/min] 2.26 Max ANA Po	[m O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA	SS/hr]
3.96 Highest	SRT Ae	33 75 54 57 63 64 53 44 45 45 39 41 48 mobic MLSS [mg/L] 1780	[mg/	L]	[unitl 0.0	ER pH AU	R (NH4-N SI [mg/L/hr] 2.17		SA [mg/g Ml	LVSS/hr] 60	NPR (N [m	IOx-N Slope g/L/min] 2.26	[m O ₄ -P	SNPF ng/g MLV 1.67	SS/hr]
[days 3.96 Highest Conc	SSRT Ae. SSRT Ae. SSRT AAE.	33 75 54 57 63 64 53 44 45 45 39 41 48 mobic MLSS [mg/L] 1780	[mg/ 1.6 -P Uptake pe)	[L] 5	[unitl 0.0	ER pH AU ess]	R (NH4-N SI [mg/L/hr] 2.17 NH ₄ -N Clari		SA [mg/g MI 1.	LVSS/hr] 60 Clarifier ID	NPR (N [m	JOx-N Slope g/L/min] 2.26 Max ANA Po	[m O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA	SS/hr] NO ₃ -N tration
[days 3.96 Highest Conc [r	SSRT Aes	83 75 54 57 63 64 53 44 45 45 45 45 45 45 45 45 45 47 48 Fobic MLSS [mg/L] 1780	[mg/ 1.6 -P Uptake pe)	L] 5 SPUF	[unitl 0.0 R SS/hr]	ER pH AU ess] 00 Clarifier IDC	R (NH4-N SI [mg/L/hr] 2.17 NH ₄ -N Clari	fier ID0	SA [mg/g MI 1.0 C NOx-N	LVSS/hr] 60 Clarifier ID Concen	NPR (N [m	IOx-N Slope g/L/min] 2.26 Max ANA Po Release	[m O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent	SS/hr] NO ₃ -N tration
[days 3.96 Highest Conc	SSRT Ae.	83 75 54 57 63 64 53 44 45 45 45 45 45 45 45 45 45 47 48 Fobic MLSS [mg/L] 1780	[mg/ 1.6 -P Uptake pe) L/hr]	SPUF	[unitl 0.0 R SS/hr]	ER pH AU ess] 00 Clarifier IDC [mg/L 5.840	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari	fier ID0 [mg/ 13.32	SA [mg/g MI] 1.	LVSS/hr] 60 Clarifier ID Concen [mg 0.	NPR (N [m	JOx-N Slope g/L/min] 2.26 Max ANA Po Release [mg/L] 20.56	[m	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/ 0.32	NO ₃ -N tration L]
[days 3.96 Highest Conc	SRT Ae:	33 75 54 57 63 64 53 44 45 45 45 45 41 48 mobic MLSS [mg/L] 1780 N PUR (PO ₄ Slo	[mg/ 1.6 -P Uptake pe) L/hr] 28	SPUF Smg/g MLV 5.38	[unitl 0.0 SS/hr]	ER pH AU ess] 00 Clarifier IDC [mg/L 5.840 cific PO ₄ -	R (NH4-N SI [mg/L/hr] 2.17 NH ₄ -N Clari] 0	fier ID0 [mg/ 13.32	SA [mg/g MI 1 1 1 1 1 1 1 1	Clarifier ID Concen [mg 0.ecific PO	NPR (N [rr	JOx-N Slope g/L/min] 2.26 Max ANA P Release [mg/L] 20.56	O ₄ -P	SNPF ng/g MLVV 1.67 Avg. ANA Concent [mg/0.32	NO ₃ -N tration
[days 3.96 Highest Conc [r	SRT Ae:	83 75 54 57 63 64 53 44 45 45 45 45 45 45 45 45 45 47 48 mobic MLSS [mg/L] 1780 N PUR (PO ₄ Slo	[mg/ 1.6 -P Uptake pe) L/hr]	SPUF Smg/g MLV 5.38	[unitl 0.0 SS/hr]	ER pH AU ess] 00 Clarifier IDC [mg/L 5.840	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari	fier ID0 [mg/ 13.32	SA [mg/g MI 1 1 1 1 1 1 1 1	LVSS/hr] 60 Clarifier ID Concen [mg 0.	NPR (N [rr	JOx-N Slope g/L/min] 2.26 Max ANA Po Release [mg/L] 20.56	O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/ 0.32	NO ₃ -N tration L]
[days 3.96 Highest Conc	SSRT Ae. SSRT Ae. AER NO ₂ -I entration mg/L] 4.02	33 75 54 57 63 64 53 44 45 45 45 45 41 48 mobic MLSS [mg/L] 1780 N PUR (PO ₄ Slo	[mg/ 1.6 -P Uptake pe) L/hr] 28	SPUF mg/g MLV 5.38 -P AA	[unitl 0.0 SS/hr]	ER pH AU ess] 00 Clarifier IDC [mg/L 5.840 cific PO ₄ -	R (NH4-N SI [mg/L/hr] 2.17 NH ₄ -N Clari] 0	fier ID0 [mg/ 13.32 -P ope	SA [mg/g MI 1.0 C NOx-N /L] 200 AA2 Sp P Rele	Clarifier ID Concen [mg 0.ecific PO	NPR (N [mr C PO ₄ -P otration VL] 4 AA Rele Rele	JOx-N Slope g/L/min] 2.26 Max ANA P Release [mg/L] 20.56	O ₄ -P	SNPF ng/g MLVV 1.67 Avg. ANA Concent [mg/0.32	NO ₃ -N tration L] 22 c PO ₄ - Rate
[days 3.96 dighest Conc [r	SSRT Ae: SSRT Ae: AER NO ₂ -I entration mg/L] 4.02 Dx-N AA	83 75 54 57 63 64 53 44 45 45 45 45 45 41 48 mobic MLSS [mg/L] 1780 N PUR (PO ₄ Slo	[mg/ 1.6 -P Uptake pe) L/hr] 28 AA1 PO ₄ Release SI	SPUF mg/g MLV 5.38 -P AA	[unitl 0.0 SS/hr] A1 Spe Relea	ER pH AU ess] 00 Clarifier IDC [mg/L 5.8400 cific PO ₄ - use Rate	R (NH4-N SI [mg/L/hr] 2.17 NH ₄ -N Clari] 0 AA2 PO ₄ Release SI	fier ID0 [mg/ 13.32 -P ope	SA [mg/g MI 1.0 C NOx-N //L] 2000 AA2 Sp P Rele [mg/g M	Clarifier ID Concen [mg 0.ecific PO ease Rate	NPR (N [mr C PO ₄ -P otration VL] 4 AA Rele Rele	Max ANA Port Release [mg/L] 20.56	O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/g 0.32 3 Specifi Release	NO ₃ -N tration L] 22 c PO ₄ - Rate
[days 3.96 dighest Conc [r	SRT Ae SRT NO2-1 entration mg/L] 4.02 Dx-N AA	33 75 54 57 63 64 53 44 45 45 39 41 48 **robic MLSS** [mg/L] 1780 **PUR (PO ₄ Slo [mg/L] 7. 6 NOx-N **mg/L] 0.32	[mg/L/hi 1.6 -P Uptake pe) L/hr] 28 AA1 PO ₄ Release SI [mg/L/hi 11.00	SPUF SPUF mg/g MLV 5.38 -P AAA ope F] [n	[unitl 0.0 R SS/hr] A1 Spe P Releang/g M	ER pH AU ess] 00 Clarifier IDC [mg/L 5.840 cific PO ₄ - use Rate LVSS/hr] .45	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari] 0 AA2 PO ₄ Release SI [mg/L/h 10.48	fier ID0 [mg/ 13.32 -P ope	SA [mg/g MI 1.1 C NOx-N /L] 200 AA2 Sp P Rele [mg/g N	LVSS/hr] 60 Clarifier ID Concen [mg 0. ecific PO ease Rate MLVSS/h 2.82	NPR (N [rr] [rr]]	Max ANA Port Release [mg/L] 20.56 A3 PO ₄ -Pease Slope mg/L/hr] 7.50	O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/g 0.32 3 Specifi Release	NO ₃ -N tration L] 22 c PO ₄ - Rate
Idays 3.96 dighest Conc Ir	SSRT Ae. SSRT Ae. SSRT Ae. AER NO ₂ -1 entration mg/L] 4.02 Dx-N AA L] PO ₄ -P P	33 75 54 57 63 64 53 44 45 45 45 41 48 TO BE TO	[mg/ 1.6 -P Uptake pe) L/hr] 28 AA1 PO ₄ Release SI [mg/L/hr] 11.00	SPUF SPUF mg/g MLV 5.38 -P AAA ope F] [n	[unitl 0.0 SS/hr] A1 Spe P Releang/g M 13	Clarifier IDC [mg/L 5.840 cific PO ₄ - lse Rate LVSS/hr] .45	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari] 0 AA2 PO ₄ Release SI [mg/L/h 10.48 AA6 PO ₄ -P	fier ID0 [mg/ 13.32 -P ope	SA [mg/g MI 1.1 C NOx-N /L] 200 AA2 Sp P Rele [mg/g N 1.1	LVSS/hr] 60 Clarifier ID Concen [mg 0.: ecific PO ease Rate MLVSS/h 2.82	NPR (N [rr] [rr]]	Max ANA Port Release [mg/L] 20.56 A3 PO ₄ -P ease Slope mg/L/hr] 7.50 (AA1) Spe	O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/g 0.32 3 Specifi Release	NO ₃ -Natration L] 22 c PO ₄ Rate
[days 3.96 dighest Conc [r	SSRT Ae. SSRT Ae. SSRT Ae. AER NO ₂ -1 entration mg/L] 4.02 Dx-N AA L] PO ₄ -P P	33 75 54 57 63 64 53 44 45 45 39 41 48 **robic MLSS** [mg/L] 1780 **PUR (PO ₄ Slo [mg/L] 7. 6 NOx-N **mg/L] 0.32	[mg/ 1.6 -P Uptake pe) L/hr] 28 AA1 PO ₄ Release SI [mg/L/hr] 11.00	SPUF SPUF mg/g MLV 5.38 -P AAA ope F] [n	[unitl 0.0 SS/hr] A1 Spe P Releang/g M 13	Clarifier IDC [mg/L 5.840 cific PO ₄ - lse Rate LVSS/hr] .45	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari] 0 AA2 PO ₄ Release SI [mg/L/h 10.48	fier ID0 [mg/ 13.32 -P ope	SA [mg/g MI 1.1 C NOx-N /L] 200 AA2 Sp P Rele [mg/g N	LVSS/hr] 60 Clarifier ID Concen [mg 0.: ecific PO ease Rate MLVSS/h 2.82	NPR (N [rr] [rr]]	Max ANA Port Release [mg/L] 20.56 A3 PO ₄ -Pease Slope mg/L/hr] 7.50	O ₄ -P	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/g 0.32 3 Specifi Release	NO ₃ -N tration [L] 22 c PO ₄ Rate SS/hr]
Idays 3.96 Highest Conc Ir	SSRT Ae SSRT Ae AER NO ₂ -1 entration mg/L] 4.02 Dx-N AA L]	33 75 54 57 63 64 53 44 45 45 45 41 48 TO BE TO	[mg/L/hr] AA1 PO ₄ Release SI [mg/L/hr] 11.00 AA5 F Release	SPUF 55 SPUF 5.38 -P	[unitl 0.0 R SS/hr] A1 Spe P Relea ng/g M 13 AA5 Spe P Relea	Clarifier IDC [mg/L 5.840 cific PO ₄ - lse Rate LVSS/hr] .45	R (NH4-N SI- [mg/L/hr] 2.17 NH ₄ -N Clari] 0 AA2 PO ₄ Release SI [mg/L/h 10.48 AA6 PO ₄ -P	[mg 13.32 -P ope	SA [mg/g MI 1.1 C NOx-N /L] 200 AA2 Sp P Rele [mg/g N 1.1	LVSS/hr] 60 Clarifier ID Concen [mg 0. ecific PO ease Rate MLVSS/h 2.82	NPR (N [rr] [rr]]	Max ANA Port Release [mg/L] 20.56 A3 PO ₄ -P ease Slope mg/L/hr] 7.50 (AA1) Spe	O ₄ -P AA P [m	SNPF ng/g MLV 1.67 Avg. ANA Concent [mg/g 0.32 3 Specifi Release ng/g MLV 9.17	NO ₃ -N tration L] 22 c PO ₄ - Rate SS/hr]

# Chem Used	FeCl3 Hrs U	sed FeCl3	Addition	Active Bios Influent Fl		ARCY	′ Flow	CR	CY Flow	NRCY Flow
[lbs]	[hr]	[r	ng/L]	[MGD]		[M	GD]	[1	MGD]	[MGD]
0	0.000	C	.000	18.84		17.	17	1	1.23	0
ANA MLTSS	S ANA % VOL	ANA MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER MI	_TSS	AER % VC	DL AER MLVSS
[mg/L]	[%]	[mg/L]	[mg/L]	[%]	[n	ng/L]	[mg/	L]	[%]	[mg/L]
940	87	818	1780	76	1	353	178	0	76	1353



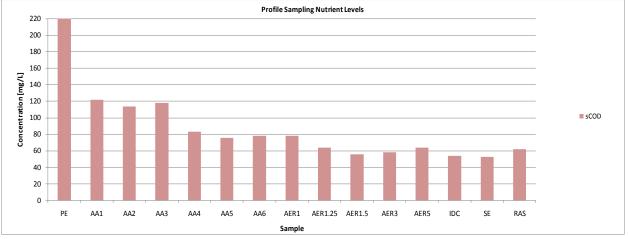


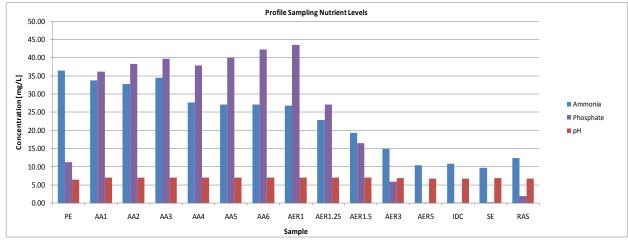


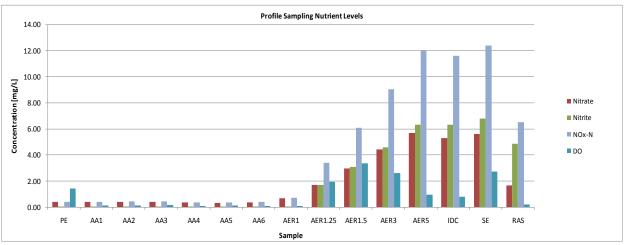
Week 13 - 10/13/09 & 10/16/09

130crs 1	PROFILE 1		ration Tank 4		0, 05													
Semmy Nam		,	HRT along BNR Process (w/out										•			•	1	
130crs 1	Date	Sample Time	Recycles)	Sample Number	Reactor Ty	pe Ammor	nia by HACH T	NT Nitrat	te by HACH	TNT N	Nitrite by HACH	NT			,	INT	DO	рН
13-01-06 726 0.90	dd-mmm-yy	h:mm	hr			n	ng/L NH3-N	n	mg/L NO3-N		mg/L NO2-N		mg/L NOx-N		mg/l	P04-P	[mg/L]	[unitless]
19-04-98 7-56 1-58 1-5	13-Oct-09	7:04	0		-		36.40		0.38		0.03		0.41		1	1.26	1.43	6.41
19.04.9 2.88 2.70 A.3 Assentic 3.49 0.41 0.02 0.3 39.55 0.09 7.01 19.04.9 3.11 4.51 A.45 Assentic 27.70 0.35 0.02 0.35 39.87 0.01 7.01 19.04.9 3.11 4.51 A.45 Assentic 27.75 0.37 0.02 0.38 39.87 0.01 7.01 19.04.9 3.11 4.51 A.45 Assentic 27.75 0.37 0.02 0.38 39.87 0.01 7.01 19.04.9 3.12 0.00 A.87 Assentic 27.75 0.37 0.02 0.38 39.87 0.01 7.01 19.04.9 3.40 0.00 A.87 Assentic 22.30 1.69 1.72 3.41 27.85 1.80 7.01 19.04.9 3.40 0.05 A.871 Assentic 9.35 2.98 3.10 1.68 1.65 3.34 7.05 19.04.9 3.40 0.35 A.871.5 Assentic 9.35 2.98 3.10 1.68 1.65 3.34 7.05 19.04.9 10.6 3.40 A.873 Assentic 19.35 2.98 3.10 1.68 1.65 3.34 7.05 19.04.9 11.50 0.00 A.875 Assentic 19.35 2.98 3.10 1.68 1.65 3.34 7.05 19.04.9 11.50 0.00 A.875 Assentic 19.35 2.98 3.10 1.68 1.65 3.34 7.05 19.04.9 11.50 0.00 A. 10 0.00 5.70 6.30 1.10 0.14 0.75 0.75 6.30 19.04.9 1.50 0.00 A.875 Assentic 1.00 0.5 0.5 0.60 0.14 0.05 0.77 6.30 19.04.9 1.50 0.00 A.875 Assentic 1.00 0.00 0.00 0.00 0.00 0.00 0.00 19.04.9 1.50 0.00					Anaerobi	;			0.39				0.41				_	
130c86 8.58 3.81					Anaerobi										3	8.34		
130498 311																	_	
19.048 9.28 5.41 M6																	_	
19.00.00																	_	
19.049 9.86																		
130ct/8 988 070																		
130ct 0 1043 348 AERS Aeroke 15.00 443 440 9.03 5.57 2.08 6.77 150ct 0 1143 6.99 AERS Aeroke 15.00 5.70 6.30 12.00 0.16 0.97 6.77 150ct 0 1150 0.00 0.16 0.97 6.77 150ct 0 1150 0.00 0.16 0.97 6.77 150ct 0 1150 0.00 0.16 0.97 6.77 150ct 0 15.00 1450 0.00 0.16 0.97 6.77 150ct 0 15.00 1450 0.00 0.16 0.97 6.77 150ct 0 15.00 0.16 0.97 6.77 15.00 0.16 0.97 0.78 0.7																		
130x86																		
130x86 1150																	_	
130tt 1450			6.99		Aerobic												_	
SOURCE Sample: SE					-													
SE EPA 365.1 EPA 350.1 EPA 353.2	13-Oct-09	14:50		SE	-		9.74		5.61		6.80		12.41		-).25	2.73	6.93
SCO By CEL	13-Oct-09	15:07		RAS	-		12.40		1.66		4.85		6.51			.88	0.19	6.75
SCOD by CEL O-PO4 NH4-N NO2.3-N NO3-N NO2-N	QA/	QC S	ample:	S	E													
Timg/L mg/L		EPA	365.1	1	EPA :	350.1	EP	A 35	3.2	EF	PA 353	.2	EPA	35 3	3.2	EP	A 35	53.2
1.1.4. 1.1.8 8.3 76 78 78 6.4 5.6 5.8 6.4 5.4 5.4 5.2 Actual SRT Aerobic MI.SS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOX-N Slope) SNPR [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [mg/L] [davs [mg/L] [davs [mg/L] [m	s	[m	g/L]	L			_			N		Г			1			
114																		
Actual SRT		- - - - -	33 76 78 78 64 56 58															
Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SNPR [days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVSS/hr] 4.43 1700 1.79 6.77 2.03 1.51 1.98 1.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Max ANA PO ₄ -P Release Avg. ANA NO ₃ -N Concentration Concentration [mg/L] AA3 PO ₄ -P Release Slope AA3 Specific PO ₄ -P Release Slope AA2 Specific PO ₄ -P Release Rate AA3 PO ₄ -P Release Slope P Release Slope P Release Slope P Release Rate [AA4 PO ₄ -P Release Rate AA5 PO ₄ -P Release Rate AA6 PO ₄ -P Release Rate					0.2	93		9.86			12.1	\Box	5.	16			6.94	
[days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVSS/hr] 4.43 1700 1.79 6.77 2.03 1.51 1.98 1.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Clarifier IDC PO ₄ -P Release Max ANA PO ₄ -P Release Avg. ANA NO ₃ -N Concentration [mg/L] [mg/			62				Т			T						Π		
Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Release Concentration Concentration Release Concentration Release Concentration Release Concentration Concentration Release Concentration Concentration Release Concentration Concentratio	Actual	SRT Aeı	obic MLSS	Average AB	R DO	Min AEI	R pH AL	JR (NH4	4-N Slop	e)	SAUR		NPR (N	Ox-N	I Slope)	SNPI	R
Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Release Concentration Release Concentration Concentration Release Concentration Concentration Concentration Release Concentration Conc										[SS/hr]	[mg		nin]	[mg		
Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Release Concentration [mg/L]	4.43	3	1700	1.79		6.77	7	2.	03		1.51			1.98			1.47	,
6.30 8.38 6.24 10.9000 11.6000 0.1 31.00 0.380 AA4 NOx-N AA6 NOx-N AA1 PO ₄ -P AA1 Specific PO ₄ - P Release Rate AA2 PO ₄ -P Release Rate AA2 Specific PO ₄ - Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate AA6 PO ₄ -P AA6 Specific PO ₄ - AA6 Specific PO ₄ - AA6 Specific PO ₄ - P Release Rate AA6 Specific PO ₄ - AA6 Specific PO ₄ - AA6 Specific PO ₄ - P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope Release Slo	Conce	entration	, ,		,				₄ -N Clar		C NOx-N	Conce	entration		Release		Concer	tration
AA4 NOx-N AA6 NOx-N AA6 NOx-N Release Slope P Release Rate [mg/L] 0.37 AA2 PO ₄ -P Release Slope P Release Slope P Release Slope P Release Slope P Release Slope P Release Rate [mg/L] 11.16 14.92 AA5 PO ₄ -P Release Slope AA6 PO ₄ -P Release Slope AA7 Specific PO ₄ - Release Slope AA8 Specific PO ₄ - Release Slope AA8 Specific PO ₄ - Release Slope AA8 Specific PO ₄ - Release Slope AA8 Specific PO ₄ - Release Slope Release Slope AA6 PO ₄ -P Release Rate Release Slope Release Slo					[mg/g				-							-		
AA4 PO ₄ -P AA4 Specific PO ₄ - AA5 PO ₄ -P AA5 Specific PO ₄ - AA6 PO ₄ -P AA6 Specific PO ₄ - (AA1) COD Uptake (AA1) Specific Release Slope P Release Rate Release Slope P Release Rate Slope COD Uptake Rate (AA1) PO ₄ -P/COD [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/mg]	AA4	NOx-N	AA6 NO	A c-N Re	ease Slo	P AA	1 Specific Release F	PO₄- Rate	Release	PO₄-P e Slop	AA2 S	pecific ease F	PO ₄ -	eleas	PO₄-P e Slope	Р	3 Specit Release	fic PO₄- e Rate
Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Slope COD Uptake Rate (AA1) PO ₄ -P/COD [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/mg]																		
	Release	Slope	P Release Rate	Release	Slope	P Releas	e Rate	Release	Slope	P Rel	ease Rate	` ′ S	Slope	COD) Uptake	Rate (
											2.63							

# Chem Use	ed	FeCl3 Use	_	FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]		[M	GD]	[MGD]	[MGD]
0		0.00	0	0.000)	18.53		17	.15	1	0.938	0
ANA MLTSS	ANA	4 % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	IXNA	MLVSS	AER MI	TSS	AER % VO	DL AER MLVSS
[mg/L]		[%]	[1	mg/L]	[mg/L]	[%]	[m	ng/L]	[mg/	L]	[%]	[mg/L]
880		85		748	1700	79	1:	343	170	0	79	1343

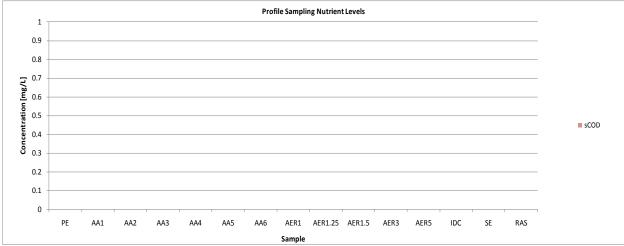


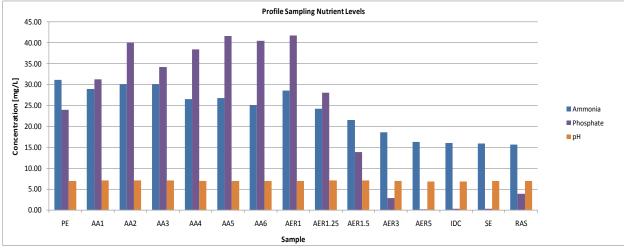


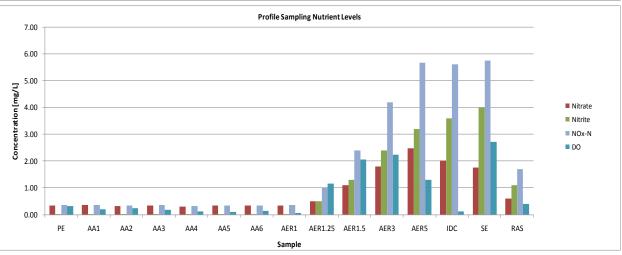


	Train 6. Ae	ration Tank 5,	Clarifier 5												
		HRT along BNR										5			
Date	Sample Time	Process (w/out Recycles)	Sample Number	Reactor Typ	e Ammonia	by HACH TNT	Nitrate by HAC	CH TNT N	Nitrite by HACH T		/ Nitrate HAC trite HACH	CH Phosphate TNT	•	DO	pН
dd-mmm-yy	h:mm	hr				L NH3-N	mg/L NO3		mg/L NO2-N		L NOx-N	mg/L l		[mg/L]	[unitless]
16-Oct-09	9:43	0	PE			1.20	0.34		0.02		0.36	23		0.32	6.89
16-Oct-09	9:58	0.83	AA1	Anaerobic		9.00	0.35		0.01		0.37	31		0.20	7.01
16-Oct-09	10:13	1.80	AA2	Anaerobic	3	0.10	0.33		0.02		0.34	40	.14	0.25	7.00
16-Oct-09	10:28	2.70	AA3	Anaerobic		0.10	0.35		0.01		0.36	34		0.18	7.00
16-Oct-09	10:43	3.61	AA4	Anaerobic		26.60	0.30		0.01		0.31	38		0.12	6.95
16-Oct-09 16-Oct-09	10:51 10:59	4.51 5.41	AA5 AA6	Anaerobic Anaerobic		.6.80 .5.10	0.34		0.01		0.35	41		0.11	6.96 6.95
16-Oct-09	11:07	0.00	AER1	Anaerobic		8.60	0.34		0.01		0.35	41		0.14	6.95
16-Oct-09	11:13	0.35	AER1.25	Aerobic		4.20	0.51		0.50		1.01	28		1.15	7.01
16-Oct-09	11:19	0.70	AER1.5	Aerobic		1.60	1.09		1.30		2.39	13	.87	2.06	7.04
16-Oct-09	12:04	3.49	AER3	Aerobic	1	8.65	1.79		2.40		4.19	2.	77	2.23	6.92
16-Oct-09	13:04	6.99	AER5	Aerobic	1	6.25	2.47		3.20		5.67	0.	09	1.30	6.76
16-Oct-09	13:10		IDC	-		6.05	2.02		3.60		5.62		24	0.12	6.81
16-Oct-09	16:10		SE	-	1	5.90	1.75		4.00		5.75	0.	25	2.72	6.96
16-Oct-09	16:30		RAS	-	1	5.65	0.60		1.10		1.70	3.	92	0.40	6.93
QA/	QC Sa	ample:	S	E											
	EPA	365.1		EPA 3	50.1	EPA	353.2	EF	PA 353.	2 E	PA 35	53.2	EP	A 35	3.2
s	COD	by CE	_	O-P	\mathcal{O}_4	NH	I ₄ -N	N	NO _{2,3} -N		NO ₃ -	·N	N	IO ₂ -1	N
	[m	g/L]		mg/	L	mg	g/L		mg/L		mg/l	L	r	ng/L	
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		0 0 0 0 0 0													
Actual S		0 0 0 0 0 0	Average A	ER DO	Min AER ;	bH AUR	(NH4-N Sic	ppe)	SAUR	N	PR (NOx	T-N Slope)	<u> </u>	SNPF	3
Actual S	SRT Aer	O O O O O O O	Average A		√lin AER [unitless		(NH4-N Slo [mg/L/hr]		SAUR mg/g MLVS		PR (NOx [mg/L		[mg/	SNPF	
	SRT Aer	O O O O O O O O O O O O O O O O O O O		-]			•				`	/min]	[mg/		
[days	SRT Aer	O O O O O O O O O O O O O O O O O O O	[mg/l	-]	[unitless		[mg/L/hr]		mg/g MLVS 1.18	SS/hr]	[mg/L 1.0	/min] 00		g MLV 0.83	SS/hr]
[days 4.90 Highest A	SRT Aer	O O O O O O O O O O O O O O O O O O O	[mg/l	-]	[unitless]	[mg/L/hr] 1.42	[1	mg/g MLVS 1.18		[mg/L 1.0 PO ₄ -P M	/min]) ₄ -P A	g MLV	SS/hr] NO ₃ -N
[days 4.90 Highest A	SRT Aer] AER NO ₂ -N entration	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Slope	_] 5	[unitless 6.76 PUR	Clarifier ID	[mg/L/hr] 1.42 C NH ₄ -N	[u	mg/g MLVS 1.18 COC NOx-N	SS/hr] larifier IDC	[mg/L 1.0 PO ₄ -P M	/min] 00 lax ANA PC Release) ₄ -P A	0.83 vg. ANA Concent	SS/hr] NO ₃ -N
[days 4.90 Highest A Conce	SRT Aer	O O O O O O O O O O O O O O O O O O O	[mg/l	S [mg/g M	[unitless 6.76]	[mg/L/hr] 1.42 C NH ₄ -N C	[1	1.18 1.0C NOx-N	SS/hr]	[mg/L 1.0 PO ₄ -P M	/min] 00 lax ANA PC) ₄ -P A	g MLV 0.83 vg. ANA	SS/hr] NO ₃ -N ration
[days 4.90 Highest A Conce	SRT Aer J AER NO ₂ -N entration g/L]	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Slope g/L/hr]	E) S [mg/g N	[unitless 6.76 PUR //LVSS/hr]	Clarifier ID	[mg/L/hr] 1.42 C NH ₄ -N C /L]	larifier ID	1.18 0C NOx-N 1/L]	larifier IDC Concentra [mg/L] 0.2	[mg/L 1.0 PO ₄ -P M tion	/min] 00 lax ANA PC Release [mg/L] 17.62) ₄ -P A	yg. ANA Concent	SS/hr] NO ₃ -N ration [L]
[days 4.90 Highest A Conce [m	SRT Aer] AER NO ₂ -N nntration 19/L]	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Slope g/L/hr] 0.01	S S [mg/g N 7	[unitless 6.76 PUR MLVSS/hr] 7.50	Clarifier ID [mg 16.05	[mg/L/hr] 1.42 C NH ₄ -N C /L] 500	larifier ID [mg 5.62	1.18 1.18 0C NOx-N 17/L] 200 AA2 Sp	larifier IDC I Concentra [mg/L] 0.2	[mg/L 1.0 PO ₄ -P M tion	/min] 00 lax ANA PC Release [mg/L] 17.62) ₄ -P A	/g MLV 0.83 vg. ANA Concent [mg/ 0.33	NO ₃ -N cration
[days 4.90 Highest A Conce [m 3	SRT Aer AER NO ₂ -N entration 19/L 20 NOx-N	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Slope g/L/hr] 0.01	S S [mg/g N 7	[unitless 6.76 PUR MLVSS/hr] 7.50 AA1 PR	Clarifier ID [mg/ 16.09 Specific PC	[mg/L/hr] 1.42 C NH ₄ -N Cl /L] 500 D ₄ - AA2 Relea	Igentifier ID Img 5.62 PO ₄ -P ase Slop	1.18 1.18 COC NOx-N 19/L1 200 AA2 Sp P Rel	larifier IDC I Concentra [mg/L] 0.2 Decific PO4 ease Rate	[mg/L 1.0 PO ₄ -P M tion AA Relea	/min] 00 lax ANA PC Release [mg/L] 17.62 3 PO ₄ -P ase Slope	AA3	/g MLV 0.83 vg. ANA Concent [mg/ 0.33	NO ₃ -N rration L] 33 c PO ₄ - Rate
[days 4.90 Highest # Conce [m 3	SRT Aer] AER NO ₂ -N nntration 19/L]	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Slope g/L/hr] 0.01	S S [mg/g N 7	[unitless 6.76 PUR MLVSS/hr] 7.50 AA1 PR	Clarifier ID [mg 16.05	[mg/L/hr] 1.42 C NH ₄ -N Cl /L] 500 D ₄ - AA2 e Relea hr] [m	larifier ID [mg 5.62	1.18 1.18 COC NOx-N 19/L] 200 AA2 Sp P Religing/g	larifier IDC I Concentra [mg/L] 0.2	Img/L 1.(PO ₄ -P M tion AA Rele:	/min] 00 lax ANA PC Release [mg/L] 17.62	AA3	/g MLV 0.83 vg. ANA Concent [mg/ 0.33	NO ₃ -N rration L] 33 c PO ₄ - Rate SS/hr]
[days 4.90 Highest # Conce [m 3	SRT Aer] AER NO ₂ -N ontration [g/L] .20 NOx-N [g/L] .31	O O O O O O O O O O O O O O O O O O O	[mg/l 1.36 Uptake Sloppi J/L/hr] 0.01	S [mg/g N 7 7 7 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	[unitless 6.76 PUR MLVSS/hr] 7.50 AA1 P R [mg/	Clarifier ID [mg/ 16.06 Specific PC telease Rate g MLVSS/II -0.38	[mg/L/hr] 1.42 C NH ₄ -N C /L] 500 D ₄ - AA2 e Relea hr] [m	Igraphic ID Image 5.62 PO ₄ -P Isse Slop g/L/hr] 2.62	1.18 1.18 OC NOx-N 1.200 AA2 Sp P Rele [mg/g	larifier IDC Concentra [mg/L] 0.2 ecific PO ₄ ease Rate MLVSS/hr 16.88	[mg/L 1.0 PO ₄ -P M tition AA Rele:	/min] 200 dax ANA PC Release [mg/L] 17.62 3 PO ₄ -P ase Slope ng/L/hr] 4.48	AA3 P F	/g MLV 0.83 vg. ANA Concent [mg/ 0.33 s Specifii Release	NO ₃ -N rration L] 33 c PO ₄ - Rate SS/hr]
[days 4.90 A.90 AA4 P	SRT Aer] AER NO ₂ -N ontration 19/L] .20 NOx-N g/L] .31 O ₄ -P A	O O O O O O O O O O O O O O O O O O O	[mg/l] 1.36 Uptake Sloppi 3/L/hr] 9.01 4- AA5 P	S [mg/g N 7 7 7 7 8 8 8 8 9 9 9 9 9 9 9 9 9 9 9 9	[unitless 6.76 PUR MLVSS/hr] AA1 P R [mg/	Clarifier ID [mg/ 16.05 Specific PC telease Rat /g MLVSS/I -0.38 PO ₄ - A	[mg/L/hr] 1.42 C NH ₄ -N Cl /L] 500 D ₄ - AA2 e Relea hr] [m 1	larifier ID [mg 5.62 2 PO ₄ -P use Slop g/L/hr] 2.62 AA6 Sp	1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18	larifier IDC Concentra [mg/L] 0.2 Decific PO4 Dease Rate MLVSS/hr 16.88	[mg/L 1.(PO ₄ -P M tion AA Rele:	/min] Dio dax ANA PC Release [mg/L] 17.62 3 PO ₄ -P ase Slope ng/L/hr] 4.48 (AA1) Speci	AA3 P F [mg	/g MLV 0.83 vg. ANA Concent [mg/ 0.33 S Specific Release y/g MLV 5.99	NO ₃ -N ration L] 33 c PO ₄ - Rate SS/hr]
[days 4.90 Highest # Conce [m 3	SRT Aer] AER NO ₂ -N ontration 19/L] .20 NOx-N g/L] .31 O ₄ -P A	O O O O O O O O O O O O O O O O O O O	[mg/l] 1.36 Uptake Sloppi 3/L/hr] 9.01 Re-N Re	S (mg/g N 7 7 7 7 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	[unitless 6.76 PUR MLVSS/hr] 7.50 AA1 P R [mg/	Clarifier ID [mg/ 16.05 Specific PC telease Rat /g MLVSS/I -0.38 PO ₄ - A	[mg/L/hr] 1.42 C NH ₄ -N C /L] 500 D ₄ - AA2 e Relea hr] [m	larifier ID [mg 5.62 2 PO ₄ -P use Slop g/L/hr] 2.62 AA6 Sp	1.18 1.18 OC NOx-N 1.200 AA2 Sp P Rele [mg/g	larifier IDC Concentra [mg/L] 0.2 ecific PO ₄ ease Rate MLVSS/hr 16.88	[mg/L 1.(PO ₄ -P M tion AA Rele:	/min] 200 dax ANA PC Release [mg/L] 17.62 3 PO ₄ -P ase Slope ng/L/hr] 4.48	AA3 P F [mg	/g MLV 0.83 vg. ANA Concent [mg/ 0.33 s Specifii Release	NO ₃ -N ration L] 33 c PO ₄ - Rate SS/hr]
[days 4.90 Highest A Conce [m 3 3 AA4 [m 0.44 P]	SRT Aer AER NO2-N entration G/L .20 NOx-N G/L .31 O ₄ -P A	O O O O O O O O O O O O O O O O O O O	Uptake Slope g/L/hr] 0.01 4-AA5 P Release	Solution Solution	[unitless 6.76 PUR MLVSS/hr] AA1 P R [mg/	Clarifier ID [mg/ 16.05 Specific PC celease Rat g MLVSS/I -0.38 PO ₄ - A Rate Rel	[mg/L/hr] 1.42 C NH ₄ -N Cl /L] 500 D ₄ - AA2 e Relea hr] [m 1	larifier ID [mg 5.62 PO ₄ -P ase Slop g/L/hr] 2.62 AA6 Sp P Rel	1.18 1.18 1.18 1.18 1.18 1.18 1.18 1.18	larifier IDC Concentra [mg/L] 0.2 Decific PO4 Dease Rate MLVSS/hr 16.88	[mg/L 1.(PO ₄ -P Mtion ARele:	/min] Dio dax ANA PC Release [mg/L] 17.62 3 PO ₄ -P ase Slope ng/L/hr] 4.48 (AA1) Speci	AA3 P F [mg	/g MLV 0.83 vg. ANA Concent [mg/ 0.33 S Specific Release y/g MLV 5.99	NO ₃ -N ration L] 33 c PO ₄ -Rate SS/hr]

# Chem Us	ed	FeCl3 Use	_	FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]		[M	GD]]	MGD]	[MGD]
0		0.00	0	0.000)	20.22		17	.21	1	1.905	0
ANA MLTSS	ANA	A % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	IXNA	MLVSS	AER ML	TSS	AER % VC	DL AER MLVSS
[mg/L]		[%]	[mg/L]	[mg/L]	[%]	[m	ig/L]	[mg/	L]	[%]	[mg/L]
840		89		748	1540	78	1:	201	1540	0	78	1201



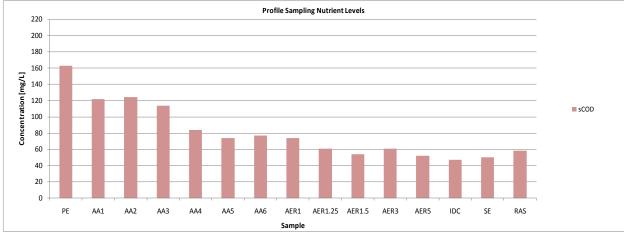


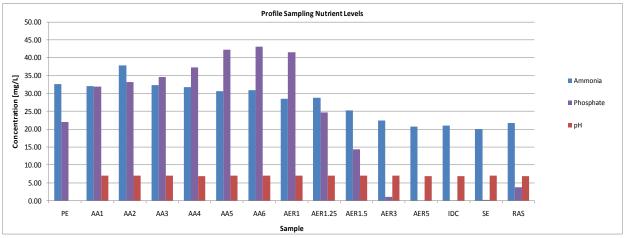


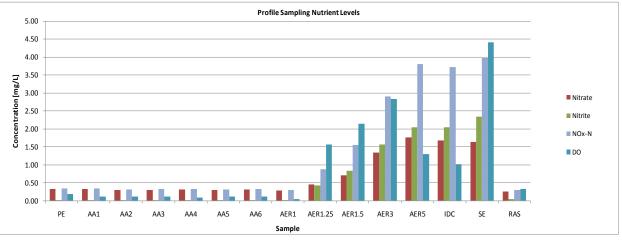
Week 14 – 10/19/09 & 10/21/09

Fit along Bits Process years Sample Number Roader Type Ammonia by MCN RT Niese by MCN RT			ration Tank 4,	Clarifier 4													
Description Processing Name Processing Name Report Type Ammoria by HOH NIT Name by HOH NIT			HRT along BNR														
Sementy Imm											NO	0x-N by Nitrate I	HACH	Phosphate	by HACH	4	
Second Sep 0	Date	Sample Time	Recycles)	Sample Number	r Reactor Typ	e Ammonia	y HACH TNT	Nitrate by H.	ACH TNT	Nitrite by HAC	1 TNT	+ Nitrite HACI	Н	TN	Ť	DO	рН
90-049 937	dd-mmm-yy	h:mm	hr			mg/l	. NH3-N	mg/L No	D3-N	mg/L NO2	N	mg/L NOx-N		mg/L P	04-P	[mg/L]	[unitless]
190-049 1007 2.56 A31	19-Oct-09	8:50	0	PE	-	3	2.60	0.32	2	0.02		0.34		22.0)3	0.18	701
19-00-89 10-07 2-56																	7.02
99-07-08 1937 3-41 A44 Areacobic 318.0 0.31 0.01 0.33 9.38 0.09				1													7.02
1904.06 1050																	7.01 6.93
90-08-9 11-93 512 A66 Arearchic 30.90 0.31 0.01 0.32 4.07 0.11																	6.96
1904-09 11-16 0.00																	6.96
1904-96 11-32		11:16	0.00	AER1	Aerobic			0.29	9	0.01		0.30		41.6	61	0.05	6.95
1900;190;17 331																	7.06
19-00-96 1317 6-61				1													7.08
190c49 1326 1326 1326 1524 152 154																	6.96
90ct/9 1656			6.61														6.93
SOOD by CEL					-												6.91
CAPQC Sample: SE					-											_	7.08
SCOD by CEL			mple		· ·		1.00	0.20)	0.04		0.30		3.1	0	0.32	6.88
SCOD by CEL	QA.					50.1	EDA	252.0		ED A 25	2.0	EDA	252		ED	A 25	2.2
Times/L Time		EPA	303.1		EPA 3	30.1	EPA	333.2	+-	EPA 33.	5.4	EPA	333		EP	A 33	3.2
Img/L																	
Img/L mg/L	s	COD	by CE	L	O-P	O_4	NH	I ₄ -N		NO _{2.3} -	Ν	NO	3-N	-	N	1 O 2-I	V
163 122 124 114 84 774 777 74 61 54 61 52 47 50 0.112 21.2 3.49 1.05 2.44 Sequence of the proof								-									
124								2'		<u></u>			,			8	
114 84 74 77 74 77 74 61 54 61 52 47 50 58																	
R4 Fig. Fi																	
74 77 74 61 54 61 52 47 50 0.112 21.2 3.49 1.05 2.44 58 Actual SRT																	
T-4																	
61 54 61 52 477 50 O.112 21.2 3.49 1.05 2.44 58 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SNPR [days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS] 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.71 0.47 Highest AER NO _{z-N} PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Release Rote Release Slope P Release Rate Rote Release Slope P Release Rote Rote Release Slope P Release Rote Rote Rote Rote Rote Rote Rote Rot				_													
S-4 G-1 S-2		-	74														
61 52 47 50 0.112 21.2 3.49 1.05 2.44 58 Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SNPR [days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS] 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration [mg/L] [mg/L		- - -	74 77 74														
S2		- - -	74 77 74 51														
Actual SRT Aerobic MLSS Average AER DO Min AER pH AUR (NH4-N Slope) SAUR NPR (NOx-N Slope) SNPR		-	74 77 74 61 54														
Actual SRT		-	74 77 74 61 54														
Actual SRT		-	74 77 74 61 54 61														
[days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Max ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P Release Rate Release Slope P Release Rate Relea		-	74 77 74 61 54 61 52		0.11	12	2:	1.2	T	3.49		1.0	05			2.44	
[days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Max ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P Release Rate Release Slope P Release Rate Relea		- - - - - - - - - - - - - - - - - - -	74 77 74 61 54 61 52 47		0.11	12	2:	1.2		3.49		1.0	05			2.44	
[days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Max ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P Release Rate Release Slope P Release Rate Relea		- - - - - - - - - - - - - - - - - - -	74 77 74 61 54 61 52 47		0.1	2	2:	1.2	T	3.49	I	1.0	05	T		2.44	
[days] [mg/L] [mg/L] [unitless] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/min] [mg/g MLVS 4.93 1910 1.58 6.93 1.16 0.77 0.71 0.47 Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Max ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Avg. ANA PO ₄ -P Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P Release Rate Release Slope P Release Rate Relea		- - - - - - - - - - - - - - - - - - -	74 77 74 61 54 61 52 47		0.11	12	2.	1.2		3.49	I	1.0	05			2.44	
A.93	Actual S		74 77 74 61 61 62 47 60	Average A					Slope)		TR.			Slope)			
Highest AER NO ₂ -N Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Clarifier IDC PO ₄ -P Concentration Max ANA PO ₄ -P Avg. ANA Concentration Avg. ANA Concentration [mg/L] AA3 PO ₄ -P Release Slope P Release Slope </td <td></td> <td>SRT Aer</td> <td>74 77 74 61 63 61 65 62 47 60 68</td> <td></td> <td>ER DO</td> <td>Min AER I</td> <td>oH AUR</td> <td>(NH4-N S</td> <td>Slope)</td> <td>SAL</td> <td></td> <td>NPR (N</td> <td>Ox-N</td> <td></td> <td></td> <td>SNPF</td> <td></td>		SRT Aer	74 77 74 61 63 61 65 62 47 60 68		ER DO	Min AER I	oH AUR	(NH4-N S	Slope)	SAL		NPR (N	Ox-N			SNPF	
Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P AA2 Specific PO ₄ -P AA3 PO ₄ -P AA3 PO ₄ -P AA3 Specific P Release Release Slope P Release Release Slope P Release Release Slope P Release Release Release Slope P Release Release <td>[days</td> <td></td> <td>74 77 74 61 64 61 52 47 50 58</td> <td>[mg/</td> <td>ER DO</td> <td>ا Min AER [unitless</td> <td>oH AUR</td> <td>(NH4-N S [mg/L/hr]</td> <td>Slope)</td> <td>SAL [mg/g ML</td> <td>/SS/hr]</td> <td>NPR (N</td> <td>Ox-N g/L/mi</td> <td></td> <td></td> <td>SNPF /g MLV</td> <td></td>	[days		74 77 74 61 64 61 52 47 50 58	[mg/	ER DO	ا Min AER [unitless	oH AUR	(NH4-N S [mg/L/hr]	Slope)	SAL [mg/g ML	/SS/hr]	NPR (N	Ox-N g/L/mi			SNPF /g MLV	
Concentration PUR (PO ₄ -P Uptake Slope) SPUR Clarifier IDC NH ₄ -N Clarifier IDC NOx-N Concentration Release Concentration [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P AA2 Specific PO ₄ -P AA3 PO ₄ -P AA3 PO ₄ -P AA3 Specific P Release Release Slope P Release Release Slope P Release Release Slope P Release Release Release Slope P Release Release <td>[days</td> <td></td> <td>74 77 74 61 64 61 52 47 50 58</td> <td>[mg/</td> <td>ER DO</td> <td>ا Min AER [unitless</td> <td>oH AUR</td> <td>(NH4-N S [mg/L/hr]</td> <td>Slope)</td> <td>SAL [mg/g ML</td> <td>/SS/hr]</td> <td>NPR (N</td> <td>Ox-N g/L/mi</td> <td></td> <td></td> <td>SNPF /g MLV</td> <td></td>	[days		74 77 74 61 64 61 52 47 50 58	[mg/	ER DO	ا Min AER [unitless	oH AUR	(NH4-N S [mg/L/hr]	Slope)	SAL [mg/g ML	/SS/hr]	NPR (N	Ox-N g/L/mi			SNPF /g MLV	
[mg/L] [mg/L]hr] [mg/g MLVSS/hr] [mg/L] AA3 PO ₄ -P AA3 Specific PO ₄ -P AA2 PO ₄ -P AA2 Specific PO ₄ -P AA3 PO ₄ -P AA3 Specific P Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Release Rate Release Slope P Release Rat	[days 4.93	SRT Aer	74 77 74 61 64 61 52 47 50 58	[mg/	ER DO	ا Min AER [unitless	oH AUR	(NH4-N S [mg/L/hr]	Slope)	SAL [mg/g ML	VSS/hr] 7	NPR (N	Ox-N g/L/mi 0.71	in]	[mg/	SNPF /g MLV 0.47	SS/hr]
2.04 9.71 6.43 21.0000 3.7200 0.1 21.05 0.30 AA4 NOx-N AA6 NOx-N AA1 PO ₄ -P Release Slope AA1 Specific PO ₄ - Release Slope AA2 PO ₄ -P Release Rate Release Slope P Release Slope P Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release	[days 4.93	SRT Aet	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910	[mg/	ER DO	Min AER p [unitless 6.93	oh Aur	(NH4-N S [mg/L/hr] 1.16	. ,	SAL [mg/g ML [*] 0.7	/SS/hr] 7 Clarifier	NPR (NI	Ox-N g/L/mi 0.71 Max	in] ANA PO	[mg/	SNPF /g MLV 0.47	SS/hr]
AA4 NOx-N	[days 4.93 Highest A	SRT Aer	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5	ER DO I	Min AER programmer [unitless 6.93]	OH AUR	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N	Clarifier	SAL [mg/g ML 0.7 IDC NOx-N	VSS/hr] 7 Clarifier Conce	NPR (NI [mç	Ox-N g/L/mi 0.71 Max	ANA PO	[mg/	SNPF /g MLV 0.47 vg. ANA Concent	SS/hr] NO ₃ -N
AA4 NOx-N AA6 NOx-N Release Slope P Release Rate Release Slope P Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Slope P Release [mg/L] [mg/L] [mg/L/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/g	[days 4.93 Highest A Conce	SRT Aer	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5 Uptake Slop g/L/hr]	ER DO L] B S e) S	Min AER I [unitless 6.93 PUR MLVSS/hr]	OH AUR Clarifier ID	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N	Clarifier [r	SAL [mg/g ML ⁻ 0.7 IDC NOx-N ng/L]	VSS/hr] 7 Clarifier Conce	NPR (NI [mç IDC PO ₄ -P entration ng/L]	Ox-N g/L/mi 0.71 Max	ANA PO	[mg/	SNPF /g MLV 0.47 avg. ANA Concent	SS/hr] NO ₃ -N ration
AA4 NOx-N AA6 NOx-N Release Slope P Release Rate Release Slope P Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Rate Release Slope P Release [mg/L] [mg/L] [mg/L/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/g	[days 4.93 Highest A Conce	SRT Aer	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5 Uptake Slop g/L/hr]	ER DO L] B S e) S	Min AER I [unitless 6.93 PUR MLVSS/hr]	OH AUR Clarifier ID	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N	Clarifier [r	SAL [mg/g ML ⁻ 0.7 IDC NOx-N ng/L]	VSS/hr] 7 Clarifier Conce	NPR (NI [mç IDC PO ₄ -P entration ng/L]	Ox-N g/L/mi 0.71 Max	ANA PO	[mg/	SNPF /g MLV 0.47 avg. ANA Concent	SS/hr] NO ₃ -N ration
[mg/L] [mg/L] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr]	[days 4.93 Highest A Conce	SRT Aer	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5	ER DO L] 3 (e) S [mg/g l	Min AER [unitless 6.93 PUR MLVSS/hr] 6.43	Clarifier ID [mg/21.00	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N 'L']	Clarifier [r	SAL [mg/g ML] 0.7 IDC NOx-N ng/L] 7200	VSS/hr] 7 Clarifier Conce	NPR (NI [mg] IDC PO ₄ -P entration mg/L] 0.1	Ox-N g/L/mi 0.71 Max	ANA PO. Release [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 wg. ANA Concent [mg/ 0.30	NO ₃ -N cration
0.33 0.32 0.10 0.13 9.17 11.58 6.45 8.14 AA4 PO ₄ -P AA4 Specific PO ₄ - AA5 PO ₄ -P AA5 Specific PO ₄ - AA6 PO ₄ -P AA6 Specific PO ₄ - (AA1) COD Uptake (AA1) Specific	[days 4.93 Highest A Conce	SRT Aer	74 77 74 61 61 62 47 60 68 60 68 60 68 60 69 69 69 69 69 69 69 69 69 69 69 69 69	[mg/ 1.5	ER DO	Min AER ; [unitless 6.93 PUR MLVSS/hr] 6.43	Clarifier ID [mg/21.00	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N /L] 1000	Clarifier [r 3.	SAL [mg/g ML' 0.7 IDC NOx-N ng/L] 7200	VSS/hr] 7 Clarifier Conce	NPR (NI [mg IDC PO ₄ -P entration ng/L] 0.1	Ox-N g/L/mi 0.71 Max F	ANA PORRelease [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 vg. ANA Concent [mg/ 0.30	NO ₃ -N cration
AA4 PO ₄ -P AA4 Specific PO ₄ - AA5 PO ₄ -P AA5 Specific PO ₄ - AA6 PO ₄ -P AA6 Specific PO ₄ - (AA1) COD Uptake (AA1) Specific	[days 4.93 Highest A Conce	SRT Aer SBRT Ae	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5 Uptake Slop g/L/hr] 3.71	ER DO L] 3 e) S [mg/g l	Min AER [unitless 6.93 PUR MLVSS/hr] 6.43 AA1 PR	Clarifier ID [mg/21.0((NH4-N S [mg/L/hr] 1.16 C NH ₄ -N /L] 000 D ₄ - AA e Relo	Clarifier [r 3. A2 PO ₄ ease Sle	SAL [mg/g ML' 0.7 IDC NOx-N mg/L] 7200 P AA2 P F	7 Clarifier Conce	NPR (NI [mg] IDC PO ₄ -P entration mg/L] 0.1 PO ₄ - Rate Rivers	Ox-N g/L/mi 0.71 Max F	ANA PORelease [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 vg. ANA Concent [mg/ 0.30	NO ₃ -N tration L] 09 c PO ₄ - Rate
	[days 4.93 Highest A Conce [m 2	SRT Aer sil	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5 Uptake Slop g/L/hr] 3.71	ER DO L] 3 e) S [mg/g] AA1 PO ₄ -Felease Slop [mg/L/hr]	Min AER [unitless 6.93 PUR MLVSS/hr] 6.43 AA1 PR	Clarifier ID [mg/21.00	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N /L] 000 D ₄ - AA e Relo	Clarifier [r 3. A2 PO ₄ - ease Slo	SAL [mg/g ML' 0.7 IDC NOx-N mg/L] 7200 P AA2 P F	7 Clarifier Conce	NPR (NI [mg] IDC PO ₄ -P entration mg/L] 0.1 PO ₄ - Rate Rivers	Ox-N Ox-N Ox-N Ox-N Max F AA3 P elease [mg/L	ANA PO Release [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 vg. ANA Concent [mg/ 0.30 8 Specifi Release	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]
	[days 4.93 Highest A Conce [m 2	SRT Aer sil	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P	[mg/ 1.5 Uptake Slop g/L/hr] 3.71	ER DO L] 3 e) S [mg/g] AA1 PO ₄ -Felease Slop [mg/L/hr]	Min AER [unitless 6.93 PUR MLVSS/hr] 6.43 AA1 PR	Clarifier ID [mg/21.00	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N /L] 000 D ₄ - AA e Relo	Clarifier [r 3. A2 PO ₄ - ease Slo	SAL [mg/g ML' 0.7 IDC NOx-N mg/L] 7200 P AA2 P F	7 Clarifier Conce	NPR (NI [mg] IDC PO ₄ -P entration mg/L] 0.1 PO ₄ - Rate Rivers	Ox-N Ox-N Ox-N Ox-N Max F AA3 P elease [mg/L	ANA PO Release [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 vg. ANA Concent [mg/ 0.30 8 Specifi Release	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]
Release Slope P Release Rate Release Slope P Release Rate Release Slope P Release Rate Slope COD Uptake Rate (AA1) PO4	Edays 4.93 Highest A Conce [m 2 AA4	SRT Aer SRT Aer NOz-N NOx-N 19/L] 33	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P [mg/L] 0.32	[mg/ 1.5	ER DO L] B S [mg/g l] AA1 PO ₄ -Felease Slop [mg/L/hr] 0.10	Min AER p [unitless 6.93 PUR MLVSS/hr] 6.43 AA1 P R [mg/	Clarifier ID [mg 21.00 Specific PC elease Rat g MLVSS/I 0.13	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N /L] 000 D ₄ - AA Relent	Clarifier [r 3. A2 PO ₄ ease Slo	SAL [mg/g ML' 0.7 IDC NOx-N ng/L] 7200 P AA2 P F F	Clarifier Conce [m] Specific telease F g MLVS 11.58	NPR (NI [mg] IDC PO ₄ -P entration mg/L] 0.1 PO ₄ - Rate Rivers	Ox-N Ox-N Ox-N Ox-N Max F AA3 P elease [mg/L	ANA PO Release [mg/L] 21.05	[mg/	SNPF /g MLV 0.47 vg. ANA Concent [mg/ 0.30 8 Specifi Release	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]
	Edays 4.93 Highest A Conce [m 2 AA4 AA4 AA4 AA4 AA4	SRT Aer (1) (2) (3) (4) (4) (4) (4) (4) (4) (4) (4) (4) (4	74 77 74 61 61 654 61 652 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P [m] 9 AA6 NO [mg/L] 0.32	[mg/ 1.5] Uptake Slop g/L/hr] 9.71 c-N R	ER DO L] B S [mg/g l] AA1 PO ₄ -F elease Slop [mg/L/hr] 0.10	Min AER p [unitless 6.93 PUR MLVSS/hr] 5.43 AA1 P R [mg/	Clarifier ID [mg/21.00 Specific PC elease Rat g MLVSS/I 0.13 PO ₄ - A	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N [L] 1000 D ₄ - AA Relenting [mathematical part of the control of the c	Clarifier [r 3. A2 PO ₄ ease Sle mg/L/hr 9.17 AA6	SAL [mg/g ML' 0.7 IDC NOx-N ng/L] 7200 P AA2 P F F [mg	Clarifier Conce [m] Specific telease F g MLVS 11.58	NPR (NI [mg] IDC PO ₄ -P entration ng/L] 0.1 PO ₄ -Rate Resides Reside	Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N	ANA PO. Release [mg/L] 21.05 PO ₄ -P Slope _/hr] 45	a-P A	SNPF /g MLV 0.47 ovg. ANA Concent [mg/ 0.30 8 Specifit Release /g MLV 8.14	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]
[mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/L/hr] [mg/g MLVSS/hr] [mg/m	Edays 4.93 Highest A Conce [m 2 AA4 AA4 AA4 AA4 AA4	SRT Aer (1) (2) (3) (4) (4) (4) (4) (4) (4) (4) (4) (4) (4	74 77 74 61 61 654 61 652 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P [m] 9 AA6 NO [mg/L] 0.32	[mg/ 1.5] Uptake Slop g/L/hr] 9.71 c-N R	ER DO L] B S [mg/g l] AA1 PO ₄ -F elease Slop [mg/L/hr] 0.10	Min AER p [unitless 6.93 PUR MLVSS/hr] 5.43 AA1 P R [mg/	Clarifier ID [mg/21.00 Specific PC elease Rat g MLVSS/I 0.13 PO ₄ - A	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N [L] 1000 D ₄ - AA Relenting [mathematical part of the control of the c	Clarifier [r 3. A2 PO ₄ ease Sle mg/L/hr 9.17 AA6	SAL [mg/g ML' 0.7 IDC NOx-N ng/L] 7200 P AA2 P F F [mg	Clarifier Conce [m] Specific telease F g MLVS 11.58	NPR (NI [mg] IDC PO ₄ -P entration ng/L] 0.1 PO ₄ -Rate Resides Reside	Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N	ANA PO. Release [mg/L] 21.05 PO ₄ -P Slope _/hr] 45	a-P A	SNPF /g MLV 0.47 ovg. ANA Concent [mg/ 0.30 8 Specifit Release /g MLV 8.14	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]
2.98 1.97 1.15 0.76 2.87 1.90 0.81 1.03 0.10	Edays A.93 Highest A Conce France France Conce France Conce France France Conce France Conce France SRT Aer SRT	74 77 74 61 61 62 47 60 68 obic MLSS [mg/L] 1910 PUR (PO ₄ -P [mg/L] 0.32 A4 Specific PC P Release Ratumg/g MLVSS/h	[mg/ 1.5 Uptake Slop g/L/hr] 9.71 c-N R	ER DO L] 3 e) S [mg/g l] AA1 PO ₄ -F elelease Slop [mg/L/hr] 0.10 PO ₄ -P e Slope	Min AER F [unitless 6.93 PUR MLVSS/hr] 5.43 AA1 P R [mg/	Clarifier ID [mg/21.00] Specific PC elease Rat g MLVSS/I 0.13 PO4- A Rel	(NH4-N S [mg/L/hr] 1.16 C NH ₄ -N [L] 1000 A6 PO ₄ -P ease Slope mg/L/hr]	Clarifier [r 3. A2 PO ₄ - ease Sle mg/L/hr 9.17 AA66	SAL [mg/g ML' 0.7 IDC NOx-N ng/L] 7200 P AA2 ope P F [mg Specific PO4 telease Rate	Clarifier Conce [m] Specific Release F (g MLVS) 11.58	NPR (NI [mg IDC PO4-P entration ng/L] 0.1 PO4-Rate Resolution Robert Resolution	Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N Ox-N	ANA PO. Release [mg/L] 21.05 PO ₄ -P Slope [/hr] 45 Uptake R	Img/	SNPF /g MLV 0.47 ovg. ANA Concent [mg/s 0.30 8 Specific Release p/g MLV 8.14	NO ₃ -N tration L] 99 c PO ₄ -Rate SS/hr]	

# Chem Use	ed	FeCl3 Use	_	FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]		[M	GD]]	MGD]	[MGD]
0		0.00	0	0.000)	19.57		17	.26	1	1.602	0
ANA MLTSS	ANA			MLVSS	ANX MLTSS			MLVSS				DL AER MLVSS
[mg/L]		[%]	[1	mg/L]	[mg/L]	[%]	[m	ng/L]	[mg/	L]	[%]	[mg/L]
890		89		792	1910	79	1:	509	191	0	79	1509

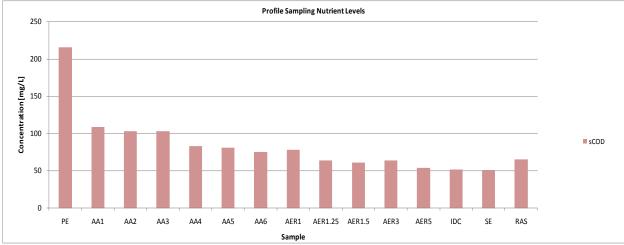


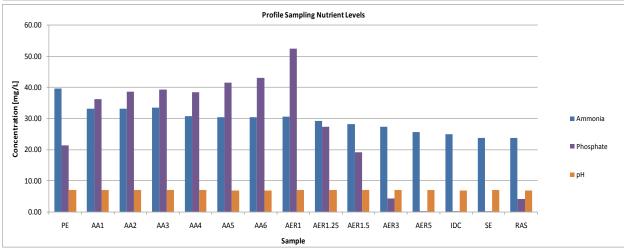


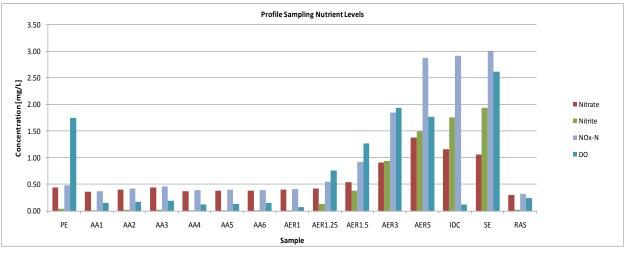


ROFILE 2	Train 5. Ae	ration Tank 5,	Siai iliti 3												
		HRT along BNR													
Date	Sample Time	Process (w/out Recycles)	Sample Numbe	Reactor Ty	pe Ammonia	a by HACH TNT	Nitrate by HACH	H TNT Nitrit	e by HACH TNT	NOx-N by Ni + Nitrite			e by HA(T844	CH DO	pН
dd-mmm-yy	h:mm	hr			mg	JL NH3-N	mg/L NO3-N	N r	mg/L NO2-N	mg/L N	IOx-N	mg/L	P04-P	[mg/L]	[unitless
21-Oct-09	6:53	0	PE	-	,	39.70	0.44		0.04	0.4	8	·	1.37	1.75	7.00
21-Oct-09	7:07	0.89	AA1	Anaerobi	С	33.10	0.36		0.01	0.3	7	36	5.22	0.15	6.99
21-Oct-09	7:37	1.71	AA2	Anaerobi	С	33.10	0.40		0.02	0.4	2	38	3.67	0.17	7.00
21-Oct-09	8:07	2.56	AA3	Anaerobi		33.50	0.44		0.02	0.4	6		9.32	0.19	6.98
21-Oct-09	8:37	3.41	AA4	Anaerobi		30.70	0.37		0.01	0.3	19	38	3.51	0.12	6.97
21-Oct-09	8:52	4.27	AA5	Anaerobi	С	30.40	0.38		0.01	0.4	0	41	1.61	0.13	6.91
21-Oct-09	9:07	5.12	AA6	Anaerobi	С	30.40	0.38		0.01	0.3	19	43	3.07	0.15	6.93
21-Oct-09	9:22	0.00	AER1	Aerobic		30.60	0.40		0.01	0.4	1	52	2.54	0.07	6.94
21-Oct-09	9:30	0.33	AER1.25	Aerobic		29.30	0.42		0.13	0.5	5	27	7.33	0.76	6.97
21-Oct-09	9:38	0.66	AER1.5	Aerobic		28.20	0.54		0.38	0.9	12	19	9.17	1.27	7.02
21-Oct-09	10:23	3.31	AER3	Aerobic		27.40	0.91		0.94	1.8	5	4	.21	1.94	6.98
21-Oct-09	11:23	6.61	AER5	Aerobic		25.70	1.38		1.50	2.8	8	0	.10	1.77	6.94
21-Oct-09	11:37		IDC	-		24.90	1.16		1.76	2.9	12	0	.15	0.12	6.91
21-Oct-09	14:37		SE			23.70	1.06		1.94	3.0	10	0	.17	2.62	7.01
21-Oct-09	14:59		RAS	-		23.70	0.30		0.03	0.3			.11	0.24	6.92
		ample:		E .		23.10	0.30		0.03	0.0	12	4	.11	0.24	0.92
QA		365.1		EPA:	350.1	EPA	353.2	EPA	X 353.2	EP	A 35	3.2	EP	PA 35	3.2
					350.1		555.2		1 000.2			2.2		11 00	<u> </u>
_		 OF		O-P		NII	I4-N	NIC	D _{2.3} -N		1 O 3-I	ν. Τ		VO 2-l	N.T
S	SCOD	by CE	_	О-Р	O ₄	1/1	14-17	110	J2,3-1N	1.	103-1	. 7	1	VO 2-1	
		_			/T		/ T		/ T		/T			/1	
	[m	ıg/L]		mg	/L	mg	g/L	n	ng/L	1	mg/L	_		mg/L	_
	[m	_		mg	/L	mş	g/L	n	ng/L	1	mg/L	_		mg/L	_
	[m 2 1 1	1g/L] 216 09 03		mg	/L	mg	g/L	n	ng/L		mg/L	,		mg/L	_
	[m 2 1 1	ng/L] 216 09 03 03		mg	/L	mş	g/L	m	ng/L	1	mg/L			mg/L	
	[m 2 1 1 1	19/L] 216 09 03 03 83		mg	/L	mg	g/L	n	ng/L		mg/L			mg/L	
	[m 2 1 1 1	ng/L] 216 09 03 03		mg	/L	mg	g/L	n	ng/L	1	mg/L			mg/I	
	[m 2 1 1 1	19/L] 216 09 03 03 83 81 75		mg	/L	mş	g/L	n	ng/L	,	mg/L			mg/I	
	[m 2 1 1 1	19/L] 216 09 03 03 83 81 75 78		mg	/L	mş	g/L	n	ng/L		mg/L			mg/I	
	[m 2 1 1 1 3	19/L] 216 09 03 03 83 81 75		mg	/ L	mş	g/L	n	ng/L	,	mg/I			mg/L	
	[m 2 1 1 1 3	9/L] 216 09 03 03 83 81 75 78 64		mg	<u>/L</u>	mş	g/L	m	ng/L	,	mg/I			mg/L	-
	[m 2 1 1 1 1 	9/L] 216 09 03 03 83 81 75 78 64 61 64 54		mg	/ L									mg/L	
	[m 2 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54		mg	/L		g/L 3.8		ng/L		mg/I			<u>mg/I</u>	
	[m 2 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54		mg	/L										
	[m 2 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54		mg	/L										
Actual	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54 52	Average A			23	3.8	2	2.35		0.35			2	
Actual	[m 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54 52 51	Average A	ER DO	Min AER	pH AUR	3.8 (NH4-N Slop	pe)	2.35 SAUR	NPF	O.35	N Slope)		2 SNPF	₹
[days	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19/L] 216 09 03 03 83 81 75 78 64 61 64 552 51 65	[mg/	ER DO	Min AER	pH AUR	3.8 (NH4-N Slop [mg/L/hr]	pe)	2.35 SAUR n/g MLVSS.	NPF	O.35	N Slope)		2 SNPF g/g MLV	R (SS/hr]
	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 54 52 51		ER DO	Min AER	pH AUR	3.8 (NH4-N Slop	pe)	2.35 SAUR	NPF	O.35	N Slope)		2 SNPF	R (SS/hr)
[days	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19/L] 216 09 03 03 83 81 75 78 64 61 64 54 55 51 65	[mg/	ER DO	Min AER	pH AUR	3.8 (NH4-N Slop [mg/L/hr]	pe)	SAUR g/g MLVSS 0.40	NPF	O.35 (NOx- [mg/L/ 0.4:	N Slope)	[mg	2 SNPF g/g MLV	₹ (SS/hr]
[days	[m 2 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19/L] 216 09 03 03 83 81 75 78 64 61 64 54 55 51 65	[mg/	ER DO	Min AER	pH AUR	3.8 (NH4-N Slop [mg/L/hr] 0.61	pe)	SAUR 3/g MLVSS 0.40 Clar	NPF	O.35 (NOx- [mg/L/ 0.4: 4-P Ma	N Slope)	[m ₉	2 SNPP g/g MLV 0.28	₹ (SS/hr] 1 1 1 1 1 1 1 1 1 1 1 1 1
[days 5.25 Highest / Conce	2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19/L] 216 09 03 03 83 81 75 78 64 61 64 54 55 51 65 robic MLSS [mg/L] 1940 PUR (PO ₄ -P	[mg/ 1.10 Uptake Slop	ER DO	Min AER [unitles 6.94	pH AUR s] Clarifier ID	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla	pe) [mg	SAUR 3/g MLVSS 0.40 Clar	NPF	O.35 (NOx- [mg/L/ 0.4: 4-P Ma	N Slope) min] 2 ax ANA PC Release	[m ₉	SNPF g/g MLV 0.28 Avg. ANA Concen	RSS/hr
[days 5.25 Highest / Conce	2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	19/L] 216 09 03 03 83 81 75 78 64 61 64 54 52 51 65 robic MLSS [mg/L] 1940 PUR (PO ₄ -P	[mg/	ER DO	Min AER [unitles 6.94	pH AUR s] Clarifier ID	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla	pe) [mg	SAUR 2/g MLVSS 0.40 Clar NOx-N C	NPF	O.35 (NOx- [mg/L/ 0.4: 4-P Ma	N Slope) min] 2 ax ANA PO	[m ₉	2 SNPF g/g MLV 0.28 Avg. ANA	R SSS/hr A NO ₃ -N tration /L]
[days 5.25 Highest / Conce	SRT Aei	19/L] 216 09 03 03 83 81 75 78 64 61 64 54 52 51 65 robic MLSS [mg/L] 1940 PUR (PO ₄ -P	[mg/ 1.10 Uptake Slop g/L/hr] 0.98	ER DO	Min AER [unitles 6.94 SPUR MLVSS/hr] 7.16	pH AUR s] Clarifier ID [mg 24.90	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Clar (L]	pe) [mc	SAUR g/g MLVSS. 0.40 Clar	NPF /hr] ifier IDC PC oncentratio [mg/L] 0.2	O.35 R (NOx- [mg/L/i 0.4: 4-P Man	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70	[mg	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33	R (SS/hr) A NO ₃ -N ttration /L]
[days 5.25 Highest / Conce [m	SRT Aer	robic MLSS [mg/L] 216 09 03 03 83 81 75 78 64 61 64 55 51 65 robic MLSS [mg/L] 1940 PUR (PO ₄ -P	[mg/ 1.11 Uptake Slop g/L/hr] 0.98	ER DO L.]	Min AER [unitles 6.94 SPUR MLVSS/hr 7.16 P AA1	pH AUR s] Clarifier ID I [mg 24.90	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Clar /L] 000	pe) [mc] arifier IDC 1 [mg/L] 2.9200 PO ₄ -P	SAUR g/g MLVSS. 0.40 Clar NOx-N Clar	NPF /hr] ifier IDC PC oncentratio [mg/L] 0.2 ifier PO ₄ -	O.35 R (NOx- [mg/L/i 0.4: -P Man	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P	[m ₉	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33	R (SS/hr) A NO ₃ -N Attration (/L) 900
[days 5.25 Highest / Conce [m 1	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 55 51 65 mobic MLSS [mg/L] 1940 PUR (PO ₄ -P [m] 1	[mg/ 1.11 Uptake Slop g/L/hr] 0.98	ER DO L] 6 [mg/g AA1 PO ₄ -lelease Slo	Min AER [unitles 6.94 SPUR MLVSS/hr 7.16 P AA1	pH AUR s] Clarifier ID [mg 24.9) Specific PC Release Rat	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Clar 'L] 000 D ₄ - AA2 e Releas	pe) [mg/L] 2.9200 PO ₄ -P se Slope	SAUR g/g MLVSS, 0.40 Clar NOx-N C AA2 Spec	NPF /hr] ifier IDC PC oncentratio [mg/L] 0.2 ifier PO ₄ - se Rate	O.35 R (NOx- [mg/L/ 0.4: -P Man AA3 Relea	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P sse Slope	[mg	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33	R (SS/hr) A NO ₃ -N tration //L] g0 Gic PO ₄ -
[days 5.25 Highest / Conce [m 1	Page Page	robic MLSS [mg/L] 1940 AA6 NO) [mg/L] AA6 NO)	[mg/ 1.10 Uptake Slop g/L/hr] 0.98	ER DO L] S [mg/g AA1 PO ₄ -lelease Slo	Min AER [unitles 6.94 SPUR MLVSS/hr 7.16 P AA1	pH AUR s] Clarifier ID [mg 24.90 I Specific PC Release Rat g/g MLVSS/l	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Clar /L] 000 D ₄ - AA2 e Releas hr] [mg/L/hr]	pe) [mg/L] 2.9200 PO ₄ -P se Slope y/L/hr]	SAUR g/g MLVSS, 0.40 Clar NOx-N C AA2 Spec P Relea: [mg/g ML	NPF /hr] ifier IDC PC oncentratio [mg/L] 0.2 ifier PO ₄ - se Rate _VSS/hr]	O.35 R (NOx- [mg/L/i 0.4: -P Man AA3 Relea	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P sse Slope g/L/hr]	[mg	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33 3 Specif Release	R (SS/hr) A NO ₃ -N tration //L] 90 tic PO ₄ te Rate (SS/hr)
[days 5.25 Highest / Conce [m 1	[m 2 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1 1	9/L] 216 09 03 03 83 81 75 78 64 61 64 55 51 65 mobic MLSS [mg/L] 1940 PUR (PO ₄ -P [m] 1	[mg/ 1.10 Uptake Slop g/L/hr] 0.98	ER DO L] 6 [mg/g AA1 PO ₄ -lelease Slo	Min AER [unitles 6.94 SPUR MLVSS/hr 7.16 P AA1	pH AUR s] Clarifier ID [mg 24.9) Specific PC Release Rat	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Clar /L] 000 D ₄ - AA2 e Releas hr] [mg/L/hr]	pe) [mg/L] 2.9200 PO ₄ -P se Slope	SAUR g/g MLVSS, 0.40 Clar NOx-N C AA2 Spec	NPF /hr] ifier IDC PC oncentratio [mg/L] 0.2 ifier PO ₄ - se Rate _VSS/hr]	O.35 R (NOx- [mg/L/i 0.4: -P Man AA3 Relea	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P sse Slope	[mg	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33	R (SS/hr) A NO ₃ -N tration //L] g0 Gic PO ₄ - B Rate
[days 5.25 Highest / Conce [m 1	Property Property	PUR (PO ₄ -P AA6 NO) [mg/L] AA6 NO) 1940 AA6 NO) 1940 AA6 NO)	[mg/ 1.10 Uptake Slop g/L/hr] 0.98	ER DO L] 6 [mg/g AA1 PO ₄ -lelease Sic [mg/L/hr] 5.04	Min AER [unitles 6.94] SPUR MLVSS/hrj 7.16 P AA1	pH AUR s] Clarifier ID I mg 24.9(I Specific PC Release Rat g/g MLVSS/I 6.47	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla 'L] 000 D ₄ - AA2 e Releas nr] [mg	pe) [mç arifier IDC 1 [mg/L] 2.9200 PO ₄ -P se Slope y/L/hr] 1.28	SAUR g/g MLVSS. 0.40 Clar NOx-N C AA2 Spec P Relea: [mg/g ML	NPF Interpolation NPF Inte	O.35 R (NOx- [mg/L/i 0.4 -P Man AA3 Relea	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P se Slope g/L/hr] 7.28	[m ₅]	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33 3 Specif Release	R (SS/hr) A NO ₃ -N tration //L] g0 Gic PO ₄ - B Rate
[days 5.25 Highest / Conce m 1 1 AA4 [m 0 AA4 F	Part Part	PUR (PO ₄ -P AA6 NO) [mg/L] AA6 NO) A4 Specific PC	[mg/ 1.10 Uptake Slop g/L/hr] 0.98 c-N Re	ER DO L] 3 e) [mg/g AA1 PO ₄ -lelease Slc [mg/L/hr] 5.04 O ₄ -P	Min AER [unitles 6.94] SPUR MLVSS/hrj 7.16 P AA1	pH AUR s] Clarifier ID [mg 24.9i I Specific PC Release Rat 3/g MLVSS/i 6.47 c PO ₄ - A	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla [L] 000 D ₄ - AA2 Releas nr] [mg/L/mr] 11 A6 PO ₄ -P	pe) [mg/L] 2.9200 PO ₄ -P se Slope y/L/hr] 1.28 AA6 Spec	SAUR 3/g MLVSS 0.40 Clar CO AA2 Spec P Relea: [mg/g ML 14.	NPF Interpolation of the property of the prop	O.35 R (NOx- [mg/L/i 0.4: -P Man AA3 Relea [mg/ciake]	N Slope) min] 2 Release [mg/L] 21.70 8 PO ₄ -P sse Slope g/L/hr] 7.28 AA1) Spec	[m ₅]	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33 3 Specifi Release ng/g MLV 9.35	R SS/hr A NO ₃ -N A NO ₃ -N A NO ₃ -N A NO ₃ -N A NO ₃ -N B Rate B Rate C/SS/hr]
[days 5.25] Highest / Conce [m 1] AA4 [m 0] AA4 FRelease	Page Page	PUR (PO ₄ -P AA6 NO) A4 Specific PC P Release Rate	[mg/ 1.10 Uptake Slop g/L/hr] 0.98 c-N Ro	ER DO L] S (mg/g) AA1 PO ₄ -P Slease Slo (mg/L/hr) 5.04	Min AER [unitles 6.94 SPUR MLVSS/hr 7.16 P AA1 ppe P [mg	pH AUR s] Clarifier ID [mg. 24.9i I Specific PC Release Rat g/g MLVSS/i 6.47 c PO ₄ - A Rate Rei	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla (L) 1000 04- AA2 e Releas nr) [mg, 11 A6 PO ₄ -P ease Slope	pe) [mg/L] 2.9200 PO ₄ -P se Slope y/L/hr] 1.28 AA6 Spec P Releas	SAUR g/g MLVSS. 0.40 Clar NOx-N Clar P Relea: [mg/g ML 14. ciffic PO ₄ - (AA	NPF Iffier IDC PC oncentratio [mg/L] 0.2 Iffice PO ₄ - se Rate VSS/hr] 49 Iffice PO ₄ - se Rate VSS/hr] Slope	O.35 R (NOx- [mg/L/i 0.4- P Main AA3 Relea [mi cake (/co	N Slope) min] 2 ax ANA PC Release [mg/L] 21.70 8 PO ₄ -P use Slope g/L/hr] 7.28 AA1) Spec	[ms] AA−P AA−P Incomplete the second of t	SNPP g/g MLV 0.28 Avg. ANA Concen [mg 0.33 3 Specifi Release 19/g MLV 9.35	R (SS/hr) A NO ₃ -N ttration /L] 90 cic PO ₄ - P Rate /SS/hr] 6
[days 5.25 Highest / Conce m 1 1 AA4 [m 0 AA4 F	PO P A P P P P P P P P	PUR (PO ₄ -P AA6 NO) [mg/L] AA6 NO) A4 Specific PC	[mg/ 1.10 Uptake Slop g/L/hr] 0.98 c-N Ro	ER DO L] S (mg/g) AA1 PO ₄ -P elease Sic (mg/L/hr) 5.04 O ₄ -P e Siope /hr]	Min AER [unitles 6.94] SPUR MLVSS/hrj 7.16 P AA1	pH AUR s] Clarifier ID [mg 24.9i I Specific PC Release Rat g/g MLVSS/i 6.47 c PO ₄ - A Rate Rei	(NH4-N Slop [mg/L/hr] 0.61 C NH ₄ -N Cla [L] 000 D ₄ - AA2 Releas nr] [mg/L/mr] 11 A6 PO ₄ -P	pe) [mg/L] 2.9200 PO ₄ -P se Slope y/L/hr] 1.28 AA6 Spec	SAUR g/g MLVSS. 0.40 Clar NOx-N AA2 Spec P Relea: [mg/g ML 14. cific PO ₄ - se Rate VSS/hr]	NPF Interpolation of the property of the prop	O.35 R (NOx- [mg/L/i 0.4- P Main AA3 Relea [mi cake (/co	N Slope) min] 2 Release [mg/L] 21.70 8 PO ₄ -P sse Slope g/L/hr] 7.28 AA1) Spec	[ms] AA−P AA−P Incomplete the second of t	SNPF g/g MLV 0.28 Avg. ANA Concen [mg 0.33 3 Specifi Release ng/g MLV 9.35	RS/hr A NO ₃ -hr tration /L] 90 ic PO ₄ Rate //SS/hr ic Harter //SS/hr ic mg]

# Chem Us	ed	FeCl3 Use	_	FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]		[M	GD]	[MGD]	[MGD]
0		0.00	0	0.000)	18.75		17	.13	,	11.19	0
ANA MLTSS	ANA	4 % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	IXNA	MLVSS	AER ML	TSS	AER % VC	L AER MLVSS
[mg/L]		[%]	[1	mg/L]	[mg/L]	[%]	[m	ng/L]	[mg/l	L]	[%]	[mg/L]
865		90		779	1940	79	1:	533	1940)	79	1533



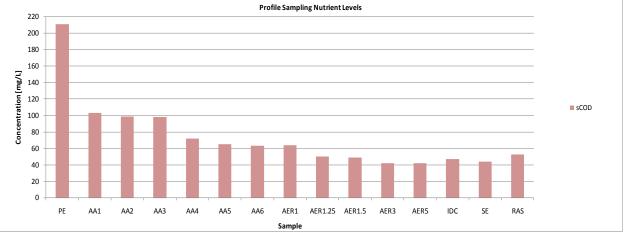


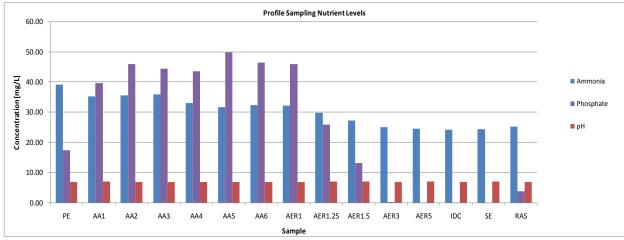


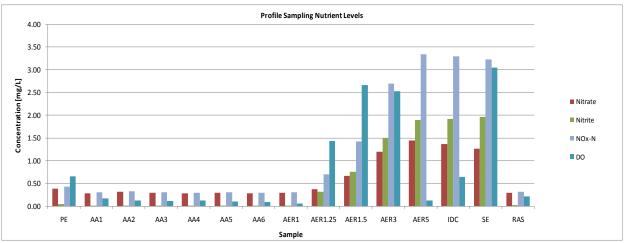
Week 15 – 10/26/09 & 10/27/09

		vation Tank (. , , ,											
KUFILE 1	ı rain 4, Ae	eration Tank 4,	Ciarifier 4												
		HRT along BNR													
		Process (w/out								NOx	-N by Nitrate H	ACH Phospha	ate by HAO	Н	
Date	Sample Time	Recycles)	Sample Number	r Reactor 1	ype Ammoni	a by HACH TNT	Nitrate by	HACH TNT	Nitrite by HACH	1 TNT	+ Nitrite HACH		TNT	DO	рН
dd-mmm-yy	h:mm	hr			mç	g/L NH3-N	mg/L	NO3-N	mg/L NO2-	N	mg/L NOx-N	mg/l	L PO4-P	[mg/L]	[unitless]
26-Oct-09	7:11	0	PE	-		39.10	0.	39	0.04		0.43	1	17.46	0.66	6.90
26-Oct-09	7:27	0.89	AA1	Anaerol	ic	35.20	0.	29	0.01		0.30	3	39.65	0.17	6.99
26-Oct-09	7:57	1.78	AA2	Anaerol		35.60		32	0.01		0.33		16.01	0.13	6.92
26-Oct-09	8:27	2.67	AA3	Anaerol		36.00		29	0.01		0.30		14.38	0.11	6.98
26-Oct-09 26-Oct-09	8:57 9:10	3.55	AA4	Anaerol		33.00		29	0.01		0.30		13.56 19.93	0.13	6.98
26-Oct-09	9:10	4.44 5.33	AA5 AA6	Anaerol Anaerol		31.60 32.40		29 29	0.01 0.01		0.31		16.50	0.10	6.97 6.96
26-Oct-09	9:38	0.00	AER1	Aerobi		32.20		30	0.01		0.31		16.01	0.03	6.93
26-Oct-09	9:46	0.34	AER1.25	Aerobi		29.80		38	0.32		0.70		25.94	1.43	7.02
26-Oct-09	9:54	0.69	AER1.5	Aerobi	;	27.20	0.	66	0.76		1.42	1	13.22	2.66	7.03
26-Oct-09	10:39	3.44	AER3	Aerobi	;	25.10	1.	20	1.50		2.70		0.31	2.53	6.94
26-Oct-09	11:47	6.88	AER5	Aerobi	;	24.50		45	1.89		3.34		0.08	0.13	7.01
26-Oct-09	11:55		IDC	-		24.20	1.	37	1.92		3.29		0.14	0.64	6.89
26-Oct-09	14:55		SE	-		24.30	1.	26	1.96		3.22		0.17	3.04	7.12
26-Oct-09	15:19		RAS	-		25.20	0.	29	0.03		0.32		3.79	0.22	6.98
QA/	QC S	ample:		E											
	EPA	365.1		EPA	350.1	EPA	353.2	2 E	EPA 353	3.2	EPA 3	353.2	EF	A 35	3.2
5	COD	by CE	_	O-I	PO_4	NE	I 4-N		NO _{2,3} -I	7	NO:	3-N	1	VO 2-1	N
	[m	ıg/L]		mg	₅ /L	m	g/L		mg/L		mg	/L		mg/L	,
		211	_												
		03 99	_												
		98													
		72													
		65													
		63 64													
		50													
		49													
		42 42	_												
		42 47													
		-, 44		0.1	01	2	4.7	Т	2.64		0.6	53		2.01	
		53		0.1	0.1		,		2.0.		0.0	,,,		2.01	
													Т		
			Λυργασο Λ	ED DO	Min AED	pH AUR	(NILIA NI	Slope)	SAU	D	NIDD (NIC	0x-N Slope		SNPF	,
Actual		robic MLSS					•				,		<i>'</i>		
[days		[mg/L]	[mg/		[unitles	s]	[mg/L/h	r]	[mg/g ML\			/L/min]	[m	g/g MLV	
5.90)	2320	1.30	<u> </u>	6.93	<u> </u>	0.92		0.5		().64	—	0.36	
_	AER NO ₂ -N									Clarifier II		Max ANA P		Avg. ANA	
Conce	entration	` '	Uptake Slop	e)	SPUR	Clarifier ID	C NH ₄ -N	Clarifier	IDC NOx-N	Conce	ntration	Release	Э	Concent	ration
	ng/L]		g/L/hr]	[mg/g	MLVSS/hr				mg/L]		g/L]	[mg/L]		[mg/	
1	1.89	1	0.30		5.76	24.2	000	3.	.2900	0	.1	32.47		0.29	94
				AA1 PO ₄		1 Specific P		AA2 PO ₄ -		Specific F		A3 PO ₄ -P		3 Specifi	
AA4	NOx-N	AA6 NO	(-N R	elease SI		Release Ra		lease Sl	· .	elease Ra		lease Slope		Release	Doto
				[mg/L/hr] [m	g/g MLVSS/	hr]	[mg/L/hr] [mg/	g MLVSS	S/hr]	[mg/L/hr]	ſm		Rate
	ng/L]	[mg/L]					- 1			7 CE		7 44	Į.··	g/g MLV	SS/hr]
	ng/L] 0.30	[mg/L] 0.30		9.45		5.20		13.91		7.65		7.41		g/g MLV 4.07	SS/hr]
0	0.30	0.30		9.45				13.91		7.05		7.41			SS/hr]
AA4 F	0.30 PO ₄ -P A	0.30		9.45 PO ₄ -P	AA5 Specifi	c PO ₄ -	A6 PO ₄ -P	13.91 AA6	Specific PO ₄ -	(AA1) C0	OD Uptake	(AA1) Spe	cific		SS/hr]
AA4 F	PO ₄ -P A	0.30 AA4 Specific PC P Release Rate	e Release	9.45 PO ₄ -P e Slope	P Release	c PO ₄ - A	lease Slop	13.91 AA6 e P R	Release Rate	(AA1) CC	lope	(AA1) Spe COD Uptake	cific Rate		SS/hr]
AA4 F	PO ₄ -P A e Slope	0.30	e Release	9.45 PO ₄ -P e Slope _/hr]		c PO ₄ - A Rate Re		13.91 AA6 e P R		(AA1) CC SI [mg	lope	(AA1) Spe	cific Rate	4.07	SS/hr] _{-P/COD}

# Chem Use	ed	FeCl3 Used [hr]	b	FeCl3 Additio	on I	Biosolid Influ Flow [MGD]	uent	[M	Y Flow GD]]	CY Flow MGD]	NRCY Flow [MGD]
0		0.00	0	0.000)	18.80		17	.01	1	1.129	0
ANA MLTSS	ANA	\ % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	IXNA	MLVSS	AER ML	_TSS	AER % VC	L AER MLVSS
[mg/L]		[%]	[۱	mg/L]	[mg/L]	[%]	[m	ng/L]	[mg/	L]	[%]	[mg/L]
2140		85		1819	2320	77	11	786	2320	0	77	1786

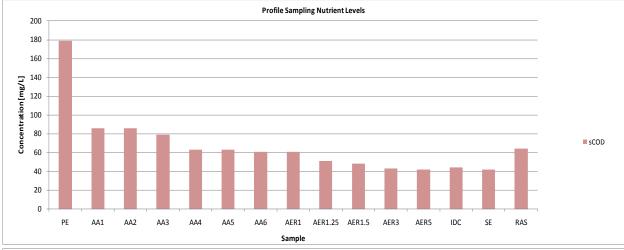


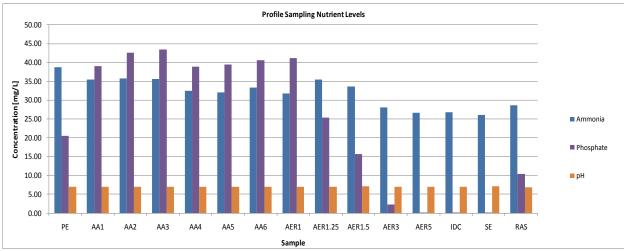


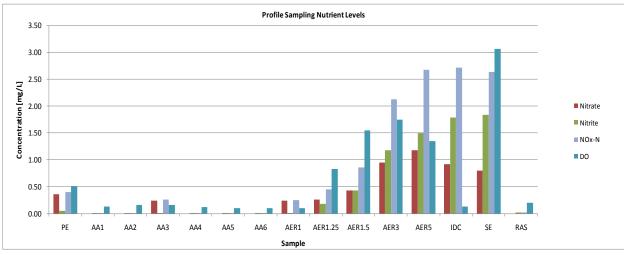


ROFILE 2	Train 5, Ae	eration Tank 5,	Clarifier 5												
		HRT along BNR													
Date	Sample Time	Process (w/out Recycles)	Sample Number	Reactor Ty	pe Ammonia	by HACH TNT	Nitrate by HA0	CH TNT	Nitrite by HACH TN		y Nitrate HA0 itrite HACH		ate by HAC NT844	H DO	pН
dd-mmm-yy	h:mm	hr		,		L NH3-N	mg/L NO3		mg/L NO2-N		/L NOx-N		L PO4-P	[mg/L]	[unitless]
27-Oct-09	7:09	0	PE	-		38.80	0.36	/ 18	0.05	,	0.40		20.56	0.51	7.01
27-Oct-09	7:26	0.88	AA1	Anaerobio		35.40	0.00		0.01		0.01		38.99	0.13	7.05
27-Oct-09	7:56	1.78	AA2	Anaerobio	:	35.70	0.00		0.01		0.01	4	12.58	0.16	7.05
27-Oct-09	8:26	2.67	AA3	Anaerobio	; ;	35.60	0.24		0.01		0.26	4	13.40	0.16	7.02
27-Oct-09	8:56	3.55	AA4	Anaerobio		32.50	0.00		0.01		0.01	_	38.83	0.12	7.01
27-Oct-09	9:03	4.44	AA5	Anaerobio		32.00	0.00		0.01		0.01		39.48	0.10	6.95
27-Oct-09	9:15	5.33	AA6	Anaerobio		33.30	0.00		0.01		0.01		10.63	0.10	6.96
27-Oct-09 27-Oct-09	9:37 9:46	0.00 0.34	AER1 AER1.25	Aerobic Aerobic		31.80 35.50	0.24 0.26		0.01		0.25		11.12 25.29	0.10	6.97 7.01
27-Oct-09	9:55	0.69	AER1.5	Aerobic		33.60	0.43		0.43		0.45		15.66	1.55	7.07
27-Oct-09	10:40	3.44	AER3	Aerobic		28.10	0.94		1.18		2.12	_	2.27	1.75	7.02
27-Oct-09	11:45	6.88	AER5	Aerobic		26.70	1.18		1.50		2.68		0.08	1.35	6.95
27-Oct-09	11:53		IDC	-		26.80	0.92		1.79		2.71		0.15	0.13	6.92
27-Oct-09	14:53		SE	-		26.00	0.80		1.84		2.64		0.16	3.06	7.10
27-Oct-09	15:13		RAS	_		28.70	0.00		0.02		0.02		10.47	0.20	6.89
		1 mg/L NH ₄ -N	1010			20.70	0.00		0.02		0.02		10.11	0.20	0.03
		0.23 mg/L NO ₃ -N													
		ample:	S	E											
		365.1		EPA 3	350.1	EPA	353.2	E	PA 353.	2 E	PA 3	53.2	EP	A 35	3.2
5	COD	by CE	L	О-Р	\mathbf{O}_4	NH	I4-N	1	1O _{2,3} -N		NO ₃	-N	r	اO ₂ -ا	Z
		ig/L]		mg.	/L	m	g/L		mg/L		mg/	L		mg/L	,
		79													
		86													
		86													
		79 63													
		63													
		61													
		61													
		51 48	_												
		48 43													
		42													
		44													
		42				20	5.6		2.1		0.23	3		1.87	
		64													
								,	0.4410		IDD (1)0		,	01100	_
Actual	SRT Ae	robic MLSS	Average Al	R DO	Min AER	pH AUR	(NH4-N SI	ope)	SAUR	N	IPR (NO)	c-N Slope	e)	SNPF	₹
[days]	[mg/L]	[mg/L	.]	[unitless	s]	[mg/L/hr]		[mg/g MLVS	S/hr]	[mg/L	_/min]	[mo	/g MLV	SS/hr]
5.20		2140	1.12	Í	6.95		1.11		0.67		0.	53		0.32	
· · · · · · · · · · · · · · · · · · ·															
Highest A	AER NO ₂ -N					1			Cl	arifier IDC	PO ₄ -P	Max ANA P	O ₄ -P	Avg. ANA	NO ₃ -N
Conce	entration	DLID (DO . D	Uptake Slope	۹ (SPUR	Clarifier ID	C NI⊒. NI C	larifier IF	OC NOx-N	Concentr	ation	Release	۵	Concen	tration
		FUN (FU4-P	opiane Siope	,		Clarifier ID	·	idinioi IL	JO HOA IV			11010031			
[m	ıg/L]	[m	g/L/hr]	[mg/g	MLVSS/hr]	[mg	/L]	[m	g/L]	[mg/L		[mg/L]		[mg/	<u>[L]</u>
1	.50	(9.02		5.47	26.80	000	2.7	'110	0.2		22.84		0.04	41
1				- I											
			А	A1 PO ₄ -F	P AA1	Specific PO	O ₄ - AA2	2 PO ₄ -P	AA2 Sp	ecific PO	4- AA	\3 PO₄-P	AA	3 Specif	ic PO ₄ -
AA4	NOx-N	AA6 NO		ease Slo		Release Rat		ase Slop		ase Rate		ase Slope		Release	
									'						
	g/L]	[mg/L]		[mg/L/hr]	Įmg	/g MLVSS/I		g/L/hr]		/LVSS/h	rj [r	ng/L/hr]	Įm	g/g MLV	
0	.01	0.01		10.31		11.03		11.99	1	2.82		7.58		8.11	
	, l	A 4 0		, , I,	۰.۸۲ ۵	DO .	4 C D C D):f- DO				.		
	UP 1	A4 Specific PC	O ₄ - AA5 P	J ₄ -P A	AA5 Specific	PU ₄ - A	A6 PO₄-P	I AA6 S	Specific PO ₄ - ()	A1) COD		(AA1) Spe			
AA4 P													_		
AA4 P Release		P Release Rate		Slope	P Release	Rate Rel	ease Slope		elease Rate	Slop	e C	OD Uptake	_	AA1) PO	₄ -P/COD
Release	Slope		e Release				ease Slope	P Re	elease Rate				Rate (-	
	Slope /hr]	P Release Rate mg/g MLVSS/r 0.58	e Release	/hr]	P Release [mg/g MLVS 0.09			P Re		Slop [mg/L/ 43.1	hr] [ı	OD Uptake mg/g MLVS 46.14	Rate (AA1) PO, [mg/i 0.2	mg]

# Chem Us		FeCl3 l Used	_	FeCl3		Biosolid Influ Flow	ient	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]		[M	GD]	[MGD]	[MGD]
0		0.00	0	0.000)	18.97		16	.57	1	1.362	0
ANA MLTSS	ANA	% VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER ML	TSS	AER % VO	L AER MLVSS
[mg/L]		[%]	1]	mg/L]	[mg/L]	[%]	[m	g/L]	[mg/l	L]	[%]	[mg/L]
1100		85		935	2140	77	16	648	2140)	77	1648



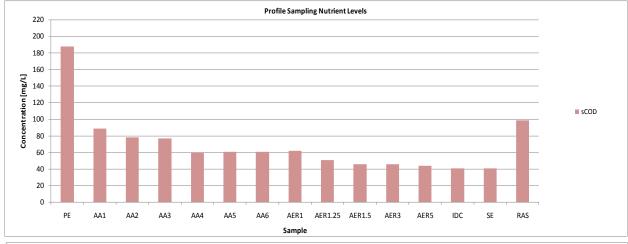


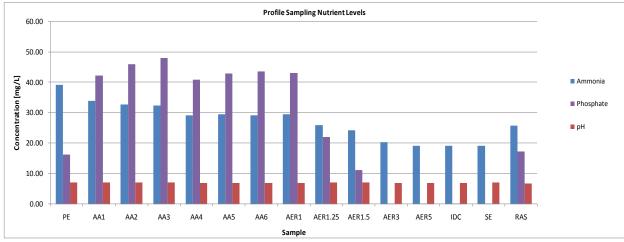


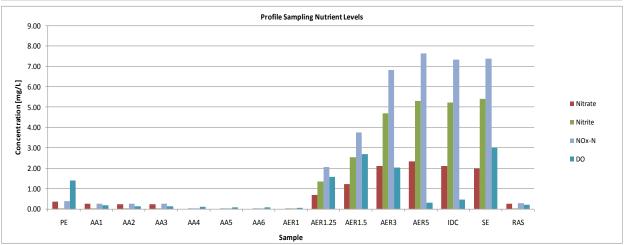
Week 16 – 11/3/09 & 11/5/09

	10 – 1														
PROFILE 1	Train 4, Ae	eration Tank 4,	Clarifier 4												
Date	Sample Time	HRT along BNR Process (w/out Recycles)	Sample Number	Reactor Typ	e Ammonia	by HACH TNT	Nitrate by HACH	HTNT Nitrite	e by HACH TNT	NOx-N by N + Nitrite	litrate HACH e HACH		ite by HAC	DO DO	pН
dd-mmm-yy	h:mm	hr			mg/	/L NH3-N	mg/L NO3-N	N n	ng/L NO2-N	mg/L N	NOx-N	mg/L	_P04-P	[mg/L]	[unitless]
3-Nov-09	7:25	0	PE	-		39.20	0.36		0.04	0.3	39	1	6.32	1.40	7.11
3-Nov-09	7:38	0.89	AA1	Anaerobic	;	33.90	0.25		0.01	0.2	27	4	2.26	0.17	7.05
3-Nov-09	8:08	1.77	AA2	Anaerobic		32.70	0.24		0.01	0.2			6.01	0.13	7.04
3-Nov-09	8:38	2.66	AA3	Anaerobic		32.40	0.24		0.01	0.2			7.97	0.12	7.02
3-Nov-09	9:08	3.55	AA4	Anaerobic		29.10	0.00		0.01	0.0			10.79	0.11	6.98
3-Nov-09 3-Nov-09	9:23 9:38	4.43 5.32	AA5 AA6	Anaerobic Anaerobic		29.40 29.20	0.00 0.00		0.01	0.0			12.91 13.56	0.08	6.97 6.97
3-Nov-09	9:53	0.00	AER1	Alaerobic		29.40	0.00		0.01	0.0			13.07	0.03	6.96
3-Nov-09	10:01	0.34	AER1.25	Aerobic		25.90	0.70		1.36	2.0			2.03	1.57	7.02
3-Nov-09	10:09	0.69	AER1.5	Aerobic		24.20	1.23		2.53	3.7	76	1	1.09	2.68	7.05
3-Nov-09	10:54	3.43	AER3	Aerobic		20.30	2.12		4.70	6.8	82	(0.12	2.03	6.86
3-Nov-09	11:54	6.87	AER5	Aerobic		19.10	2.33		5.30	7.6	63	- (0.08	0.30	6.84
3-Nov-09	12:06		IDC	-		19.10	2.10		5.22	7.3	32	- 1	0.09	0.45	6.83
3-Nov-09	15:08		SE	-		19.20	1.98		5.40	7.3	38	(0.14	3.01	7.01
3-Nov-09	15:30		RAS	-		25.70	0.26		0.03	0.2	29	1	7.29	0.21	6.81
NOTE: Thes	se values are <	1 mg/L NH ₄ -N									_				
		:0.23 mg/L NO ₃ -N													
		ample:	s	E											
		365.1		EPA 3	50.1	EPA	353.2	EPA	353.2	EP	A 35	3.2	EP	A 35	3.2
		by CE	_	O-P	-		I ₄ -N) _{2,3} -N		NO ₃ -N	7		NO ₂ -1	
s				mg/	L	ms	g/L	n	ng/L		mg/L	,		mg/L	,
S	1	ng/L] 188 89 78 77													
S	1	188 89 78 77 60 61 61 62 51 46 46													
S	1	188 89 78 77 60 61 61 62 51 46 46 44		0.1	7		3.1	6	5.41		0.9			5.61	
S	1	188 89 78 77 60 61 61 62 51 46 46		0.1	7		3.1	ě	5.41		0.9			5.61	
S	1	188 89 78 77 60 61 61 62 51 46 46 44 41		0.1	7		3.1	· ·	5.41		0.9			5.61	
		188 89 78 77 60 61 61 62 51 46 46 44 41	Average Al			18		pe)	SAUR		R (NOx-N			SNPF	
Actual S	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99	[mg/l	ER DO N	Min AER [unitless	pH AUR	(NH4-N Slop [mg/L/hr]	pe)	SAUR /g MLVSS/		R (NOx-N [mg/L/n	nin]		SNPF g/g MLV	
Actual S	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99		ER DO N	Min AER	pH AUR	(NH4-N Slop	pe)	SAUR		R (NOx-N	nin]		SNPF	
Actual S [days 4.00 Highest A	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99	[mg/l 1.33	ER DO I	Min AER [unitless 6.84	pH AUR	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla	pe) [mg	SAUR y/g MLVSS, 0.64 Clar	fier IDC PC	R (NOx-N [mg/L/n 1.70 0 ₄ -P Max	nin] x ANA P Release	[mg	SNPF g/g MLV 0.85 Avg. ANA Concent	SS/hr] . NO ₃ -N
Actual S [days 4.00 Highest A Conce	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99	[mg/l 1.33 Uptake Slope g/L/hr]	ER DO I	Min AER [unitless 6.84 PUR //LVSS/hr]	pH AUR Clarifier ID [mg/	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla	[mg	SAUR y/g MLVSS, 0.64 Clar	fier IDC PC oncentratio	R (NOx-N [mg/L/n 1.70 0 ₄ -P Max	x ANA P Release	[mg	SNPF g/g MLV 0.85 Avg. ANA Concent	SS/hr] NO ₃ -N tration
Actual S [days 4.00 Highest A Conce	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99	[mg/l 1.33	ER DO I	Min AER [unitless 6.84	pH AUR	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla	pe) [mg	SAUR y/g MLVSS, 0.64 Clar	fier IDC PC	R (NOx-N [mg/L/n 1.70 0 ₄ -P Max	nin] x ANA P Release	[mg	SNPF g/g MLV 0.85 Avg. ANA Concent	SS/hr] NO ₃ -N tration
Actual S [days 4.00 Highest A Conce	SRT Ae	88 89 78 77 60 61 61 62 51 46 46 44 41 41 99	[mg/L] 1.33 Uptake Slope Uptake Slope 0.40	ER DO I	Min AER [unitless 6.84 PUR MLVSS/hr] 1.71 AA1 P F	pH AUR si] Clarifier IDI [mg/ 19.10 Specific PC Release Rate	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla [L] 000 D ₄ - AA2 e Release	[mg/L] 7.3200 PO ₄ -P se Slope	SAUR /g MLVSS/ 0.64 Clari	fifer IDC PC concentratio [mg/L] 0.1 diffic PO ₄ - se Rate	R (NOx-N [mg/L/n 1.70 1.70 Max on AA3 Release	x ANA P Release [mg/L] 31.65	[mg	SNPF a/g MLV 0.85 Avg. ANA Concent [mg/ 0.12 3 Specifi	SS/hr] NO ₃ -N tration L] 22 cc PO ₄ - Rate
Actual S [days 4.00 Highest # Conce [m 5	SRT Ae: Sign Aer NO ₂ -N AER NO ₂ -N ng/L] Sign NOx-N ng/L]	888897887760616161625146464444141999	[mg/L] 1.33 Uptake Slope Uptake Slope 0.40	ER DO	Min AER [unitless 6.84 PUR MLVSS/hr] 1.71 AA1 P F	DH AUR Clarifier IDI [mg/ 19.10 Specific PC Release Rate //g MLVSS/H	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla [L] 000 D ₄ - AA2 e Releas hr] [mg	[mg/L] 7.3200 PO ₄ -P se Slope	SAUR //g MLVSS, 0.64 Clari NOx-N C AA2 Spec P Releas	fier IDC PC oncentratio [mg/L] 0.1 cific PO ₄ - se Rate VSS/hr]	R (NOx-N [mg/L/n 1.70 1.70 AA3 Releas [mg	x ANA P Release [mg/L] 31.65 PO ₄ -P se Slope	[mg	SNPF g/g MLV 0.85 Avg. ANA Concent [mg/ 0.12 3 Specifi Release	SS/hr] NO ₃ -N tration L] 22 cc PO ₄ - Rate SS/hr]
Actual S [days 4.00 Highest # Conce [m 5	SRT Ae. SI O O AER NO ₂ -N SI O O NOx-N	888897887760616161625146464444141999robic MLSS	[mg/L] 1.33 Uptake Slope Uptake Slope 0.40	ER DO I	Min AER [unitless 6.84 PUR MLVSS/hr] 1.71 AA1 P F	pH AUR si] Clarifier IDI [mg/ 19.10 Specific PC Release Rate	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla [L] 000 D ₄ - AA2 e Releas hr] [mg	[mg/L] 7.3200 PO ₄ -P se Slope	SAUR /g MLVSS/ 0.64 Clari	fier IDC PC oncentratio [mg/L] 0.1 cific PO ₄ - se Rate VSS/hr]	R (NOx-N [mg/L/n 1.70 1.70 AA3 Releas [mg	x ANA P Release [mg/L] 31.65	[mg	SNPF a/g MLV 0.85 Avg. ANA Concent [mg/ 0.12 3 Specifi	SS/hr] NO ₃ -N tration L] 22 cc PO ₄ - Rate SS/hr]
Actual S [days 4.00 Highest A Conce [m 5 AA4 [m 0.0 AA4 P Release	SRT Ae: SSRT Ae	8888978 78 77 60 61 61 62 51 46 44 41 41 99 8 60 60 60 60 60 60 60 60 60 60 60 60 60	[mg/l 1.33 Uptake Slope g/L/hr] 0.40 A-N Re	ER DO	Viin AER [unitless 6.84] PUR VILVSS/hr] 1.71 AA1 PF [mg] A5 Specific P Release	DH AUR Clarifier IDI [mg/ 19.10 Specific PC Release Rate /g MLVSS/t 12.44 CPO4- A. Rate Rel	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla (L] 000 D ₄ - AA2 e Releas nr] [mg 13 A6 PO ₄ -P ease Slope	mg/L] 7.3200 PO ₄ -P se Slope //L/hr] AA6 Spec P Release	SAUR J/g MLVSS, 0.64 Clar Clar AA2 Spec P Releas [mg/g ML 11. ific PO4- (AAe Rate	fier IDC PC oncentratio [mg/L] 0.1 iific PO ₄ - se Rate NSS/hr] 22 11) COD Up Slope	R (NOx-N [mg/L/n 1.70 1.70 AA3 Releas [mg 8	x ANA P Release [mg/L] 31.65 PO ₄ -P se Slope t/L/hr] .79	O ₄ -P AA P [m	SNPF a/g MLV 0.85 Avg. ANA Concent [mg/ 0.12 3 Specifit Release g/g MLV 7.48	SS/hr] NO ₃ -N tration [L] 22 cc PO ₄ - Rate SS/hr]
Actual S [days: 4.00 Highest A Conce [m] 5	SRT Aes AER NO ₂ -N entration ng/L] .01 NOx-N 1g/L] .01 PO ₄ -P e Slope L/hr]	88889788776061616162514644441999 robic MLSS [mg/L] 2660 PUR (PO ₄ -P [mg/L] 0.01	[mg/l 1.33 Uptake Slope g/L/hr] 0.40 A-N Re	ER DO	Viin AER [unitless 6.84] PUR VILVSS/hr] I-71 AA1 P F [mg] A5 Specific	DH AUR Clarifier IDI [mg/ 19.10 Specific PC Release Rate /g MLVSS/t 12.44 CPO4- A. Rate Rel	(NH4-N Slop [mg/L/hr] 1.27 C NH ₄ -N Cla L] 000 D ₄ - AA2 Release nr] [mg 13	[mg/L] 7.3200 PO ₄ -P se Slope //L/hr] AA6 Spec	SAUR J/g MLVSS, 0.64 Clar AA2 Spec P Releas [mg/g ML 11. ific PO ₄ - e Rate VSS/hr]	fier IDC PC oncentratio [mg/L] 0.1 see Rate VSS/hr] 222	R (NOx-N [mg/L/n 1.70 1.70 AA3 Releas [mg 8	x ANA P Release [mg/L] 31.65 PO ₄ -P se Slope y/L/hr] .79	O ₄ -P AA P [m	SNPF g/g MLV 0.85 Avg. ANA Conceni [mg/ 0.12 3 Specifi Release g/g MLV 7.48	SS/hr] NO ₃ -N tration [L] 22 IC PO ₄ - Rate SS/hr]

# Chem Us	ed	FeCl3 Use	_	FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L]	[MGD]	,	[M	GD]]	MGD]	[MGD]
0		0.00	0	0.000)	18.84		17	.12	1	1.795	0
ANA MLTSS	ANA	A % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	IXNA	MLVSS	AER ML	TSS	AER % VC	L AER MLVSS
[mg/L]		[%]		mg/L]	[mg/L]	[%]		ng/L]	[mg/l		[%]	[mg/L]
1400		84		1176	2660	75	19	995	2660)	75	1995

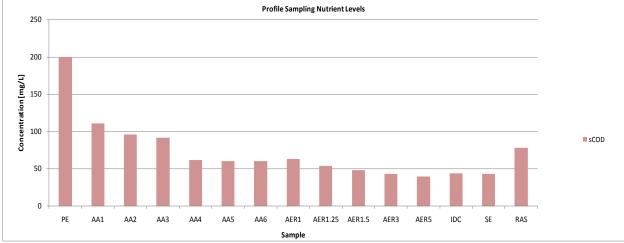


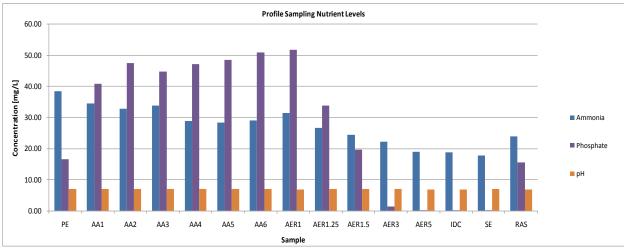


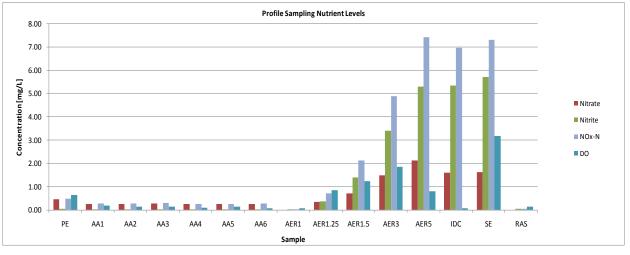


ROFILE 2	Train 5, Ae	ration Tank 5,	Claritier 5												
	,	HRT along BNR													
		Process (w/out								NOx-N by Ni	trate HACH	Phosphat	e by HACH	Н	
Date	Sample Time	Recycles)	Sample Number	Reactor Type	Ammonia	by HACH TNT	Nitrate by HAC	CH TNT Nitrit	e by HACH TNT	+ Nitrite	HACH	TN	T844	DO	pН
dd-mmm-yy	h:mm	hr			mg/l	L NH3-N	mg/L NO3	3-N I	ng/L NO2-N	mg/L N	IOx-N	mg/L	P04-P	[mg/L]	[unitless]
5-Nov-09	7:40	0	PE		3	8.50	0.45		0.04	0.4	19	16	6.64	0.65	6.98
5-Nov-09	7:54	0.88	AA1	Anaerobic	3	4.60	0.25		0.02	0.2	27	40).79	0.18	7.01
5-Nov-09	8:24	1.77	AA2	Anaerobic	3	2.80	0.26		0.01	0.2	27	47	'.48	0.13	7.00
5-Nov-09	8:54	2.66	AA3	Anaerobic	3	3.90	0.28		0.01	0.3	30	44	.71	0.14	6.99
5-Nov-09	9:24	3.55	AA4	Anaerobic		28.90	0.25		0.01	0.2			'.15	0.10	6.94
5-Nov-09	9:36	4.43	AA5	Anaerobic	_	28.40	0.25		0.01	0.2			3.62	0.14	6.94
5-Nov-09	9:48	5.32	AA6	Anaerobic	_	9.00	0.25		0.01	0.2).91	0.08	6.94
5-Nov-09	10:00	0.00	AER1	Aerobic		1.40	0.00		0.01	0.0			.72	0.07	6.93
5-Nov-09	10:08	0.34	AER1.25	Aerobic		26.70	0.35		0.36	0.7			3.86	0.84	7.02
5-Nov-09	10:16	0.69	AER1.5	Aerobic	_	4.50	0.72		1.40	2.1		-	1.74	1.24	7.05
5-Nov-09	11:01	3.43	AER3	Aerobic		2.20	1.48		3.40	4.8			.43	1.86	7.01
5-Nov-09	12:01	6.87	AER5	Aerobic		9.00	2.12		5.30	7.4			.09	0.80	6.84
5-Nov-09	12:18		IDC	-		8.80	1.60		5.35	6.9			.17	0.07	6.85
5-Nov-09	15:18		SE	-		7.80	1.61		5.70	7.3			.16	3.18	6.98
5-Nov-09	15:32		RAS		2	4.00	0.00		0.04	0.0)4	15	5.50	0.14	6.85
*NOTE: Thes	e values are <1	I mg/L NH ₄ -N													
*NOTE: Thes	e values are <0).23 mg/L NO ₃ -N													
QA/	QC Sa	ample	S	E											
	EPA	365.1	I	EPA 3:	50.1	EPA	353.2	EP/	353.2	EP	A 353	3.2	EP	A 35	3.2
	000	ь o = 1		O-PO	\mathbf{D}_4	NH	I4-N	NC	D _{2,3} -N	_	1O3-N	J	Ν	1O ₂ -I	V
١ .					-4	- 1	4 - 1	110		_	103 1	`			
s		by CEI		mg/	L	ms	z/L	n	ng/L	3	mg/L		1	ng/L	_
S	[m 2	g/L] 00		mg/	L	mg	g/L	n	ng/L	1	mg/L		r	ng/L	_
y.	[m 2 1	g/L] 00 11		mg/	L	mį	g/L	n	ng/L		mg/L		r	ng/L	<u>.</u>
S	[m 2 1	g/L] 00 11 96		mg/	L	mş	g/L	n	ng/L		mg/L		r	ng/L	_
V	[m 2 1 9	g/L] 00 11 96 92		mg/		mş	g/L	n	ng/L		mg/L		r	ng/L	
	[m 2 1 9	g/L] 00 11 96 92 62		mg/		mg	g/L	n	ng/L		mg/L			ng/L	
	[m 2 1 9	g/L] 00 11 96 92		mg/	<u> </u>	mg	g/L	n	ng/L	,	mg/L		Т	ng/L	
8	[m 2 1 9 9 9	g/L] 00 11 96 92 62 60 60 60		mg/	L	mg	g/L	n	ng/L		mg/L		Т	ng/L	_
9	[m 2 1 1 9 9 9	g/L] 00 11 96 92 62 60 60 63 64		mg/	L	mg	z/L	n	ng/L		mg/L		Г	mg/L	
S	[m 2 1 3 9 9 9	g/L] 00 11 96 92 62 60 60 60		mg/	<u> </u>	mg	g/L	n	ng/L	,	mg/L		r	mg/L	
S	[m 2 1 9 9 9 9 9 9	g/L] 00 11 96 92 62 60 60 63 54 48 43		mg/		mg	<u>z/L</u>	n	ng/L	,	mg/L				
S	[m 2 1 9 9 9 9 9 2 2	g/L] 00 11 96 92 62 60 63 54 48 43 40		mg/	L		g/L 5.9		5.92		mg/L			ng/I 5.81	
S	[m 2 1 9 9 9 9 9 2 2	g/L] 00 11 96 92 62 60 60 63 54 48 43		mg/	L										
S	[m 2 1 9 9 9 9 9 2 2	g/L] 00 11 96 92 62 60 63 54 48 43 40		mg/	L										
	[m 2 1 1 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	g/L] 00 111 96 92 62 60 63 54 48 43 40 44 43 78				16	5.9		5.92		<0.20			5.81	
Actual	[m 2 1 1 5 5 5 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 11 96 92 52 50 60 63 54 48 43 40 44 43 78	Average AE	ER DO N	flin AER	oH AUR	5.9 (NH4-N SIG	ope)	5.92 SAUR	NPF	<0.20	N Slope)		5.81 SNPF	₹
Actual (days	[m 2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 11 96 92 52 50 60 63 54 48 43 40 44 43 78	Average AE	ER DO N	flin AER	oH AUR	5.9 (NH4-N Slo [mg/L/hr]	ope)	5.92 SAUR g/g MLVSS,	NPF	<0.20 ₹ (NOx-N [mg/L/n	N Slope) nin]		5.81 SNPF /g MLV	R SS/hr]
Actual	[m 2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 11 96 92 52 50 60 63 54 48 43 40 44 43 78	Average AE	ER DO N	flin AER	oH AUR	5.9 (NH4-N SIG	ope)	5.92 SAUR	NPF	<0.20	N Slope) nin]		5.81 SNPF	R SS/hr]
Actual (days) 4.60	[m 2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 11 96 92 52 50 60 63 54 48 43 40 44 43 78	Average AE	ER DO N	flin AER	oH AUR	5.9 (NH4-N Slo [mg/L/hr]	ope)	5.92 SAUR g/g MLVSS, 0.82	NPF	<0.20 ₹ (NOx-N [mg/L/n 1.33	N Slope) nin]	[mg/	5.81 SNPF /g MLV	R SS/hr]
Actual (days 4.60)	[m 2 1 1 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5 5	9/L] OO 11 196 92 52 52 50 50 50 63 54 48 43 40 44 43 78 obic MLSS [mg/L] 2320	Average AE [mg/L 0.96	FR DO N	flin AER	pH AUR	5.9 (NH4-N Slo [mg/L/hr] 1.41	ope)	5.92 SAUR g/g MLVSS, 0.82 Clar	NPF	<0.20 R (NOx-N [mg/L/n 1.33 0 ₄ -P Max	N Slope) nin]	[mg/	5.81 SNPF /g MLV 0.77	SS/hr]
Actual: [days 4.60 Highest /	ERT Aer	9/L] OO 11 96 92 52 52 50 60 60 63 44 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96	FR DO N	fin AER [unitless 6.84	DH AUR	5.9 (NH4-N Slo [mg/L/hr] 1.41 C NH ₄ -N C	ope) [mg	5.92 SAUR g/g MLVSS, 0.82 Clar	NPF	<0.20 R (NOx-N [mg/L/n 1.33 0 ₄ -P Max	N Slope) nin] k x ANA PO Release	[mg/	SNPF /g MLV 0.77 wg. ANA Concent	SS/hr] NO ₃ -N
Actual: [days 4.60 Highest / Conce	ERT Aer	9/L] OO 11 96 92 52 52 50 60 60 63 44 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96 Uptake Slope	FR DO N	fin AER [unitless 6.84 PUR LVSS/hr]	DH AUR Clarifier ID	5.9 (NH4-N Sid [mg/L/hr] 1.41 C NH ₄ -N C	ope) [mg	5.92 SAUR 9/g MLVSS, 0.82 Clar	NPF fier IDC PC oncentratio [mg/L]	<0.20 R (NOx-N [mg/L/n 1.33 0 ₄ -P Max	N Slope) nin] x ANA PO Release [mg/L]	[mg/	SNPF /g MLV 0.77 Avg. ANA Concent [mg/	R SS/hr] NO ₃ -N
Actual: [days 4.60 Highest / Conce	ERT Aer	9/L] OO 11 196 96 92 62 63 64 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96	FR DO N	fin AER [unitless 6.84	DH AUR	5.9 (NH4-N Sid [mg/L/hr] 1.41 C NH ₄ -N C	ope) [mg	5.92 SAUR 9/g MLVSS, 0.82 Clar	NPF	<0.20 R (NOx-N [mg/L/n 1.33 0 ₄ -P Max	N Slope) nin] k x ANA PO Release	[mg/	SNPF /g MLV 0.77 wg. ANA Concent	R SS/hr] NO ₃ -N tration
Actual: [days 4.60 Highest / Conce	ERT Aer	9/L] OO 11 196 96 92 62 63 64 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03	FR DO N	flin AER [unitless 6.84 PUR ILVSS/hr]	DH AUR Clarifier ID	(NH4-N Slo [mg/L/hr] 1.41 C NH ₄ -N C L]	ope) [mg	5.92 SAUR 9/g MLVSS, 0.82 Clar	NPF /hr] fier IDC PC oncentratio [mg/L] 0.2	<0.20 R (NOx-N [mg/L/n 1.33	N Slope) nin] x ANA PO Release [mg/L]	[mg/	SNPF /g MLV 0.77 Avg. ANA Concent [mg/	SSS/hr] NO ₃ -N tration VL]
Actual (days 4.60) Highest A Conce	M 2 1 1 1 1 1 1 1 1 1	9/L] OO 11 96 92 62 62 60 60 63 54 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P [m] 1	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A	SR DO N	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1	DH AUR Clarifier ID [mg/ 18.86	(NH4-N Slo [mg/L/hr] 1.41 C NH ₄ -N C L] 000 D ₄ - AA2	ope) [mş/L] 6.9500	SAUR 3/g MLVSS, 0.82 Clar	NPF fier IDC PC oncentratio [mg/L] 0.2 iific PO ₄ -	CO.20 R (NOx-N [mg/L/n 1.33 1.4-P Max AA3	N Slope) nin] x ANA PC Release [mg/L] 34.26	[mg/	SNPF /g MLV 0.77 vg. ANA Concent [mg/ 0.25	R SS/hr] NO ₃ -N NO ₃ -N tration L] 57
Actual: [days 4.60 Highest / Conce	2 1 1 5 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 111 96 92 52 52 50 60 60 63 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel	SF DO N I I Img/g N 7 A1 PO ₄ -P ease Slop	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1 P R	Clarifier ID [mg/18.80	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 04- Relea	ope) [ms/L] 6.9500 2 PO ₄ -P ase Slope	SAUR 9/g MLVSS, 0.82 Clar CO AA2 Spec	NPF fier IDC PC oncentratio [mg/L] 0.2 diffic PO ₄ - se Rate	<0.20 R (NOx-N [mg/L/n 1.33 04-P Max n AA3 Releas	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope	[mg/ D ₄ -P A AA3 P I	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.26 3 Specific	R SS/hr] NO ₃ -N rotration VL 57 ic PO ₄ - Rate
Actual: [days 4.60 Highest / Conce [m	2 1 1 5 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 111 96 92 52 52 50 60 60 63 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P [m] 1	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel	FR DO N FR	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1 P R	Clarifier ID [mg/ 18.80 Specific PC elease Rat	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 04- AA2 e Relea nr] [m	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope gg/L/hr]	SAUR 9/g MLVSS, 0.82 Clar NOx-N C AA2 Spec P Releas [mg/g ML	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr]	(NOx-N [mg/L/n 1.33 0 ₄ -P Ma: n AA3 Releas	N Slope) nin] A ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope	[mg/ D ₄ -P A AA3 P I	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.28 3 Specifi Release	R SS/hr] NO ₃ -N NO ₃ -N Irration L] 57 ic PO ₄ - Rate SS/hr]
Actual: [days 4.60 Highest / Conce [m 5	2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 111 96 92 52 52 50 60 60 63 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel	SF DO N I I Img/g N 7 A1 PO ₄ -P ease Slop	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1 P R	Clarifier ID [mg/18.80	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 04- AA2 e Relea nr] [m	ope) [ms/L] 6.9500 2 PO ₄ -P ase Slope	SAUR 9/g MLVSS, 0.82 Clar CO AA2 Spec	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr]	(NOx-N [mg/L/n 1.33 0 ₄ -P Ma: n AA3 Releas	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope	[mg/ D ₄ -P A AA3 P I	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.26 3 Specific	R SS/hr] NO ₃ -N NO ₃ -N Irration L] 57 ic PO ₄ - Rate SS/hr]
Actual: [days 4.60 Highest / Conce [m 5.5	Part Part	9/L] OO 1 1 1 96 92 52 52 50 60 60 63 54 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P [mg/L] 0.27	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel	FR DO N SF	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1 P R [mg/	Clarifier ID [mg/ 18.80 Specific PC telease Rat [g MLVSS/6] 8.06	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 D ₄ - AA2 e Releant] [m	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope 19/L/hr] 14.42	SAUR 9/g MLVSS, 0.82 Clar CO AA2 Spec P Releas [mg/g ML	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr] 96	(NOx-N [mg/L/n 1.33 1.33 1.4-P Max n AA3 Releas	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope //L/hr]	[mg/ D ₄ -P A AA3 P I	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.28 3 Specifi Release	R SS/hr] NO ₃ -N NO ₃ -N Irration L] 57 ic PO ₄ - Rate SS/hr]
Actual: [days 4.60 Highest / Conce [m 5	2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 1 1 1 96 92 52 52 50 60 60 60 64 48 43 40 44 43 78 Obic MLSS [mg/L] 2320 PUR (PO ₄ -P [mg/L] 0.27 A4 Specific PC	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 ARel	FR DO N FR DO N	In AER [unitless 6.84 PUR LVSS/hr] 01 AA1 PR mg/	Clarifier ID [mg/ 18.80 Specific PC elease Rat g MLVSS/i 8.06 PO ₄ - A	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 0 ₄ - AA2 e Relea hr] [m 1	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope 14.42 AA6 Spec	SAUR 9/g MLVSS, 0.82 Clar C AA2 Spec P Releas [mg/g ML 12.	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr] 96	(NOx-N [mg/L/n 1.33 0 ₄ -P Ma: n AA3 Releas [mg 7	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope y/L/hr] .25 A1) Spec	[mg/	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.25 3 Specifi Release a/g MLV 6.52	SS/hr] NO ₃ -N tration VL To For PO ₄ - Rate SS/hr]
Actual: [days 4.60 Highest / Conce [m 5.5	2 1 1 5 5 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6 6	9/L] OO 1 1 1 96 92 52 52 50 60 60 63 54 48 43 40 44 43 78 obic MLSS [mg/L] 2320 PUR (PO ₄ -P [mg/L] 0.27	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 ARel	FR DO N FR DO N	fin AER [unitless 6.84 PUR LVSS/hr] .01 AA1 P R [mg/	Clarifier ID [mg/ 18.80 Specific PC elease Rat g MLVSS/i 8.06 PO ₄ - A	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 D ₄ - AA2 e Releant] [m	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope 19/L/hr] 14.42	SAUR 9/g MLVSS, 0.82 Clar C AA2 Spec P Releas [mg/g ML 12.	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr] 96	(NOx-N [mg/L/n 1.33 0 ₄ -P Ma: n AA3 Releas [mg 7	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope //L/hr]	[mg/	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.28 3 Specifi Release	SS/hr] NO ₃ -N tration VL To For PO ₄ - Rate SS/hr]
Actual S [days 4.60] Highest A Conce [m 5	Part Part	9/L] OO 11 96 92 52 52 50 60 60 63 44 43 40 44 43 78 Obic MLSS [mg/L] 2320 PUR (PO ₄ -P [mg/L] AA6 NO) [mg/L] 0.27 A4 Specific PC P Release Rate	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel 4- AA5 PC Release	SR DO M [mg/g M 7 A1 PO ₄ -P 8.96 Slope F	fin AER [unitless 6.84] PUR ILVSS/hr] .01 AA1	Clarifier ID [mg/18.86] Specific PC telease Rat g MLVSS/I 8.06 PO ₄ - A Rate Rel	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 100 D ₄ - AA2 e Relea nr] [m 1 A6 PO ₄ -P ease Slope	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope g/L/hr] 14.42 AA6 Spec P Release	SAUR a/g MLVSS 0.82 Clar C AA2 Spec P Releas [mg/g ML 12. iffic PO ₄ - (AA se Rate	NPF hr] filer IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr] 96 1) COD Upi Slope	CO.2O R (NOx-N [mg/L/n 1.33 A-P Max n AA3 Releas [mg 7 take (A COI	N Slope) nin] k ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope y/L/hr] .25 A1) Spec O Uptake	[mg/	SNPF /g MLV 0.77 Ng. ANA Concent [mg/ 0.28 3 Specifi Release g/g MLV 6.52	R SS/hr] NO ₃ -N NO ₃ -N tration 'L'] 57 ic PO ₄ - Rate SS/hr]
Actual S [days 4.60 Highest A Conce [m 5	Part Part	9/L] OO 1 1 1 96 92 52 52 50 60 60 60 64 48 43 40 44 43 78 Obic MLSS [mg/L] 2320 PUR (PO ₄ -P [mg/L] 0.27 A4 Specific PC	Average AE [mg/L 0.96 Uptake Slope g/L/hr] 2.03 A Rel 4- AA5 PC Release	SR DO N [] SF [mg/g M 7 A1 PO ₄ -P ease Slop mg/L/hr] 8.96 24-P A7 A1 A1 B1 B1 B1 B1 B1 B1 B1 B1	In AER [unitless 6.84 PUR LVSS/hr] 01 AA1 PR mg/	Clarifier ID [mg/18.86] Specific PC telease Rat g MLVSS/I 8.06 PO ₄ - A Rate Rel	(NH4-N SIG [mg/L/hr] 1.41 C NH ₄ -N C L] 000 0 ₄ - AA2 e Relea hr] [m 1	ope) [mg/L] 6.9500 2 PO ₄ -P ase Slope 14.42 AA6 Spec	SAUR 3/g MLVSS, 0.82 Clar C AA2 Spec P Releas [mg/g ML 12. iffic PO ₄ - se Rate VSS/hr]	NPF fier IDC PC oncentratio [mg/L] 0.2 iffic PO ₄ - se Rate VSS/hr] 96	CO.2O R (NOx-N [mg/L/n 1.33 A-P Max n AA3 Releas [mg 7 take (A COI	N Slope) nin] x ANA PC Release [mg/L] 34.26 PO ₄ -P se Slope y/L/hr] .25 A1) Spec	[mg/	SNPF /g MLV 0.77 Avg. ANA Concent [mg/ 0.25 3 Specifi Release a/g MLV 6.52	R SS/hr NO ₃ -N tration L To r R R R R R R R R R R R R R R R R R R R

# Chem Us	ed	FeCl3 Use		FeCl3		Biosolid Influ Flow	uent	ARC	Y Flow	CR	CY Flow	NRCY Flow
[lbs]		[hr]		[mg/L	1	[MGD]		[M	GD]	[MGD]	[MGD]
0		0.00	0	0.000)	18.97		17	.12	1	0.298	0
ANA MLTSS	ANA	4 % VOL	ANA	MLVSS	ANX MLTSS	ANX % VOL	ANX	MLVSS	AER ML	TSS	AER % VC	L AER MLVSS
[mg/L]		[%]	[1	mg/L]	[mg/L]	[%]	[m	g/L]	[mg/l	L]	[%]	[mg/L]
1340		83		1112	2320	74	17	717	2320)	74	1717

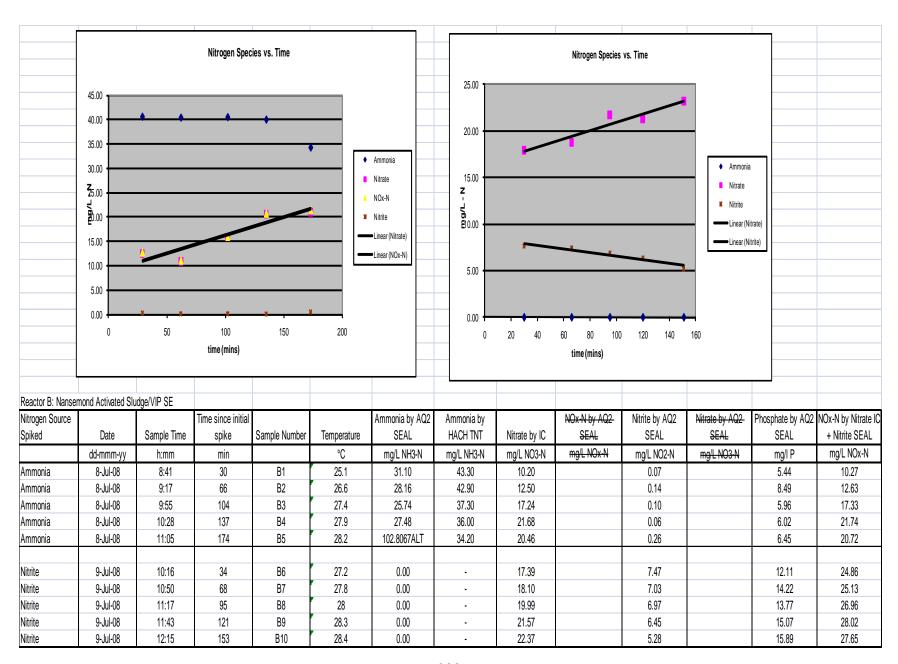


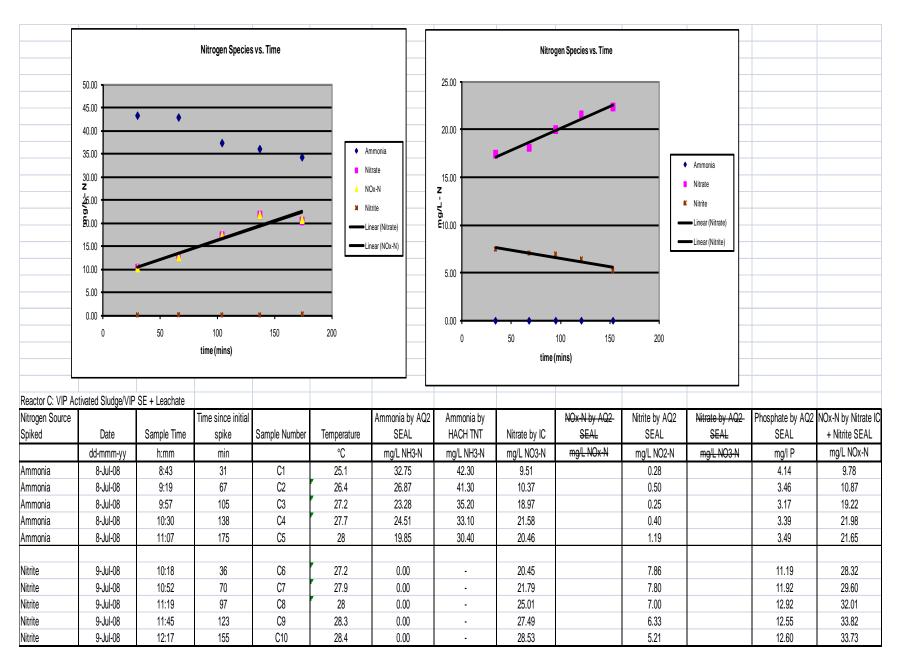


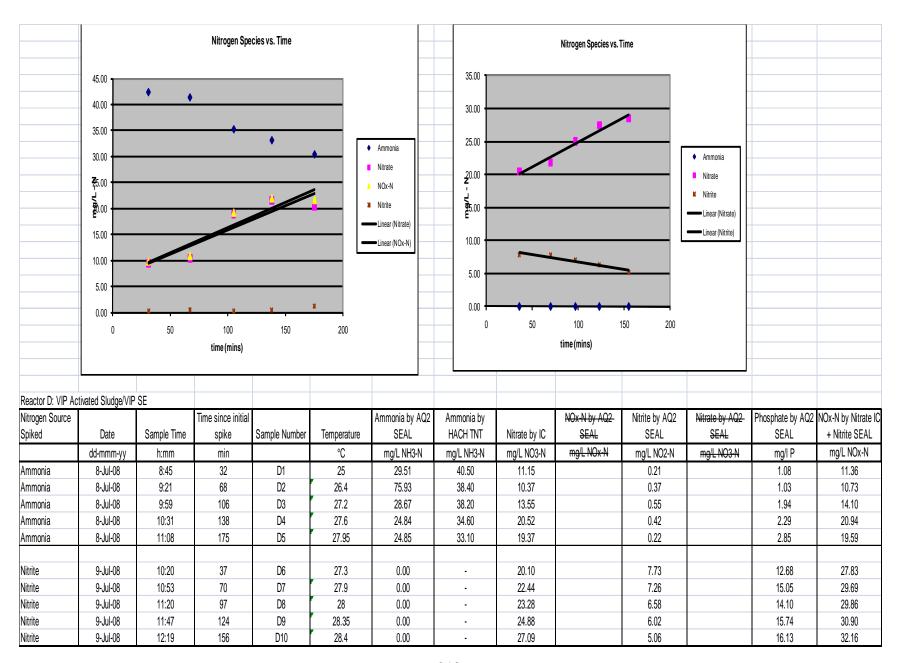


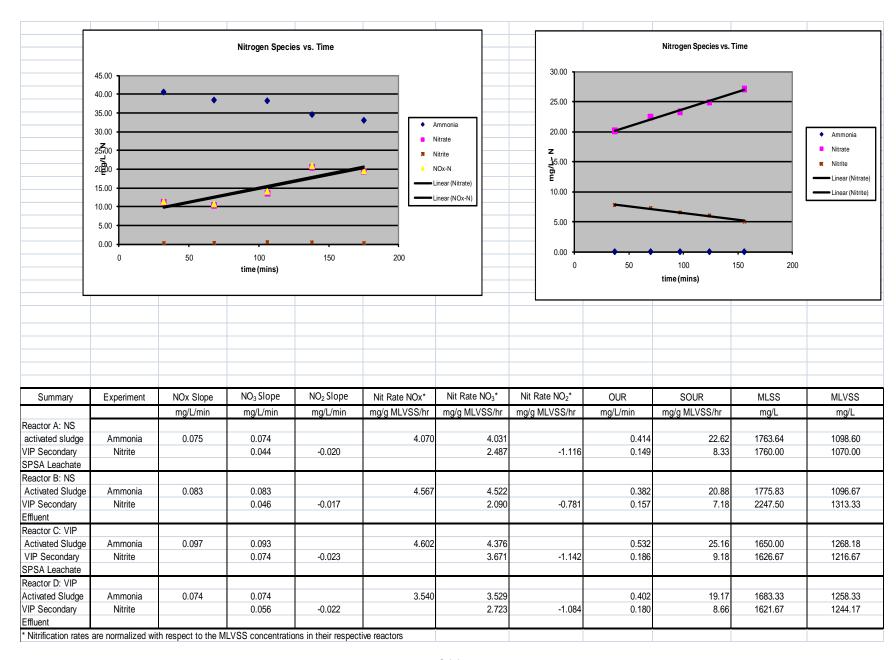
7.7 Appendix H – Batch Rate Experiment DataWeek 3 – Leachate

Sample Data R	Report for Nanse	emond Nitrificati	on innibition Stu	idy									
AOB & NOB Exp	perimentation:												
I. AOB													
a. Spiked four	3L reactors with !	50 mg/L NH ₄ .						-					
b. Each reacto	or is running conti	nously by use of st	ir bars.					-					
c. 2 L of the di	iluent source is ad	lded to the reacto	rs.					-					
d. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 8.5 L. e. Constant DO and pH were monitored and logged throughout the experiment.													
f. Oxygen/air	r blend was sparge	ed into the reactor	s to maintain DO le	evels through soler	noid valves.			-					
g. LabView so	ftware was used t	to manage DO and	pH recording as w	ell as program and	d implement the so	olenoid valves.		-					
h. 5 samples v	were collected ove	er a period of 3 ho	urs through sampli	ing ports on the re	actors and filtered	l into sample tubes th	rough millipore 0.	45 μm filters					
						1:135 dilution) based		·					
		•			,	•	•						
II. NOB													
	r 3L reactors with 3	25 mg/L NO ₂ afte	rammonia levels v	vere < 2 mg/L NH	N which was check	ked through the use o	of HACH TNT 831 A	mmonia					
method.													
	or is running conti	nously by use of st	ir hars										
				experiment.				-					
c. Constant DO and pH were monitored and logged throughout the experiment.													
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.													
						olenoid valves							
e. LabView so	ftware was used t	to manage DO and	pH recording as w	ell as program and	d implement the so		arough millinare ()	45 um filters					
e. LabView so f. 5 samples v	ftware was used t were collected ove	to manage DO and er a period of 3 ho	pH recording as w urs through sampli	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
e. LabView so f. 5 samples v	ftware was used t were collected ove	to manage DO and er a period of 3 ho	pH recording as w urs through sampli	vell as program and ing ports on the re	d implement the so actors and filtered			45 μm filters.					
e. LabView so f. 5 samples v	ftware was used t were collected ove	to manage DO and er a period of 3 ho	pH recording as w urs through sampli	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
e. LabView so f. 5 samples v	ftware was used t were collected ove	to manage DO and er a period of 3 ho	pH recording as w urs through sampli	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
e. LabView so f. 5 samples v	ftware was used t were collected ove	to manage DO and er a period of 3 ho	pH recording as w urs through sampli	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used to were collected over and C which incorp	to manage DO and er a period of 3 ho porated leachate a	pH recording as w urs through sampl ddition were spike	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used to were collected over and C which incorp	to manage DO and er a period of 3 ho	pH recording as w urs through sampl ddition were spike	vell as program and ing ports on the re	d implement the so actors and filtered	l into sample tubes th		45 μm filters.					
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e. LabView so f. 5 samples v i. Reactors A a Reactor A: Nanser Alitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	mond Activated Slamm-yy 8-Jul-08 8-Jul-08 8-Jul-08	udge/VIP SE + Lea Sample Time h:mm 8:39 9:12 9:52 10:25	pH recording as wurs through sample ddition were spike chate Time since initial spike min 29 62 102 135	Sample Number A1 A2 A3 A4	Temperature °C 24.8 26.1 27 27.6	Ammonia by AQ2 SEAL mg/L NH3-N 30.42 28.13 27.47 25.30	Ammonia by HACH TNT mg/L NH3-N 40.60 40.40 40.50 40.00	Nitrate by IC mg/L NO3-N 12.50 10.92 15.81 20.60	SEÁL	SEAL mg/L NO2-N 0.22 0.18 0.12 0.06	SEÁL	SEAL mg/I P 6.92 6.53 6.63 7.21	+ Nitrite SEAL mg/L NOx-N 12.73 11.09 15.93 20.66
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e. LabView so f. 5 samples v i. Reactors A a	mond Activated Sla dd-mmm-yy 8-Jul-08 8-Jul-08 9-Jul-08	to manage DO and er a period of 3 ho porated leachate a udge/VIP SE + Lea Sample Time h:mm 8:39 9:12 9:52 10:25 11:03	pH recording as we are through sample ddition were spike chate Time since initial spike min 29 62 102 135 173 30	Sample Number A1 A2 A3 A4 A5	Temperature °C 24.8 26.1 27 27.6 27.9	Ammonia by AQ2 SEAL mg/L NH3-N 30.42 28.13 27.47 25.30 24.55	Ammonia by HACH TNT mg/L NH3-N 40.60 40.40 40.50 40.00 34.20	Nitrate by IC mg/L NO3-N 12.50 10.92 15.81 20.60 20.94	SEÁL	SEAL mg/L NO2-N 0.22 0.18 0.12 0.06 0.41 7.66	SEÁL	SEAL mg/l P 6.92 6.53 6.63 7.21 8.62 12.17	+ Nitrite SEAL mg/L NOx-N 12.73 11.09 15.93 20.66 21.35
e. LabView so f. 5 samples v i. Reactors A a Reactor A: Nanser Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia Vitrite	mond Activated Sla dd-mmm-yy 8-Jul-08 8-Jul-08 9-Jul-08 9-Jul-08	sample Time h:mm 8:39 9:12 9:52 10:25 11:03	pH recording as we are through sample ddition were spike chate Time since initial spike min 29 62 102 135 173 30 66	Sample Number A1 A2 A3 A4 A5 A6 A7	Temperature °C 24.8 26.1 27 27.6 27.9 26.9 27.6	Ammonia by AQ2 SEAL mg/L NH3-N 30.42 28.13 27.47 25.30 24.55	Ammonia by HACH TNT mg/L NH3-N 40.60 40.40 40.50 40.00 34.20	Nitrate by IC mg/L NO3-N 12.50 10.92 15.81 20.60 20.94 17.94 18.76	SEÁL	SEAL mg/L NO2-N 0.22 0.18 0.12 0.06 0.41 7.66 7.46	SEÁL	SEAL mg/l P 6.92 6.53 6.63 7.21 8.62 12.17 12.00	+ Nitrite SEAL mg/L NOx-N 12.73 11.09 15.93 20.66 21.35 25.60 26.22









Sample Data Report for Nansemond Nitrification Inhibition Study Bio-P Experimentation: I. Bio - P a. Spiked four 3L reactors with 5 mg/L PO₄-P to raise initial PO₄-concentration to roughly 15 mg/L PO₄-P. (Initial PO₄- concentration was determined through HACH TNT PO₄- Tubes at the end of the AOB/NOB experimentation). b. The Bio-P experiment is run through an uptake/release/uptake method were the four reactors are aerobic/anaerobic/aerobic once more. During the release phase 200 mg/L of NaAc was added for COD manually to all 4 reactors. c. Each reactor is running continously by use of stir bars . d. 2L of the diluent source is added to the reactors. e. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 8.5 L.

Reactor A: Nansemond Activated Sludge/ VIP SE + Leachate

9-Jul-08

18:50

260

A12

Uptake 2

f. Constant DO and pH were monitored and logged throughout the experiment.

g. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves during the uptake phases and then these valves were turned off and the oxygen/air was cut-off and then the system was deaerated and the DO was allowed to drop to 0 before beginning the release phase. During the release phase nitrogen was sparged into the reactors.

h. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.

i. 4 samples were collected over a period of 1.5 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 µm filters during both uptake phases and 4 samples were collected over a period of 1 hour for the release phase.

j. Reactors A and C which incorporated leachate addition were spiked with 15 mL of the SPSA leachate (1:135 dilution) based on plant flow.

Phosphate by AQ2 Ammonia by AQ2 NOx-N by Nitrate IC Time since initial NOx-N by AQ2 Nitrite by AQ2 SEAL SEAL + Nitrite SEAL Sample Time Sample Number Temperature Phosphate by TNT SEAL SEAL Nitrate by IC SEAL Phosphorus Test Date spike mg/L P mg/l P mg/L NH3-N ma/L NO3-N ma/L NOx-N mg/L NO2-N mg/L NO3-N ma/L NOx-N dd-mmm-yy h:mm min 14:46 21 22 16.60 16.66 0.00 0.00 14.76 Uptake 1 9-Jul-08 A1 14.76 Uptake 1 9-Jul-08 15:05 40 A2 23.6 16.40 18.38 2.68 15.18 0.00 15.18 Uptake 1 9-Jul-08 15:26 61 A3 24.8 16.20 18.35 0.00 16.12 0.00 16.12 Uptake 1 9-Jul-08 15:53 83 A4 17.00 18.29 0.00 17.23 0.00 17.23 26 Release 9-Jul-08 16:30 120 A5 27.03 18.75 17.25 0.00 4.53 0.18 4.71 Release 9-Jul-08 16:46 136 A6 27.35 29.00 25.87 0.00 7.95 0.13 8.09 9-Jul-08 16:58 148 Α7 27.4 37.00 33.40 0.00 9.57 0.25 9.82 Release Release Α8 55.00 9.15 9-Jul-08 17:22 172 27.8 42.71 0.00 0.60 9.75 Uptake 2 9-Jul-08 207 A9 54.00 46.53 0.00 10.93 17:57 28.1 0.41 11.34 Uptake 2 9-Jul-08 18:11 221 A10 28.18 55.00 46.54 0.00 10.38 0.22 10.60 Uptake 2 9-Jul-08 18:30 240 A11 28.2 55.00 47.41 0.00 12.56 0.04 12.60

49.04

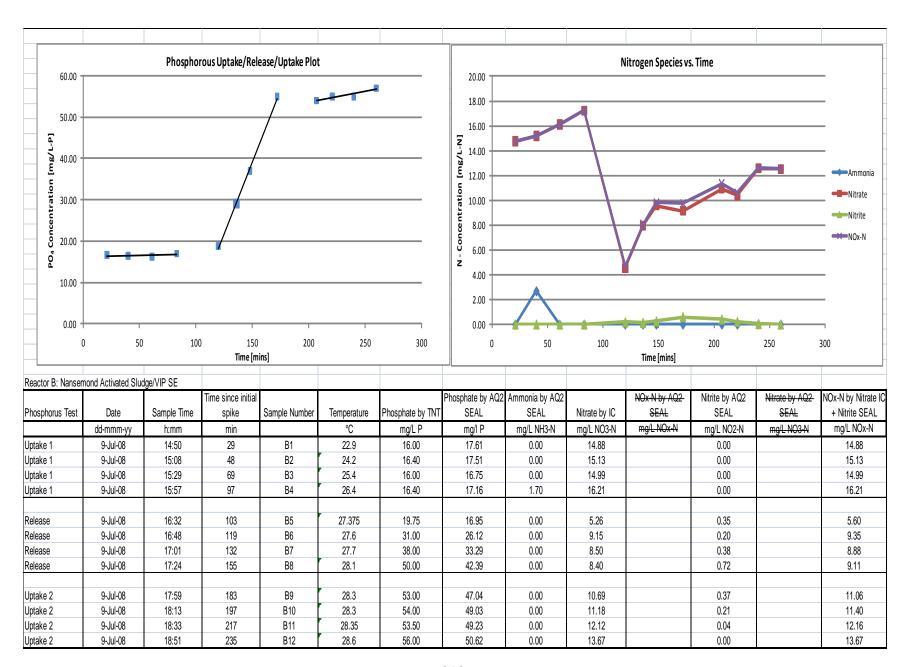
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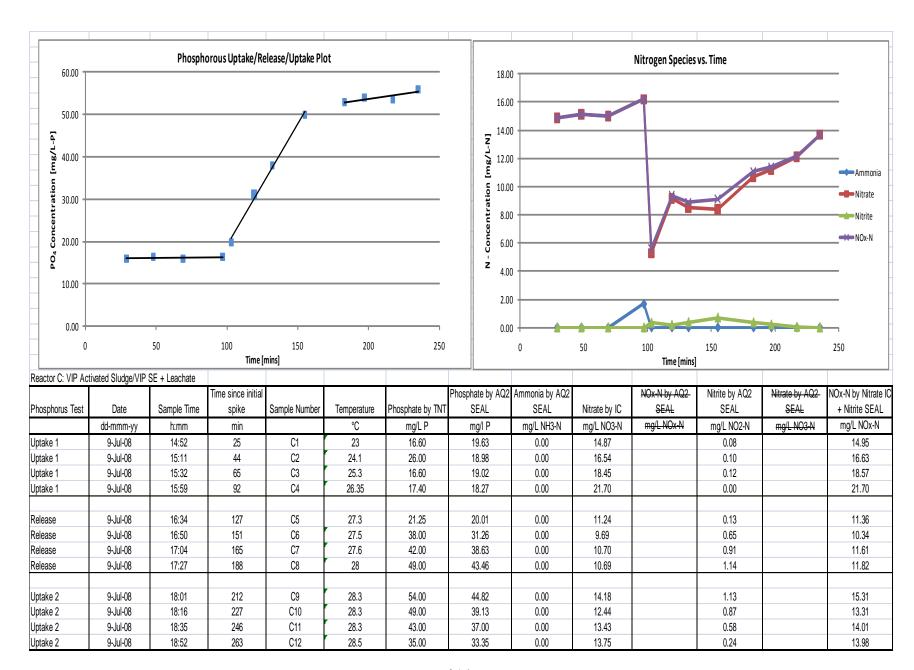
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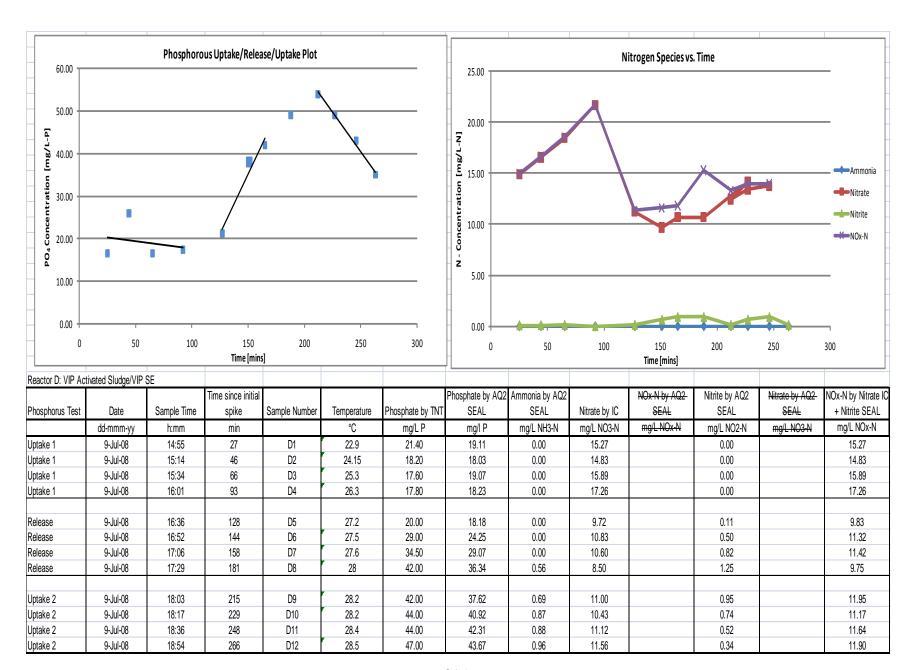
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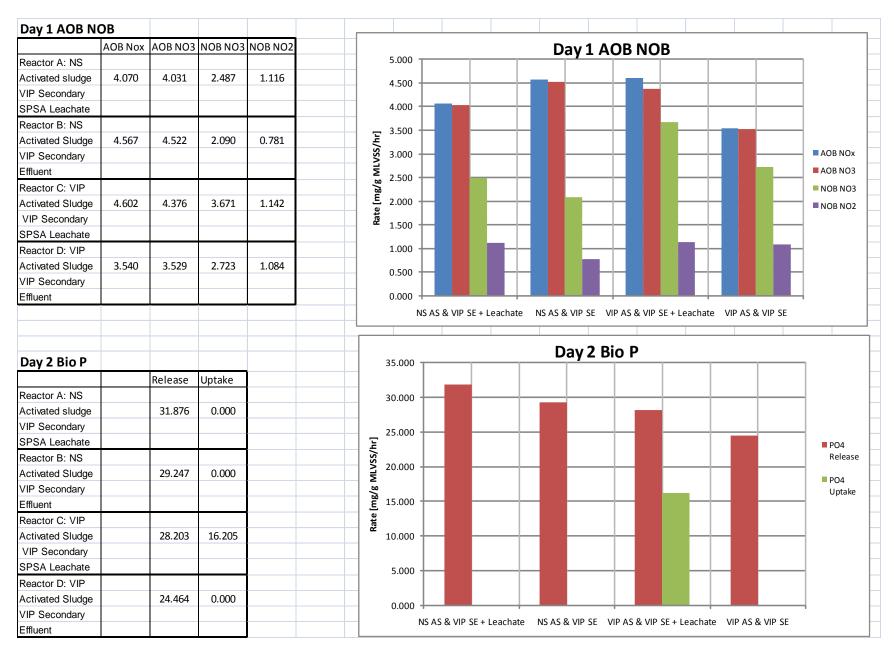
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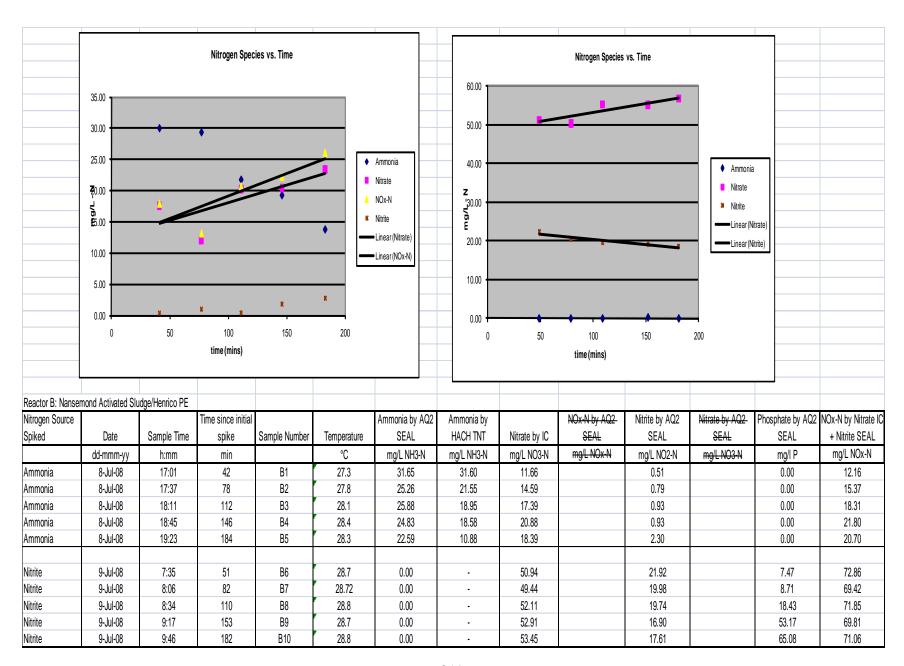


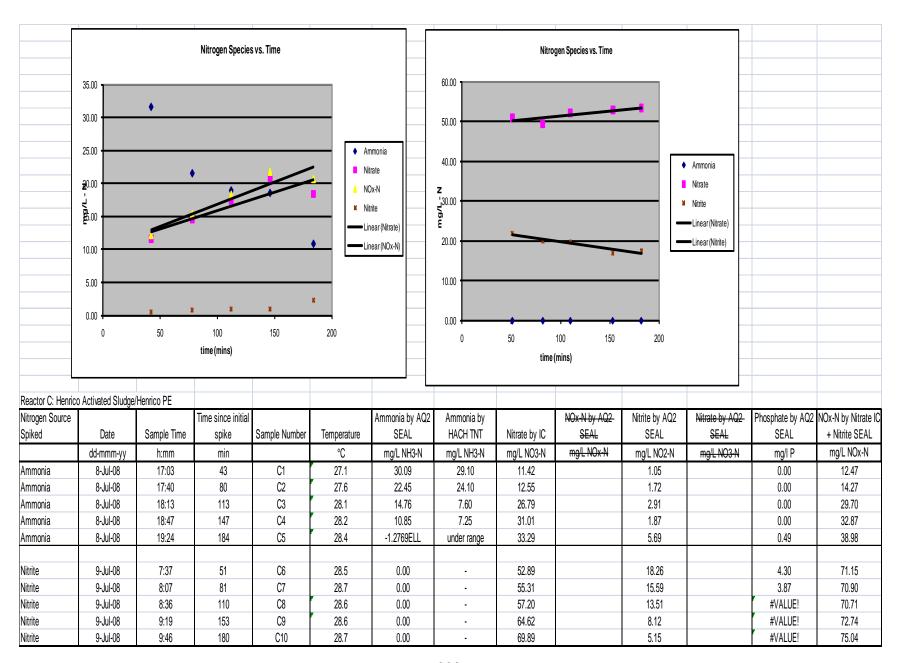


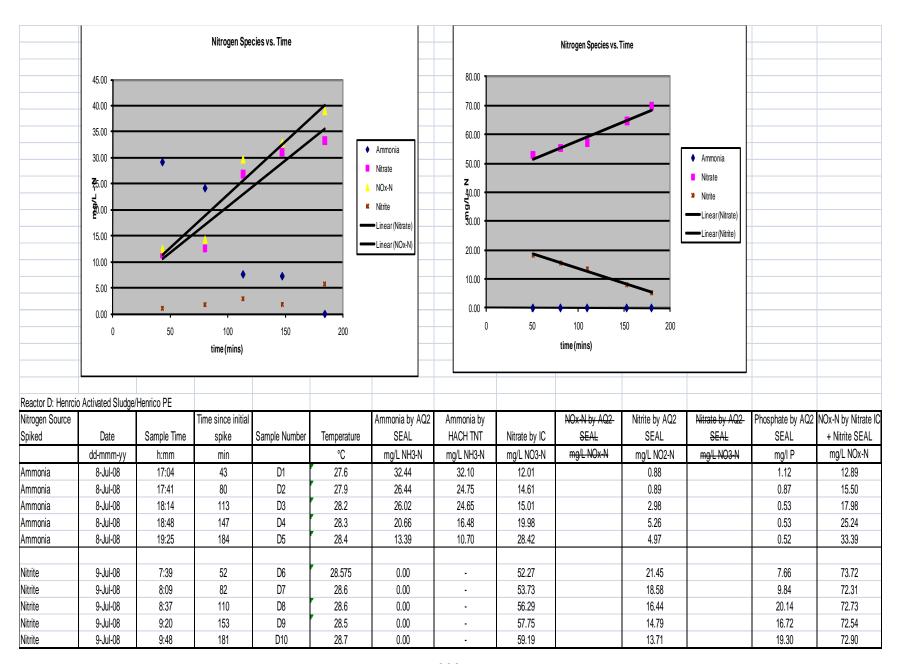


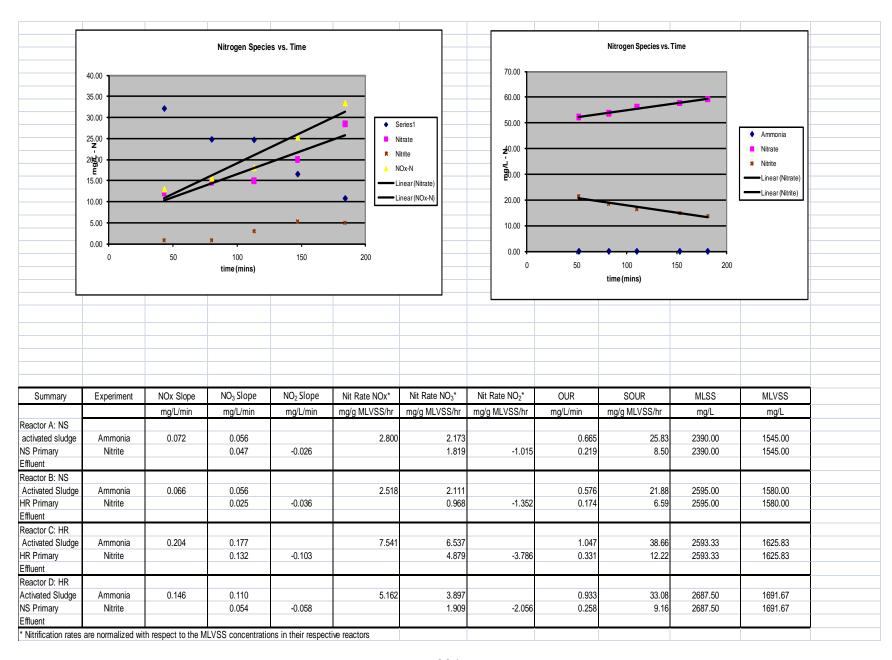
Week 4 – Henrico County Biomass Day 1 AOB/NOB

			Biomass I)R/NOB								
Sample Data R	Report for Nanse	emond Nitrificati	ion Inhibition Stu	ıdy									
AOB & NOB Exp	perimentation:												
I. AOB													
	r 3L reactors with 1	25 mg/L NH ₄ .											
		nously by use of st	tir bars .										
	-	Ided to the reacto											
	d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20L to 9 L.												
e. Constant DO and pH were monitored and logged throughout the experiment.													
			s to maintain DO l		noid valves								
					d implement the so	olenoid valves.							
						l into sample tubes t	hrough millinore 0	45 um filters					
ii. 5 sumples v	were concered ove	era periodor 5 no	rais tillough samp	iii g ports on the re	actors and mitered	anto sumple tubes t	inough minpore o.	+5 μm mens					
II. NOB													
-	r21 reactors with	25 ma/l NO - afta	rammonia lovolev	word <1 mg/L NU	N which was shool	ked through the use	of LIACLITAIT 021 A	mmonia					
			in overnight to cor			keu tili ougii tile use	UI HACH INT 651 P	illillollia					
		nously by use of st		isuille dii presenti	allillollia.								
1	•		ur bars. ed throughout the	ovnoriment									
1			•	•	n a id valva a								
			s to maintain DO			مامه منظييمان م							
					d implement the so			4E ('1)					
T. 5 samples v	were collected ove	era period of 3 no	ours through samp	ling ports on the re	eactors and flitered	l into sample tubes t	nrougn millipore v.	45 μm fliters.					
Reactor A: Nanser	mond Activated Slu	udge/Nansemond F	PE										
Nitrogen Source	D-1-	OI- T	Time since initial	O In November	T	Ammonia by AQ2	Ammonia by	Nitrata had IO	NOx-N by AQ2 SEAL	Nitrite by AQ2	Nitrate by AQ2		NOx-N by Nitrate IC
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL NUO N	HACH TNT	Nitrate by IC	_	SEAL	SEAL 37 TH NOON	SEAL	+ Nitrite SEAL
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N
Ammonia	8-Jul-08	16:59	41	A1	27.8	35.36	30.00	17.55		0.35		1.39	17.90
Ammonia	8-Jul-08	17:35	77	A2	28	33.72	29.35	12.11		0.99		0.41	13.10
Ammonia	8-Jul-08	18:09 18:44	111 146	A3 A4	28.2	29.11 26.34	21.75	20.20 20.37		0.46 1.87		0.00	20.66 22.24
Ammonia	8-Jul-08 8-Jul-08	18:44	183	A4 A5	28.3 28.4	19.46	19.25	20.37		2.75		0.00	26.09
Ammonia	0-Jul-00	19.21	103	AO	20.4	19.40	13.78	23.34		2.10		0.00	20.09
Vitrite	9-Jul-08	7:32	49	A6	28.5	0.00		51.29		22.45		6.87	73.74
Nitrite	9-Jul-08	8:02	79	A6 A7	28.7	0.00	-	51.29		20.47		6.74	70.86
Nitrite	9-Jul-08	8:32	109	A7 A8	28.6	0.00	-	55.29		19.54		#VALUE!	74.83
Nitrite	9-Jul-08	9:15	152	A0 A9	28.6	0.00	-	55.23		19.15		#VALUE!	74.03
	9-Jul-08	9:44	181	A10	_		_	56.86		18.62			
Nitrite	9-Jul-08	9:44	181	A10	28.775	0.00	-	56.86		18.62		36.15	75.48



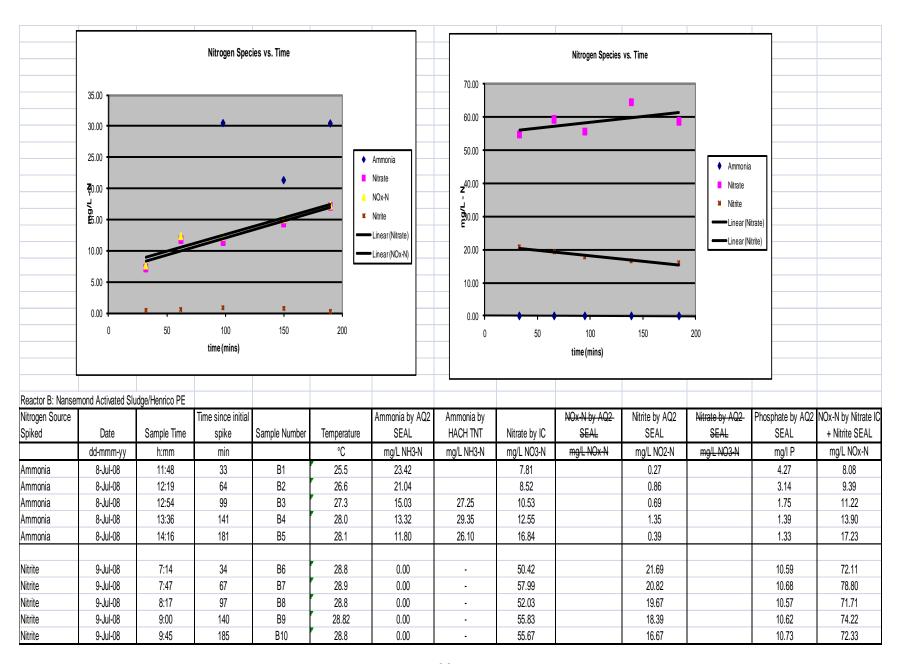


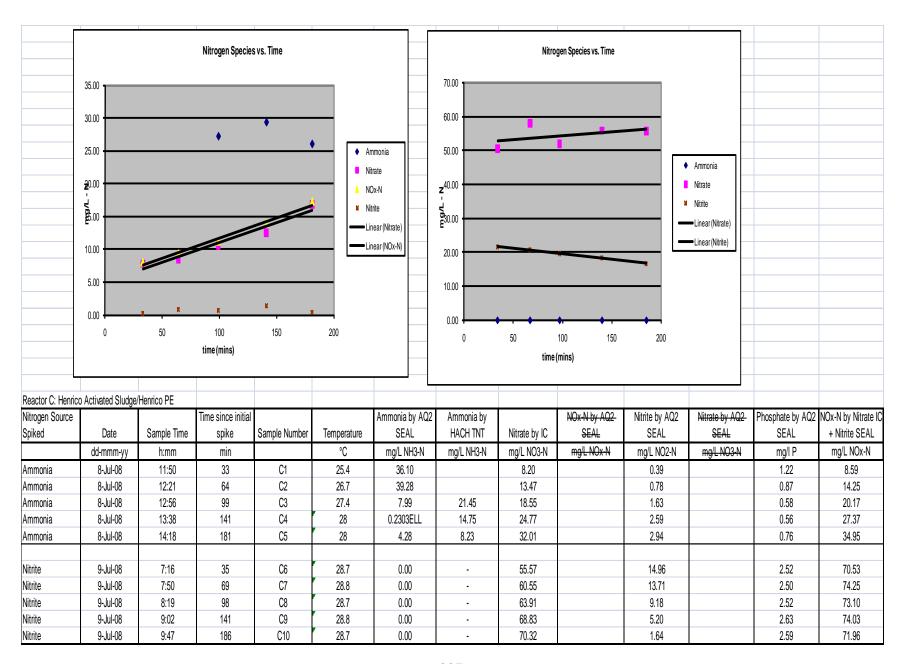


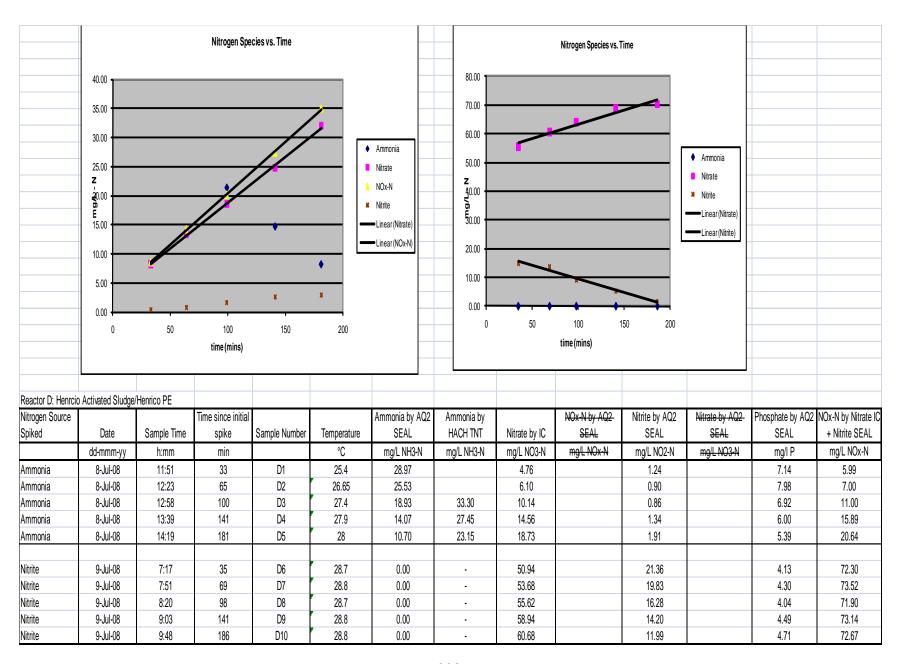


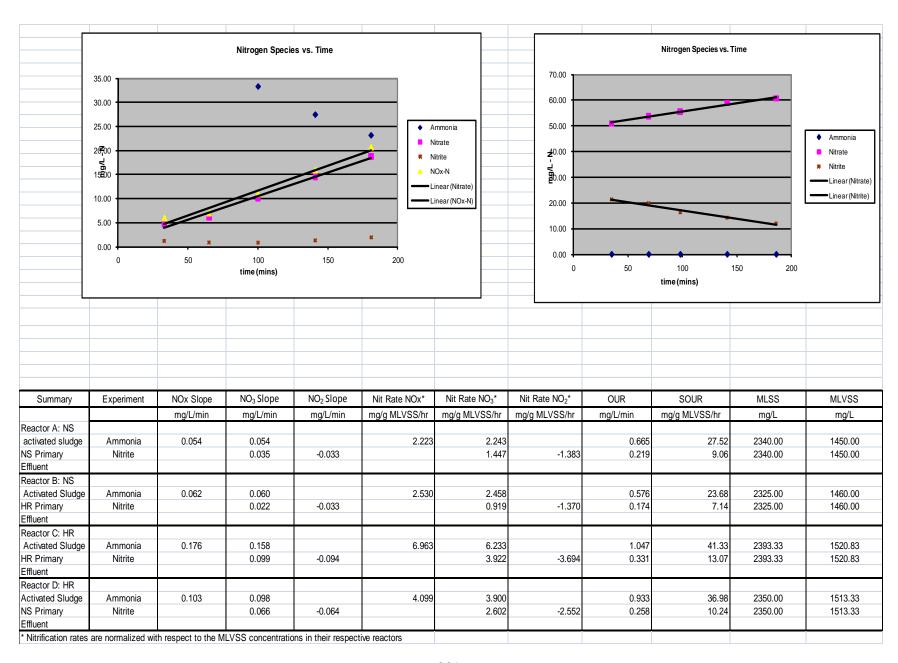
Week 4 – Henrico County Biomass Day 2 AOB/NOB

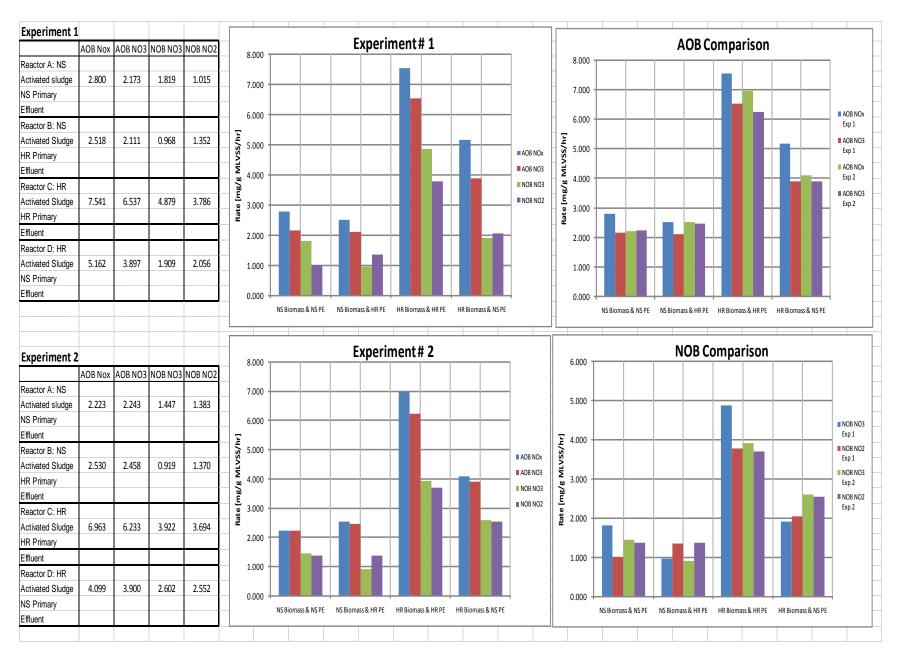
Sample Data D													
Danipie Dala N	Report for Nanse	emond Nitrificati	on Inhibition Stu	dy									
AOB & NOB Ext	perimentation:												
I. AOB													
	r 3L reactors with 2	25 mg/L NH.											
	or is running conti		ir hars										
	liluent source is ad												
	d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20L to 9 L.												
e. Constant DO and pH were monitored and logged throughout the experiment.													
	ir blend was sparge												
						alamaidualuaa							
	oftware was used t	-						ar Ch					
n. 5 samples v	were collected ove	era period of 3 no	urs through samp	ing ports on the re	eactors and filtered	d into sample tubes t	nrougn millipore U.	45 µm filters					
II. NOB													
						ked through the use	of HACH TNT 831 A	mmonia					
-	dition all reactors v		U	sume all present	ammonia.								
	or is running conti												
	O and pH were mo												
	ir blend was sparge			-									
e. LabView so	oftware was used t	o manage DO and	pH recording as w	vell as program an	d implement the s	olenoid valves.							
f. 5 samples v	were collected ove	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	d into sample tubes t	hrough millipore 0.	45 μm filters.					
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Docator & Monaco	The stand Oliver	des (Management F											
Reactor A: Nanser	mond Activated Slu	udge/Nansemond F	PE										
Reactor A: Nanser	mond Activated Slu	udge/Nansemond F	E										
	mond Activated Slu	udge/Nansemond R	E Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC
Nitrogen Source	mond Activated Slu	udge/Nansemond F		Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOX-N by AQ2- SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2- SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL
Nitrogen Source			Time since initial	Sample Number	Temperature °C	SEAL	,	Nitrate by IC	,	,	SEAL		NOx-N by Nitrate IC + Nitrite SEAL mg/L NOx-N
Nitrogen Source Spiked	Date dd-mmm-yy	Sample Time	Time since initial spike min		€	SEAL mg/L NH3-N	HACH TNT	mg/L NO3-N	SEÁL	SEAL mg/L NO2-N		SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
Nitrogen Source Spiked Ammonia	Date	Sample Time	Time since initial spike	Sample Number A1 A2		SEAL	HACH TNT		SEÁL	SEÁL	SEAL	SEAL	+ Nitrite SEAL
Nitrogen Source Spiked Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08	Sample Time h:mm	Time since initial spike min 32	A1	°C 25.3	SEAL mg/L NH3-N 29.63	HACH TNT	mg/L NO3-N 7.27	SEÁL	SEAL mg/L NO2-N 0.40	SEAL	SEAL mg/I P 13.06	+ Nitrite SEAL mg/L NOx-N 7.67
Nitrogen Source Spiked Ammonia Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08 8-Jul-08	Sample Time h:mm 11:46 12:16	Time since initial spike min 32 62	A1 A2	°C 25.3 26.4	SEAL mg/L NH3-N 29.63 40.23	HACH TNT mg/L NH3-N	mg/L NO3-N 7.27 11.80	SEÁL	SEAL mg/L NO2-N 0.40 0.60	SEAL	SEAL mg/l P 13.06 8.42	+ Nitrite SEAL mg/L NOx-N 7.67 12.40
Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08	Sample Time h:mm 11:46 12:16 12:52	Time since initial spike min 32 62 98	A1 A2 A3	°C 25.3 26.4 27.2	SEAL mg/L NH3-N 29.63 40.23 20.20	HACH TNT mg/L NH3-N 30.55	mg/L NO3-N 7.27 11.80 11.47	SEÁL	SEAL mg/L NO2-N 0.40 0.60 0.82	SEAL	SEAL mg/l P 13.06 8.42 4.08	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29
Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08	Sample Time h:mm 11:46 12:16 12:52 13:34	Time since initial spike min 32 62 98 150	A1 A2 A3 A4	°C 25.3 26.4 27.2 27.9	SEAL mg/L NH3-N 29.63 40.23 20.20 17.69	HACH TNT mg/L NH3-N 30.55 21.30	mg/L NO3-N 7.27 11.80 11.47 14.38	SEÁL	SEAL mg/L NO2-N 0.40 0.60 0.82 0.74	SEAL	SEAL mg/l P 13.06 8.42 4.08 0.88	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29 15.12
Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08	Sample Time h:mm 11:46 12:16 12:52 13:34	Time since initial spike min 32 62 98 150	A1 A2 A3 A4	°C 25.3 26.4 27.2 27.9	SEAL mg/L NH3-N 29.63 40.23 20.20 17.69	HACH TNT mg/L NH3-N 30.55 21.30	mg/L NO3-N 7.27 11.80 11.47 14.38	SEÁL	SEAL mg/L NO2-N 0.40 0.60 0.82 0.74	SEAL	SEAL mg/l P 13.06 8.42 4.08 0.88	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29 15.12
Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08	Sample Time h:mm 11:46 12:16 12:52 13:34 14:14	Time since initial spike min 32 62 98 150 190	A1 A2 A3 A4 A5	°C 25.3 26.4 27.2 27.9 28.0	SEAL mg/L NH3-N 29.63 40.23 20.20 17.69 15.30	HACH TNT mg/L NH3-N 30.55 21.30 30.50	mg/L NO3-N 7.27 11.80 11.47 14.38 17.07	SEÁL	SEAL mg/L NO2·N 0.40 0.60 0.82 0.74 0.28	SEAL	SEAL mg/l P 13.06 8.42 4.08 0.88 0.00	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29 15.12 17.34
Nitrogen Source	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08 9-Jul-08	Sample Time h:mm 11:46 12:16 12:52 13:34 14:14 7:12	Time since initial spike min 32 62 98 150 190	A1 A2 A3 A4 A5	°C 25.3 26.4 27.2 27.9 28.0	SEAL mg/L NH3-N 29.63 40.23 20.20 17.69 15.30	HACH TNT mg/L NH3-N 30.55 21.30 30.50	mg/L NO3-N 7.27 11.80 11.47 14.38 17.07	SEÁL	SEAL mg/L NO2-N 0.40 0.60 0.82 0.74 0.28	SEAL	SEAL mg/l P 13.06 8.42 4.08 0.88 0.00 7.60	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29 15.12 17.34
Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia Nitrite	Date dd-mmm-yy 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08 8-Jul-08 9-Jul-08 9-Jul-08	Sample Time h:mm 11:46 12:16 12:52 13:34 14:14 7:12 7:45	Time since initial spike min 32 62 98 150 190 33 66	A1 A2 A3 A4 A5 A6 A7	°C 25.3 26.4 27.2 27.9 28.0 28.5 28.7	SEAL mg/L NH3-N 29.63 40.23 20.20 17.69 15.30 0.00 0.00	HACH TNT mg/L NH3-N 30.55 21.30 30.50	mg/L NO3-N 7.27 11.80 11.47 14.38 17.07 54.78 59.38	SEÁL	SEAL mg/L NO2-N 0.40 0.60 0.82 0.74 0.28 20.93 19.46	SEAL	SEAL mg/l P 13.06 8.42 4.08 0.88 0.00 7.60 8.66	+ Nitrite SEAL mg/L NOx-N 7.67 12.40 12.29 15.12 17.34 75.71 78.83



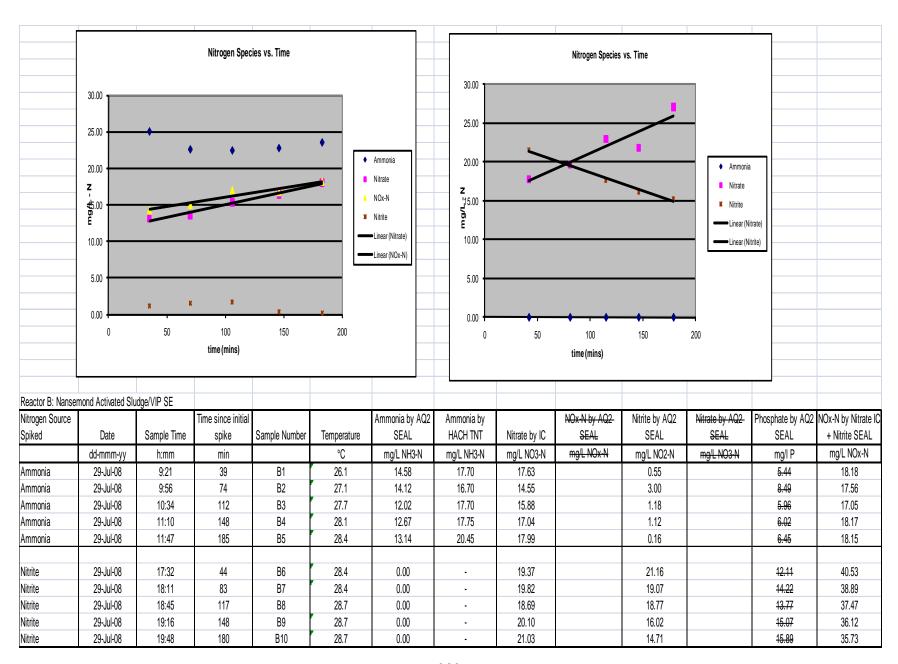


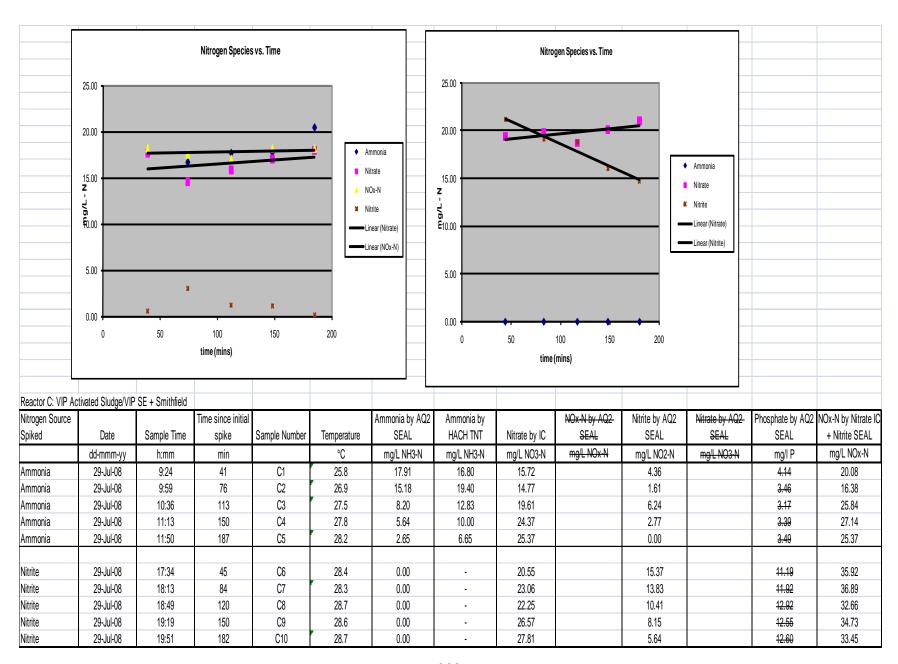


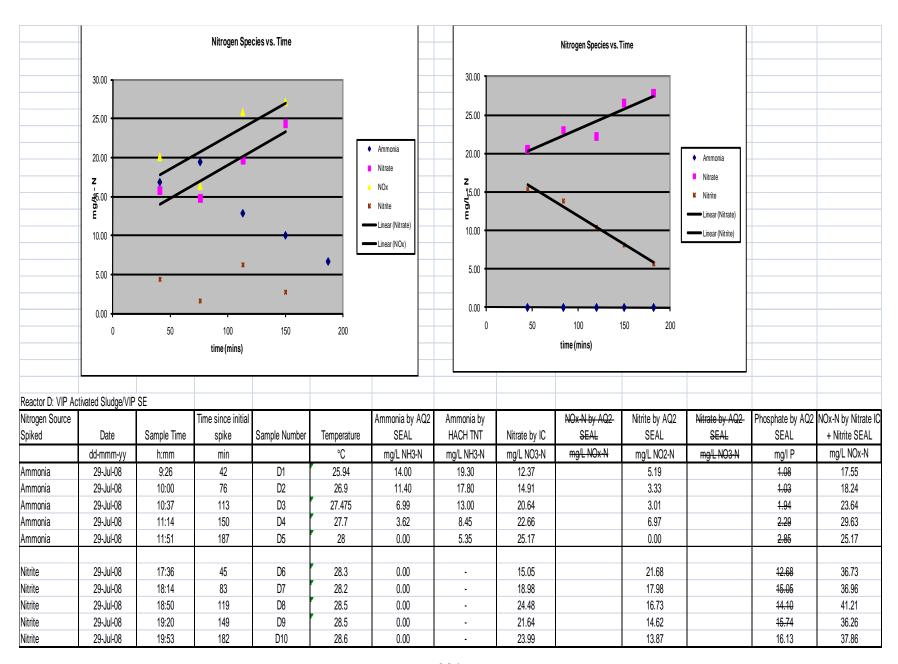


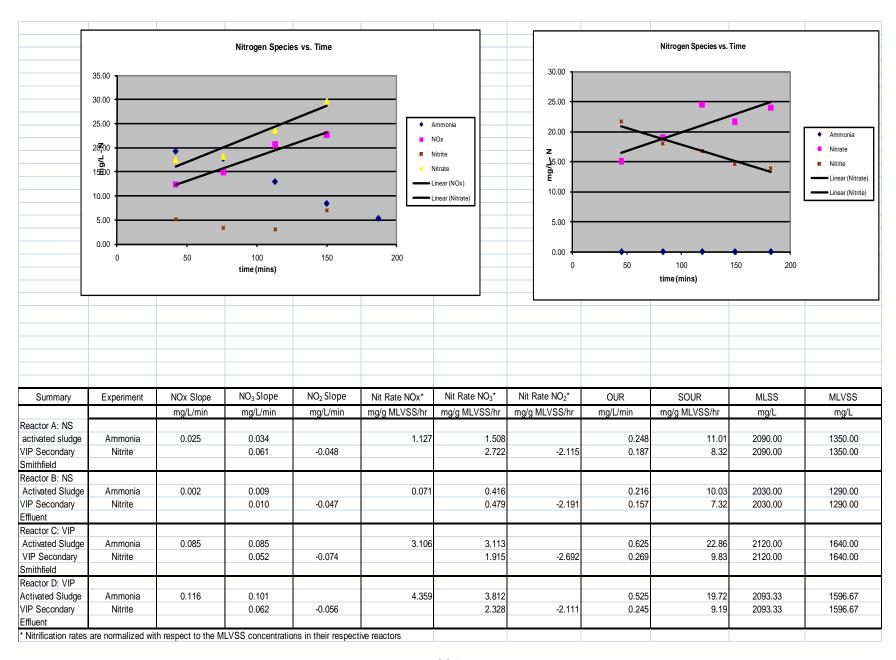


DOUBLE LIGHT K	Report for Nanse	emond Nitrificati	on Inhibition Stu	ıdv									
Jampic Dala N	CPOIL IOI TALIBO	cinoria ratambati		uy									
AOD O NOD F				1									
AOB & NOB Exp	perimentation:												
I. AOB	21	20 /											
	r 3L reactors with	O' 7											
	-	inously by use of s											
c. 2 L of the diluent source is added to the reactors. d. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 8.5 L. e. Constant DO and pH were monitored and logged throughout the experiment.													
		ed into the reactor											
					d implement the so								
h. 5 samples v	were collected ov	era period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t	hrough millipore 0.	45 μm filters					
i. Reactors A	and C which incor	rporated industrial	waste addition we	ere spiked with 20	0 mL of Smithfield	sample (1:10 dilutio	n).						
II. NOB													
a. Spiked four	r 3L reactors with	25 mg/L NO ₂ afte	rammonia levels v	vere < 2 mg/L NH ₃ -	N which was check	ed through the use	of HACH TNT 831 A	mmonia					
method.		5 .		G		· ·							
b. Each reacto	or is running conti	inously by use of st	ir bars.										
		onitored and logge		experiment.									
	•		-	•	noid valves.								
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.													
	e. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.												
e. LabView so	ftware was used		pH recording as w				hrough millinare ()	15 um filtars					
e. LabView so f. 5 samples v	oftware was used were collected ov	er a period of 3 ho	pH recording as w urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	er a period of 3 ho	pH recording as w urs through sampl	ling ports on the re	eactors and filtered			45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	er a period of 3 ho	pH recording as w urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	er a period of 3 ho	pH recording as w urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	er a period of 3 ho	pH recording as w urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used were collected over and C which incor	er a period of 3 ho porated industrial	pH recording as w urs through sampl waste addition we	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used were collected over and C which incor	er a period of 3 ho	pH recording as w urs through sampl waste addition we	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used were collected over and C which incor	er a period of 3 ho porated industrial	pH recording as w urs through sampl waste addition we	ling ports on the re	eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a	oftware was used were collected over and C which incor	er a period of 3 ho porated industrial	pH recording as w urs through sampl waste addition we ithfield	ling ports on the re	eactors and filtered	into sample tubes t ample (1:10 dilutior).	45 μm filters.					
e. LabView so f. 5 samples v i. Reactors A a leactor A: Nanser	oftware was used were collected over and C which incor	er a period of 3 ho porated industrial udge/VIP SE + Sm	pH recording as w urs through sampl waste addition we ithfield	ing ports on the re re spiked with 200	eactors and filtered O mL of Smithfield s	into sample tubes t ample (1:10 dilution	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser	oftware was used were collected over and C which incor	er a period of 3 ho porated industrial udge/VIP SE + Sm Sample Time	pH recording as w urs through sampl waste addition we ithfield Time since initial spike	ling ports on the re	eactors and filtered of mL of Smithfield s mL of Smithfield s	into sample tubes t ample (1:10 dilution Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	SEÁL	SEAL	SEÁL	SEAL	+ Nitrite SEAL
e. LabView so f. 5 samples v i. Reactors A a Reactor A: Nanser litrogen Source	oftware was used were collected over and C which incord and Activated SI Date dd-mmm-yy	er a period of 3 ho porated industrial udge/VIP SE + Sm Sample Time h:mm	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min	ing ports on the re ere spiked with 200 Sample Number	eactors and filtered of mL of Smithfield s Temperature	into sample tubes t ample (1:10 dilution Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC mg/L NO3-N	,	SEAL mg/L NO2-N	,	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser litrogen Source piked	oftware was used were collected over and C which incord and C which incord and Activated SI Date dd-mmm-yy 29-Jul-08	er a period of 3 ho porated industrial udge/VIP SE + Sm Sample Time h:mm 9:16	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35	sample Number	Temperature °C 25.6	into sample tubes t ample (1:10 dilution Ammonia by AQ2 SEAL mg/L NH3-N 18.27	Ammonia by HACH TNT mg/L NH3-N 25.10	Nitrate by IC mg/L NO3-N 13.14	SEÁL	SEAL mg/L NO2-N 1.15	SEÁL	SEAL mg/I P 6.92	+ Nitrite SEAL mg/L NOx-N 14.29
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser litrogen Source piked mmonia mmonia	oftware was used were collected over and C which incord and C which incord and Activated SI Date dd-mmm-yy 29-Jul-08 29-Jul-08	er a period of 3 ho porated industrial udge/VIP SE + Sm Sample Time h:mm 9:16 9:51	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70	Sample Number A1 A2	Temperature °C 25.6 26.76666667	Ammonia by AQ2 SEAL mg/L NH3-N 18.27 18.26	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65	Nitrate by IC mg/L NO3-N 13.14 13.56	SEÁL	SEAL mg/L NO2-N 1.15 1.49	SEÁL	SEAL mg/l P 6.92 6.53	+ Nitrite SEAL mg/L NOx-N 14.29 15.05
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser litrogen Source piked mmonia mmonia mmonia	mond Activated Sl Date dd-mmm-yy 29-Jul-08 29-Jul-08	udge/VIP SE + Sm Sample Time h:mm 9:16 9:51 10:27	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70 106	Sample Number A1 A2 A3	Temperature °C 25.6 26.76666667 27.4	Ammonia by AQ2 SEAL mg/L NH3-N 18.27 18.26 16.91	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65 22.50	Nitrate by IC mg/L NO3-N 13.14 13.56 15.31	SEÁL	SEAL mg/L NO2-N 1.15 1.49 1.70	SEÁL	SEAL mg/I P 6.92 6.53 6.63	+ Nitrite SEAL mg/L NOx-N 14.29 15.05 17.02
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser litrogen Source piked mmonia mmonia mmonia mmonia	mond Activated Sl Date dd-mmm-yy 29-Jul-08 29-Jul-08 29-Jul-08	udge/VIP SE + Sm Sample Time h:mm 9:16 9:51 10:27 11:07	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70 106 146	Sample Number A1 A2 A3 A4	Temperature °C 25.6 26.76666667 27.4 27.9	Ammonia by AQ2 SEAL mg/L NH3-N 18.27 18.26 16.91 16.95	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65 22.50 22.83	Nitrate by IC mg/L NO3-N 13.14 13.56 15.31 16.42	SEÁL	SEAL mg/L NO2-N 1.15 1.49 1.70 0.35	SEÁL	SEAL mg/I P 6.92 6.53 6.63 7.21	+ Nitrite SEAL mg/L NOx-N 14.29 15.05 17.02 16.78
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser itrogen Source piked mmonia mmonia mmonia mmonia	mond Activated Sl Date dd-mmm-yy 29-Jul-08 29-Jul-08	udge/VIP SE + Sm Sample Time h:mm 9:16 9:51 10:27	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70 106	Sample Number A1 A2 A3	Temperature °C 25.6 26.76666667 27.4	Ammonia by AQ2 SEAL mg/L NH3-N 18.27 18.26 16.91	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65 22.50	Nitrate by IC mg/L NO3-N 13.14 13.56 15.31	SEÁL	SEAL mg/L NO2-N 1.15 1.49 1.70	SEÁL	SEAL mg/I P 6.92 6.53 6.63	+ Nitrite SEAL mg/L NOx-N 14.29 15.05 17.02
e. LabView so f. 5 samples v i. Reactors A a eactor A: Nanser itrogen Source piked mmonia mmonia mmonia mmonia mmonia mmonia mmonia	mond Activated Sl Date dd-mmm-yy 29-Jul-08 29-Jul-08 29-Jul-08 29-Jul-08	udge/VIP SE + Sm Sample Time h:mm 9:16 9:51 10:27 11:07	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70 106 146 183	Sample Number A1 A2 A3 A4 A5	Temperature °C 25.6 26.76666667 27.4 27.9 28.2	Ammonia by AQ2 SEAL mg/L NH3-N 18.27 18.26 16.91 16.95 15.47	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65 22.50 22.83 23.60	Nitrate by IC mg/L NO3-N 13.14 13.56 15.31 16.42 18.01	SEÁL	SEÁL mg/L NO2-N 1.15 1.49 1.70 0.35 0.16	SEÁL	SEAL mg/l P 6-92 6-53 6-63 7-24 8-62	+ Nitrite SEAL mg/L NOx-N 14.29 15.05 17.02 16.78 18.17
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e. LabView so f. 5 samples v i. Reactors A a teactor A: Nanser teactor A: Nanser titrogen Source tipiked mmonia mmonia mmonia mmonia mmonia titrite litrite	mond Activated SI Date dd-mmm-yy 29-Jul-08 29-Jul-08 29-Jul-08 29-Jul-08 29-Jul-08 29-Jul-08	er a period of 3 ho porated industrial udge/VIP SE + Sm Sample Time h:mm 9:16 9:51 10:27 11:07 11:44 17:29 18:08	pH recording as w urs through sampl waste addition we ithfield Time since initial spike min 35 70 106 146 183 42 81	Sample Number A1 A2 A3 A4 A5	Temperature °C 25.6 26.76666667 27.4 27.9 28.1 28.2	Ammonia by AQ2 SEAL mg/L NH3-N 18.26 16.91 16.95 15.47	Ammonia by HACH TNT mg/L NH3-N 25.10 22.65 22.50 22.83 23.60	Nitrate by IC mg/L NO3-N 13.14 13.56 15.31 16.42 18.01 17.79 19.69	SEÁL	SEAL mg/L NO2-N 1.15 1.49 1.70 0.35 0.16 21.55 19.65	SEÁL	SEAL mg/l P 6-92 6-53 6-63 7-21 8-62 12-17 12-00	+ Nitrite SEAL mg/L NOx-N 14.29 15.05 17.02 16.78 18.17 39.34 39.34









Sample Data Report for Nansemond Nitrification Inhibition Study Bio-P Experimentation: I. Bio - P a. Spiked four 3L reactors with 5 mg/L PO_a-P to raise initial PO_a-concentration to roughly 15 mg/L PO_a-P. (Initial PO_a- concentration was determined through HACH TNT PO₄- Tubes at the end of the AOB/NOB experimentation). b. The Bio-P experiment was run through an uptake/release/uptake/release/uptake method were the four reactors are aerobic/an aerobic/aerobic once more.During the first release phase 100 mg/L of NaAc was added for COD manually to all 4 reactors and 200 mg/L of NaAc during the second release phase. c. Each reactor is running continously by use of stir bars. d. 2 L of the diluent source is added to the reactors. e. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 8.5 L. f. Constant DO and pH were monitored and logged throughout the experiment. g. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves during the uptake phases and then these valves were turned off and the oxygen/air was cut-off and then the system was deaerated and the DO was allowed to drop to 0 before beginning the release phase. During the release phase nitrogen was sparged into the reactors. h. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves. i. 2 samples were collected over a period of 2 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 µm filters during the second uptake phase and 5 samples were collected over a period of 1 and 1.5 hours during the second release phase and final uptake phase. The first uptake and release phases were used to eliminate any residual ammonia and nitrite. j. Reactors A and C which incorporated industrial waste addition were spiked with 200 mL of Smithfield sample (1:10 dilution). Reactor A: Nansemond Activated Sludge/VIP SE + Smithfield Time since initial Phosphate by AQ2 Ammonia by AQ2 NOx-N by AQ2 Nitrite by AQ2 NOx-N by Nitrate IC Sample Time Sample Number SEAL SEAL Nitrate by IC SEAL SEAL + Nitrite SEAL Phosphorus Test Date spike Temperature Phosphate by TNT SEAL dd-mmm-yy mg/l P mg/L NH3-N mg/L NO3-N mg/L NOx-N mg/L NO2-N mg/L NO3-N mg/L NOx-N h:mm min mg/L P Uptake 1 30-Jul-08 13:35 6 A1 28.4 55.50 40.40 0.00 4.42 9.88 14.30 Uptake 1 30-Jul-08 15:29 120 A2 28.4 37.00 24.59 0.00 11.63 2.05 13.68 Release 30-Jul-08 16:13 164 A5 28.60 37.00 24.50 0.00 5.68 2.45 8.13 Release 30-Jul-08 16:22 173 A6 28.6 44.00 29.60 0.00 4.84 5.03 9.86 Release 30-Jul-08 16:34 185 Α7 28.5 49.00 22.93 0.00 4.46 4.91 9.37 Release 30-Jul-08 16:49 200 Α8 28.5 55.00 20.06 0.00 4.77 1.54 6.31 3.11 Release 30-Jul-08 17:09 210 A9 28.6 60.00 25.13 0.00 5.66 8.76

27.33

31.23

30.45

31.90

33.15

0.00

0.00

0.00

0.00

1.57

3.62

1.79

3.05

7.32

8.17

5.44

6.52

5.72

5.15

4.37

9.07

8.32

8.77

12.47

12.54

62.00

61.00

63.00

63.00

63.00

Uptake 2

Uptake 2

Uptake 2

Uptake 2

Uptake 3

30-Jul-08

30-Jul-08

30-Jul-08

30-Jul-08

30-Jul-08

17:34

17:53

18:08

18:28

18:43

235

254

269

289

304

A10

A11

A12

A13

A14

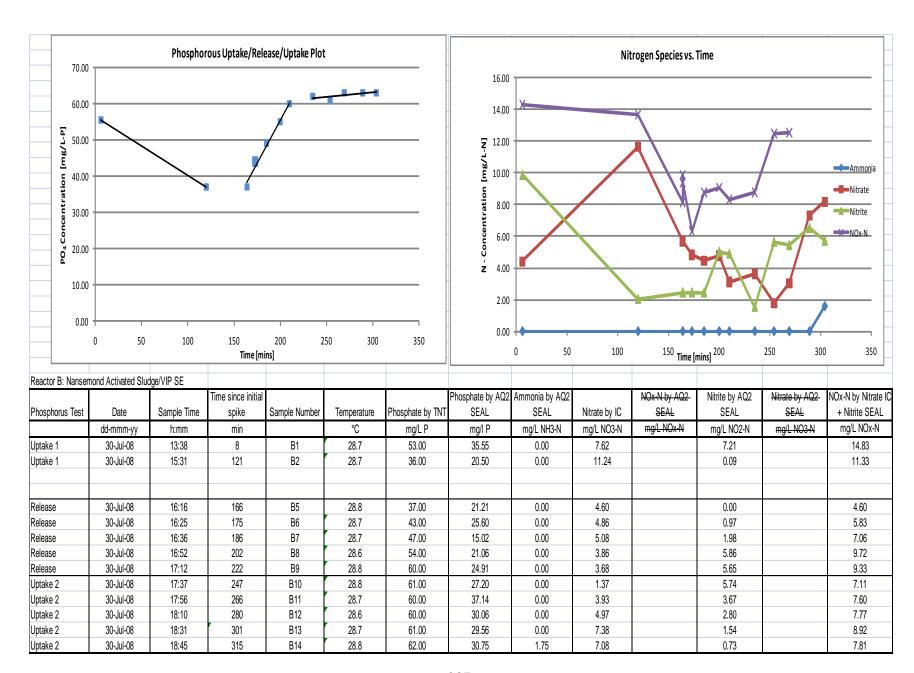
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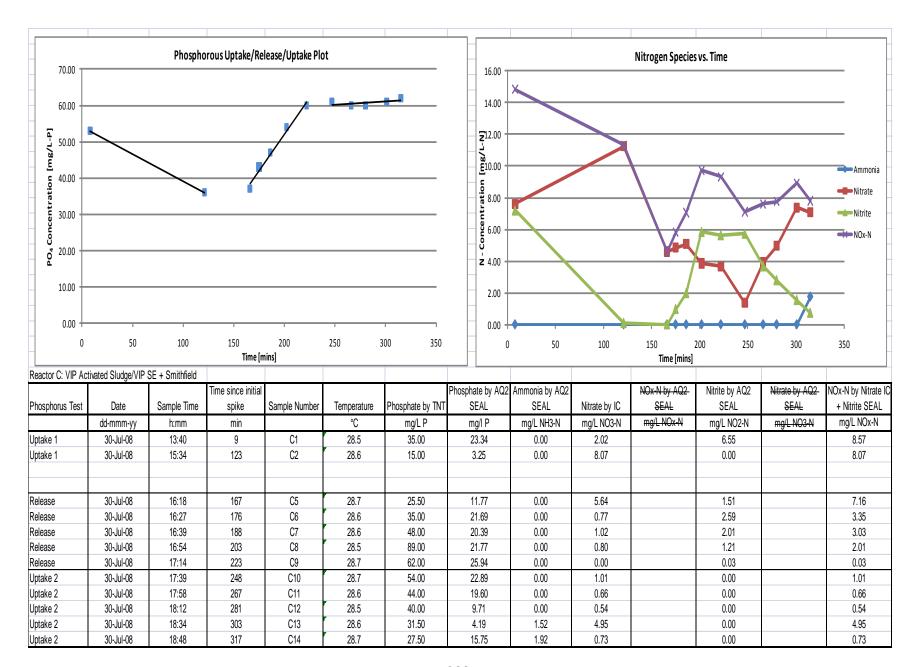
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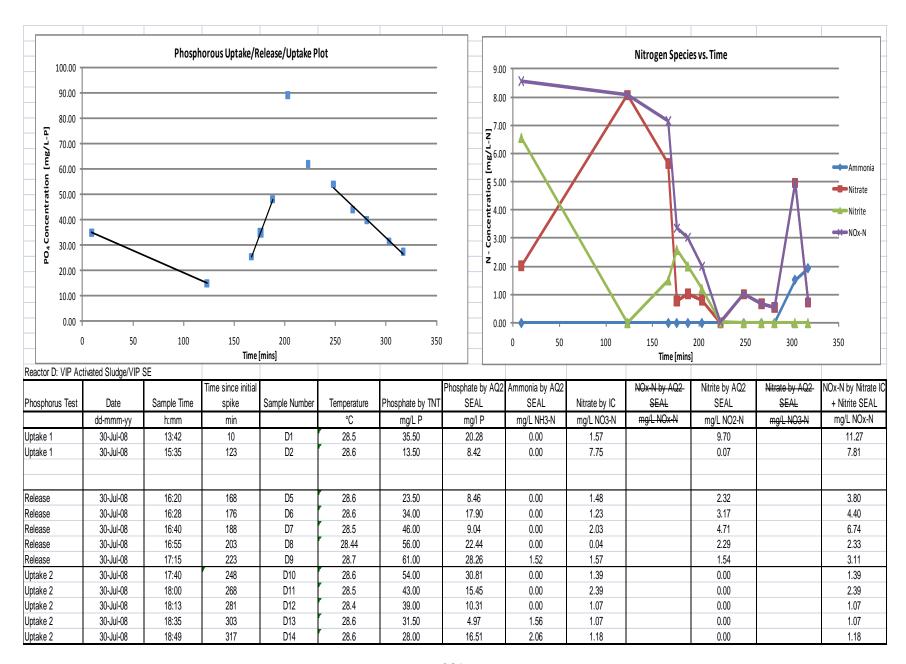
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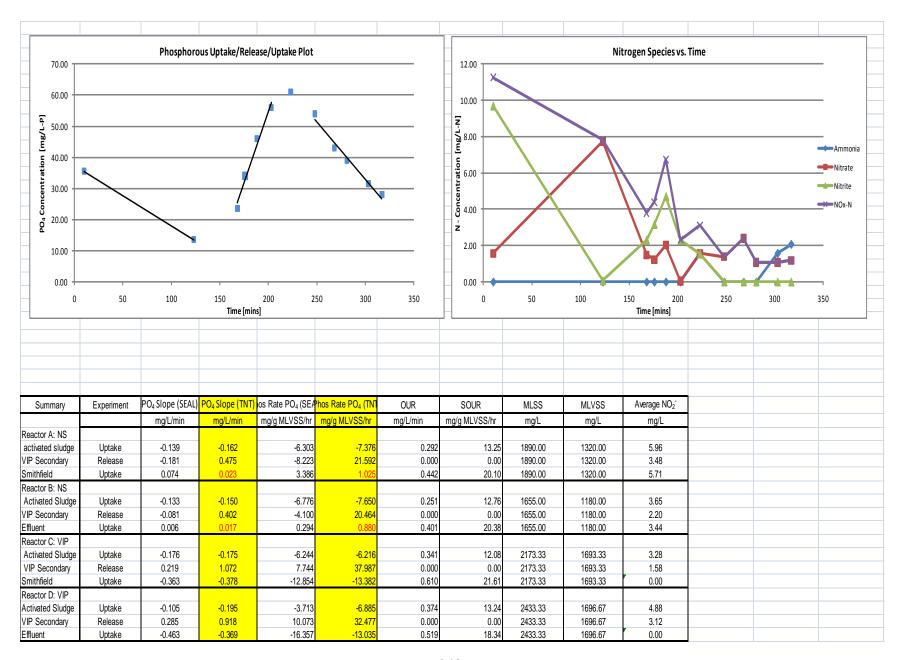
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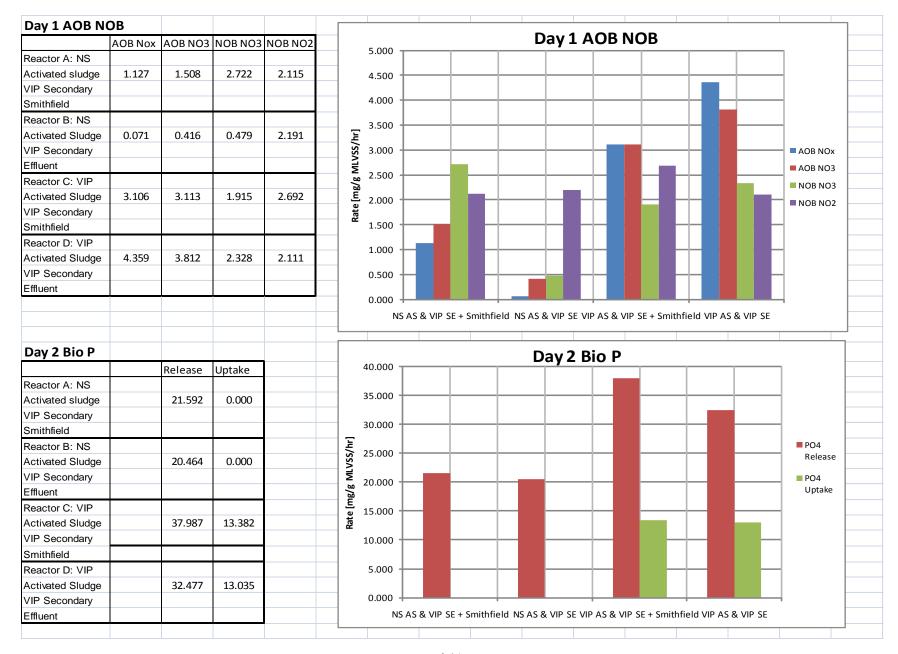
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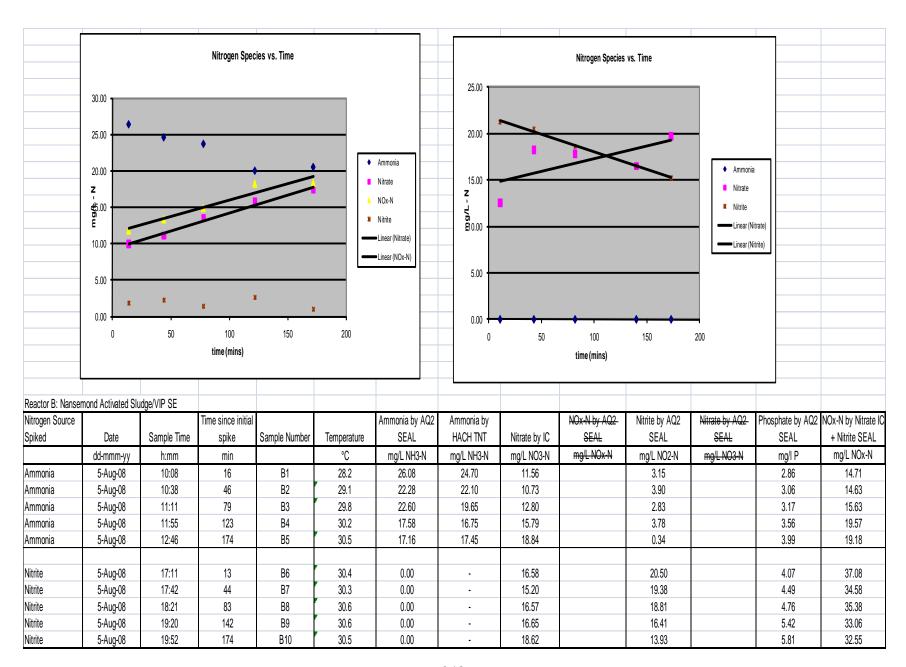


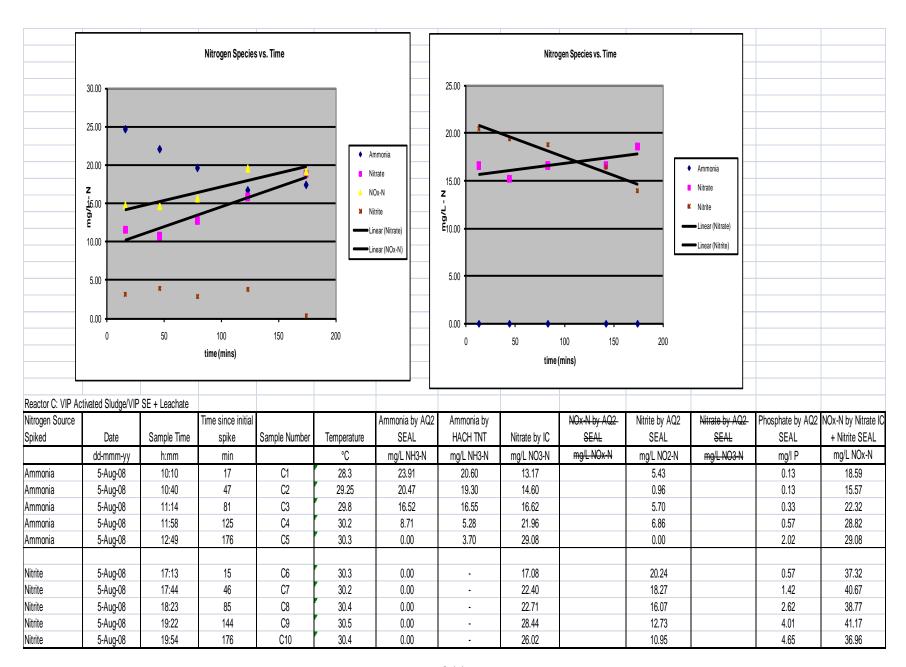


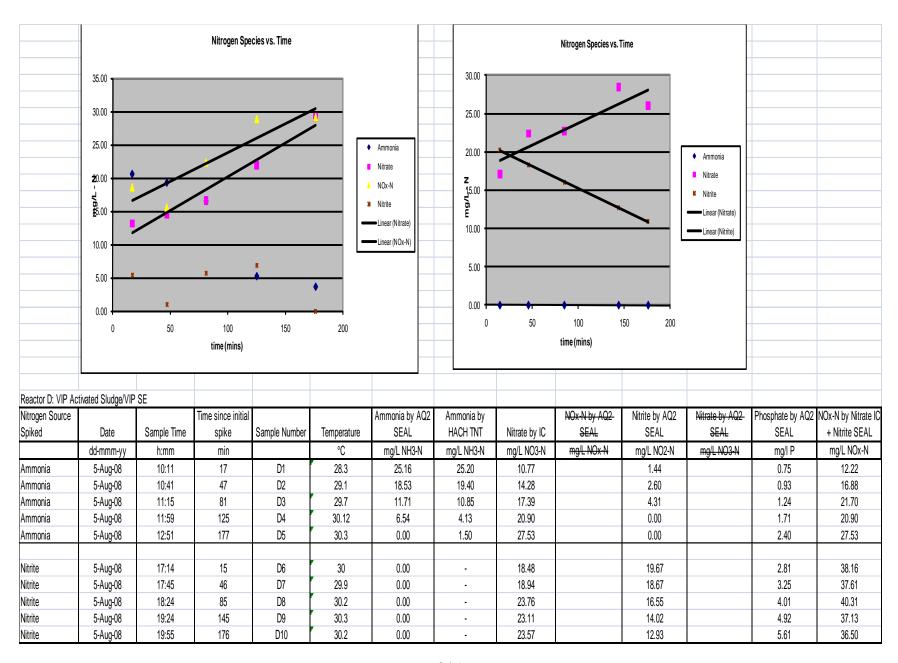
Week 6 – Landfill Leachate Day 1 AOB/NOB

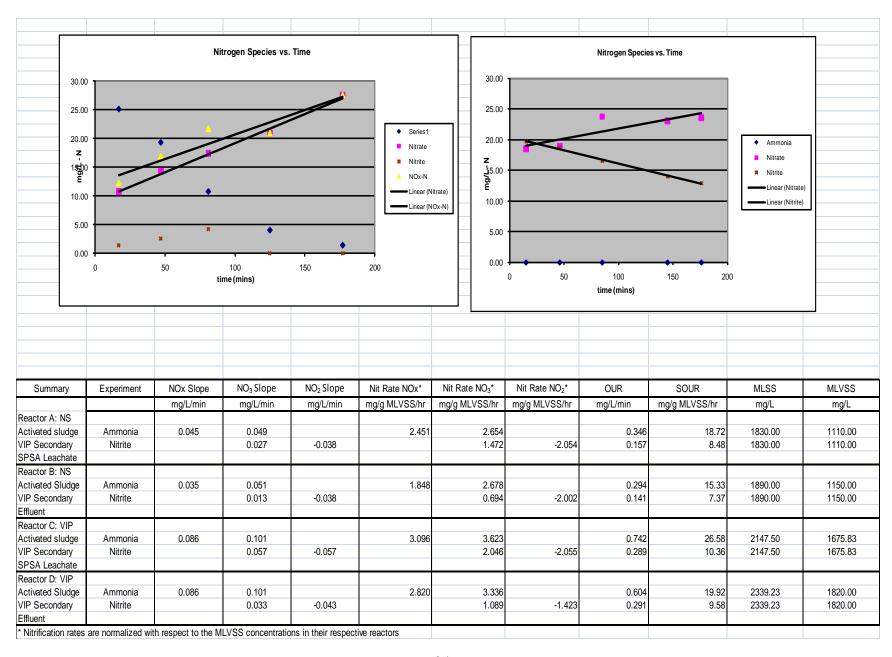
Sample Data R	Report for Nanse	mond Nitrificati	on Inhibition Stu	ıdy									
AOB & NOB Exp	perimentation:												
I. AOB	* 21	00 ma/L NIII											
	r 3L reactors with 3 or is running conti	O. 4	tir bara										
	or is running conti liluent source is ad	, ,											
			actors. For this we	aks avnarimants t									
			ed throughout the		ile biolilass was col	ncentrateu nom 20	L 10 9 L.						
			s to maintain DO le		noid valves								
,,,,	1 0			U	d implement the so	olenoid valves.							
		•				l into sample tubes t	hrough millipore 0.	45 um					
filters				01		, , , , , , , , , , , , , , , , , , , ,							
i. Reactors A	and C which incor	porated leachate	addition were spik	ed with 15 mL of t	he SPSA leachate (1:135 dilution) base	d on plant flow.						
II. NOB													
a. Spiked fou	r 3L reactors with 2	25 mg/L NO ₂ afte	r ammonia levels v	were < 1 mg/L NH ₃ -	N which was check	ed through the use	of HACH TNT 831						
				rnight to consume	all present ammor	nia.							
	or is running conti												
			ed throughout the	•									
			s to maintain DO le	-									
					d implement the so								
	were collected ove	era period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	into sample tubes t	hrough millipore 0.	45 μm					
filters.			. 1.192	. I blac	L. CDCA L L /	4.435 (2.12)	1 1 6 .						
g. Reactors A	and C wnich incor	porated leachate	addition were spik	kea with 15 mL of 1	ne SPSA leachate (1:135 dilution) base	d on plant flow.						
	11.00												
Reactor A: Nanse	mond Activated SIL	idge/VIP SE + Lea	cnate										
litrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	
piked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N
Ammonia	5-Aug-08	10:05	14	A1	28.1	26.03	26.40	9.96		1.83		2.24	11.79

Nitrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N
Ammonia	5-Aug-08	10:05	14	A1	28.1	26.03	26.40	9.96		1.83		2.24	11.79
Ammonia	5-Aug-08	10:35	44	A2	29	25.77	24.60	11.10		2.24		2.00	13.34
Ammonia	5-Aug-08	11:09	78	A3	29.7	23.09	23.70	13.50		1.40		2.14	14.90
Ammonia	5-Aug-08	11:53	122	A4	30.1	21.08	20.00	15.71		2.62		2.40	18.32
Ammonia	5-Aug-08	12:43	172	A5	30.4	18.91	20.50	17.41		0.97		2.71	18.38
Nitrite	5-Aug-08	17:08	11	A6	30.2	0.00	-	12.56		21.24		2.87	33.80
Nitrite	5-Aug-08	17:40	43	A7	30.2	0.00	-	18.28		20.49		3.68	38.77
Nitrite	5-Aug-08	18:19	82	A8	30.4	0.00	-	17.86		18.62		4.06	36.48
Nitrite	5-Aug-08	19:17	140	A9	30.5	0.00	-	16.52		16.51		4.90	33.04
Nitrite	5-Aug-08	19:50	173	A10	30.4	0.00	-	19.74		15.25		5.09	34.99



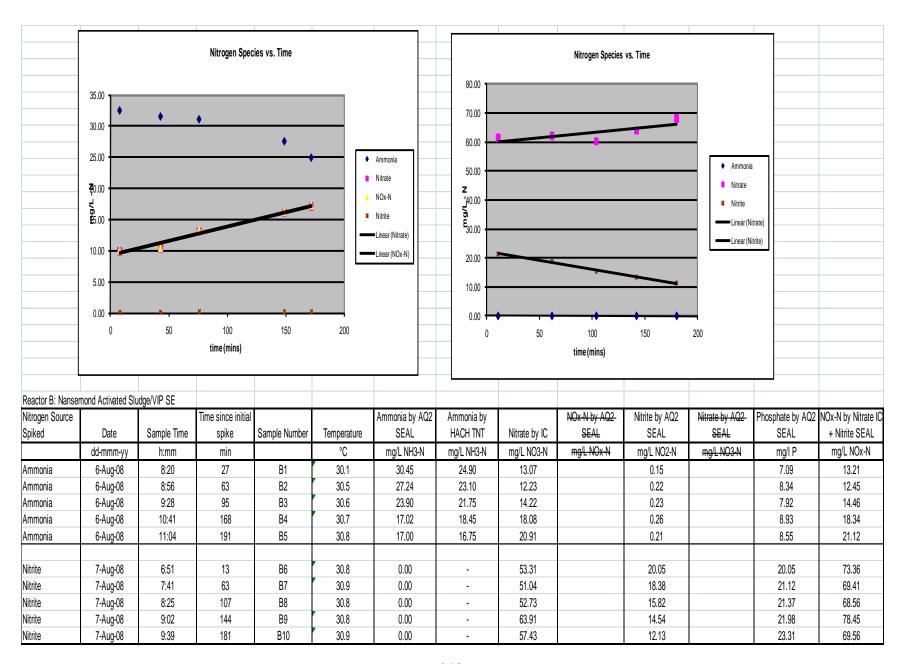


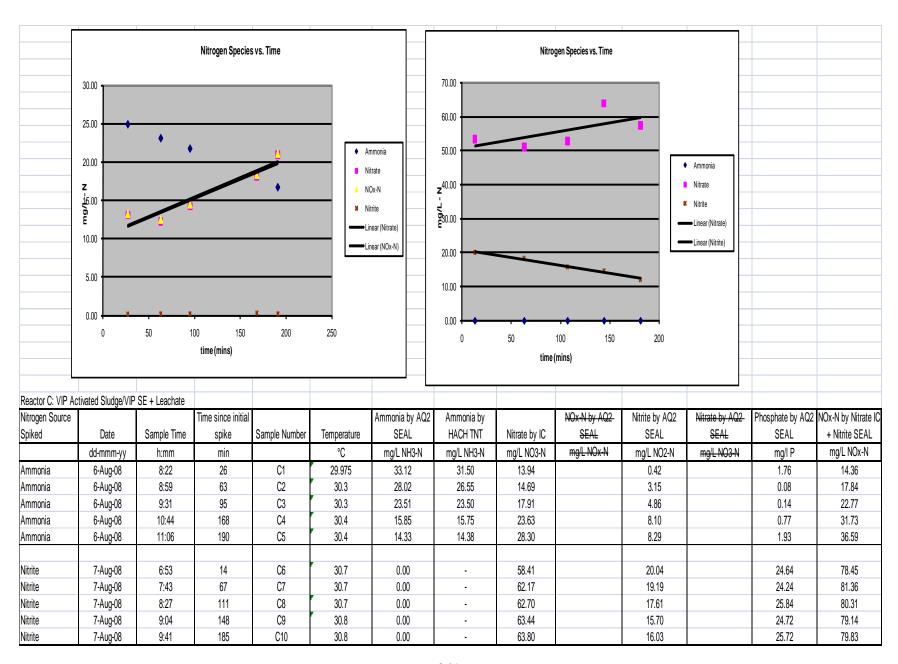


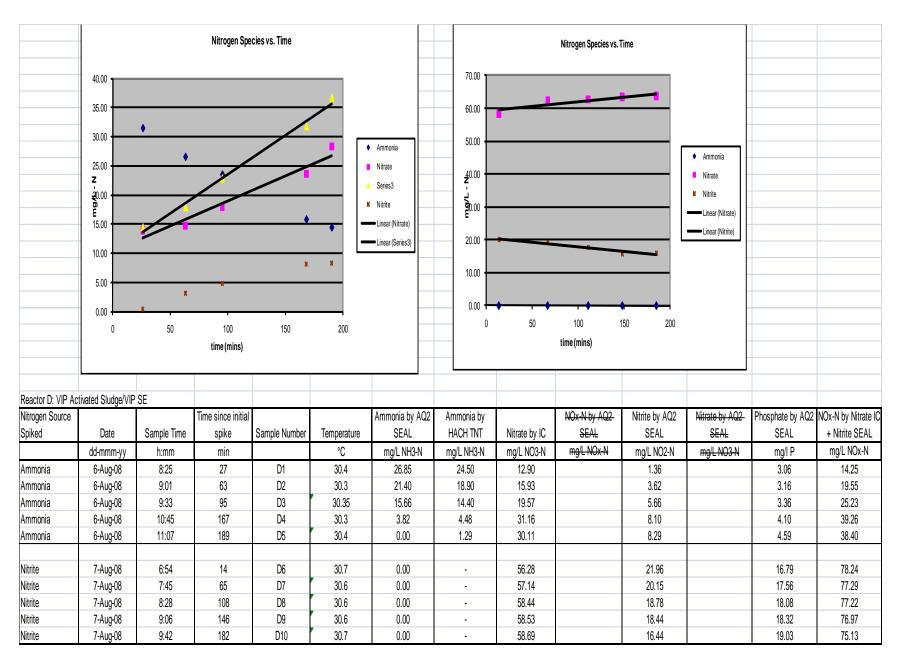


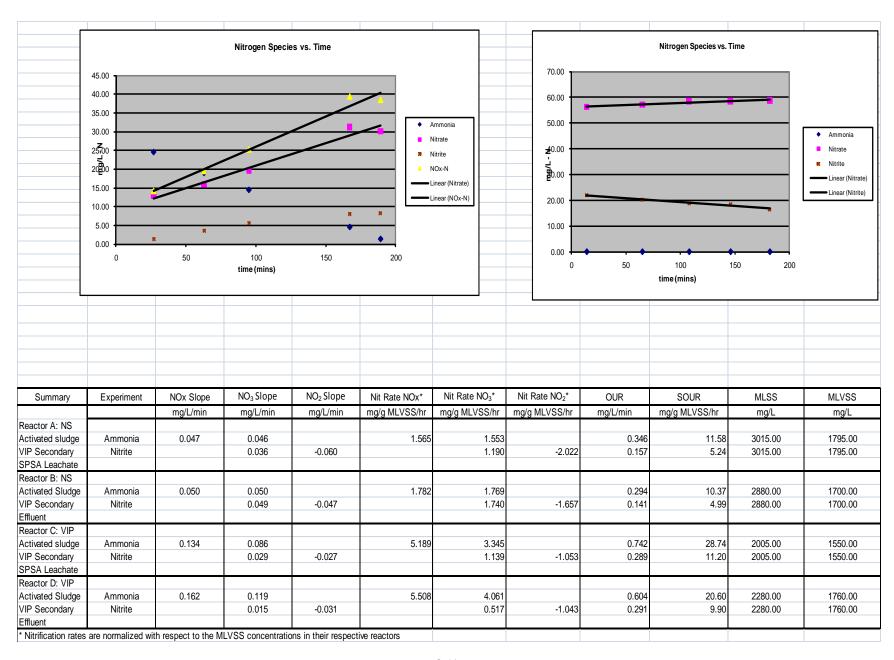
Week 6 – Landfill Leachate Day 2 AOB/NOB

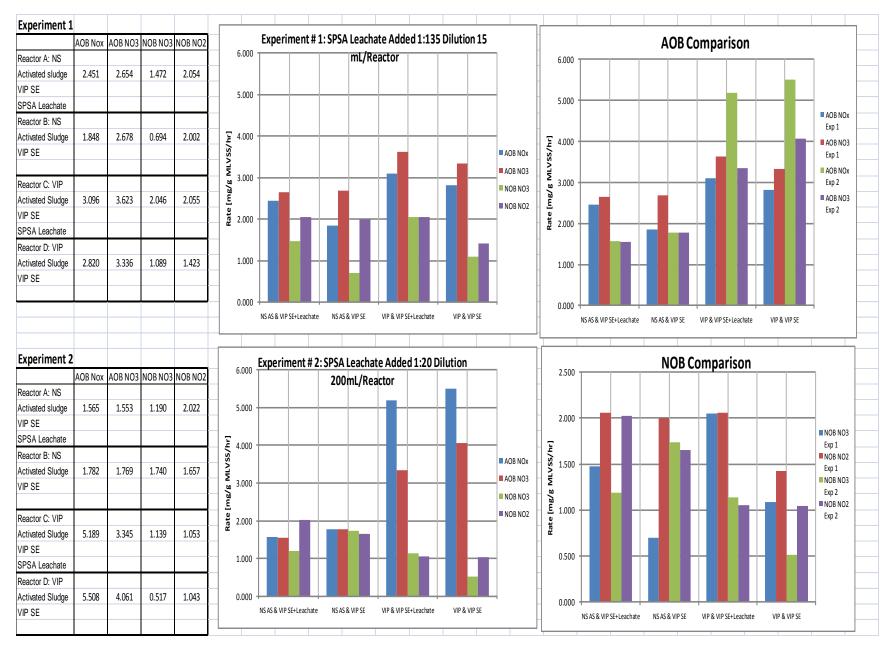
Sample Data R	Report for Nanse	emond Nitrificati	on Inhibition Stu	ıdy									
AOB & NOB Exp	perimentation:												
I. AOB	,												
a. Spiked four	r 3L reactors with	30 mg/L NH ₄ .											
'		inously by use of s	tir bars .										
	-	ce is added to the											
				eks experiments t	he biomass was co	ncentrated from 20	L to 9 L.						
		onitored and logg											
			-	evels through sole	noid valves.								
					d implement the so	olenoid valves.							
						d into sample tubes t	hrough millipore 0.	45 um filters					
						n) to simulate a slug							
						,							
II. NOB													
	r 3L reactors with	25 mg/L NO ₂ - afte	rammonia levels v	were < 1 mg/L NH ₂ -	N which was check	ked through the use	of HACH TNT 831 A	mmonia					
				isume all present a									
		inously by use of s		ioume un presente									
		onitored and logg		experiment.									
			•	evels through sole	noid valves.								
					d implement the so	olenoid valves							
							hrough millipore 0.	45 um filters					
f. 5 samples v	were collected ov	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v	were collected ov	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered			45 μm filters.					
f. 5 samples v	were collected ov	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v	were collected ov	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v	were collected ov	er a period of 3 ho	urs through sampl	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 ho rporated leachate	urs through sampl addition with 200	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 ho	urs through sampl addition with 200	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 ho rporated leachate	urs through sampl addition with 200	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 ho rporated leachate	urs through sampl addition with 200	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.	NOx-N-by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 ho rporated leachate	urs through sampl addition with 200	ling ports on the re	eactors and filtered	d into sample tubes t in) to simulate a slug	load	45 μm filters. Nitrate by IC	NOx-N by AQ2- SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2- SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL
f. 5 samples v g. Reactors A	were collected ov and C which inco	er a period of 3 hor rporated leachate udge/VIP SE + Lea	urs through sample addition with 200 chate	ling ports on the re mL of the SPSA lea	eactors and filtered echate (1:10 dilutio	d into sample tubes to in) to simulate a slug	load Ammonia by		,	•	,		
f. 5 samples v g. Reactors A	were collected ov and C which inco mond Activated SI	er a period of 3 hor rporated leachate udge/VIP SE + Lea Sample Time	urs through sample addition with 200 chate Time since initial spike	ling ports on the re mL of the SPSA lea	eactors and filtered echate (1:10 dilutio	d into sample tubes to in) to simulate a slug Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	SEÁL	SEAL	SEÁL	SEAL	+ Nitrite SEAL
f. 5 samples v g. Reactors A eactor A: Nanser itrogen Source piked	were collected ov and C which inco mond Activated SI Date dd-mmm-yy	er a period of 3 hor rporated leachate udge/VIP SE + Lea Sample Time h:mm	urs through sample addition with 200 chate Time since initial spike min	ling ports on the remL of the SPSA lea	eactors and filtered echate (1:10 dilution Temperature	d into sample tubes to in) to simulate a slug Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC mg/L NO3-N	SEÁL	SEAL mg/L NO2-N	SEÁL	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
f. 5 samples v g. Reactors A eactor A: Nanser litrogen Source piked mmonia	mond Activated SI Date dd-mmm-yy 6-Aug-08	udge/VIP SE + Lea Sample Time h:mm 8:17 8:52 9:25	chate Time since initial spike min 8 43 76	Sample Number A1 A2 A3	Temperature °C 24.95 27.7 28.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19	SEÁL	SEAL mg/l P 8.09 7.18 4.57	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32
f. 5 samples v g. Reactors A eactor A: Nanser itrogen Source piked mmonia mmonia	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38	urs through sample addition with 200 chate Time since initial spike min 8 43 76 149	Sample Number A1 A2 A3 A4	Temperature °C 24.95 27.7 28.9 30.1	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19 0.19	SEÁL	SEAL mg/I P 8.09 7.18 4.57 2.64	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27
f. 5 samples v g. Reactors A eactor A: Nanser itrogen Source piked mmonia mmonia mmonia mmonia mmonia	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08	udge/VIP SE + Lea Sample Time h:mm 8:17 8:52 9:25	chate Time since initial spike min 8 43 76	Sample Number A1 A2 A3	Temperature °C 24.95 27.7 28.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19	SEÁL	SEAL mg/l P 8.09 7.18 4.57	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32
f. 5 samples v g. Reactors A eactor A: Nanser itrogen Source piked mmonia mmonia mmonia mmonia mmonia mmonia	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38 11:01	chate Time since initial spike min 8 43 76 149 172	Sample Number A1 A2 A3 A4 A5	Temperature °C 24.95 27.7 28.9 30.1 30.3	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58 26.54	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07 17.13	SEÁL	SEÁL mg/L NO2-N 0.13 0.12 0.19 0.19 0.17	SEÁL	SEAL mg/l P 8.09 7.18 4.57 2.64 2.32	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27 17.30
f. 5 samples v g. Reactors A eactor A: Nanser itrogen Source piked mmonia mmonia mmonia mmonia mmonia mmonia itrite	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08 7-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38 11:01	rist through sample addition with 200 addition w	Sample Number A1 A2 A3 A4 A5	Temperature °C 24.95 27.7 28.9 30.1 30.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58 26.54	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07 17.13	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19 0.19 0.17	SEÁL	SEAL mg/l P 8.09 7.18 4.57 2.64 2.32	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27 17.30
f. 5 samples v g. Reactors A teactor A: Nanser litrogen Source piked mmonia mmonia mmonia mmonia litrite litrite	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08 7-Aug-08 7-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38 11:01 6:48 7:39	rist through sample addition with 200 addition w	Sample Number A1 A2 A3 A4 A5	Temperature °C 24.95 27.7 28.9 30.1 30.9 30.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58 26.54	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60 25.00	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07 17.13	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19 0.19 0.17 21.37 18.74	SEÁL	SEAL mg/l P 8.09 7.18 4.57 2.64 2.32 12.75 12.48	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27 17.30 82.82 80.75
f. 5 samples v g. Reactors A Reactor A: Nanser Ilitrogen Source piked Immonia Immonia Immonia Immonia Ilitrite Ilitrite Ilitrite	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08 7-Aug-08 7-Aug-08 7-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38 11:01 6:48 7:39 8:21	Time since initial spike min 8 43 76 149 172 11 62 104	Sample Number A1 A2 A3 A4 A5 A6 A7 A8	Temperature °C 24.95 27.7 28.9 30.1 30.9 30.9 30.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58 26.54	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60 25.00	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07 17.13 61.44 62.02 60.30	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19 0.19 0.17 21.37 18.74 15.39	SEÁL	SEAL mg/l P 8.09 7.18 4.57 2.64 2.32 12.75 12.48 14.76	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27 17.30 82.82 80.75 75.68
f. 5 samples v g. Reactors A teactor A: Nanser litrogen Source piked mmonia mmonia mmonia mmonia litrite litrite	mond Activated SI Date dd-mmm-yy 6-Aug-08 6-Aug-08 6-Aug-08 6-Aug-08 7-Aug-08 7-Aug-08	sample Time h:mm 8:17 8:52 9:25 10:38 11:01 6:48 7:39	rist through sample addition with 200 addition w	Sample Number A1 A2 A3 A4 A5	Temperature °C 24.95 27.7 28.9 30.1 30.9 30.9	Ammonia by AQ2 SEAL mg/L NH3-N 33.07 32.71 31.42 28.58 26.54	Ammonia by HACH TNT mg/L NH3-N 32.50 31.55 31.10 27.60 25.00	Nitrate by IC mg/L NO3-N 9.89 10.34 13.12 16.07 17.13	SEÁL	SEAL mg/L NO2-N 0.13 0.12 0.19 0.19 0.17 21.37 18.74	SEÁL	SEAL mg/l P 8.09 7.18 4.57 2.64 2.32 12.75 12.48	+ Nitrite SEAL mg/L NOx-N 10.02 10.46 13.32 16.27 17.30 82.82 80.75





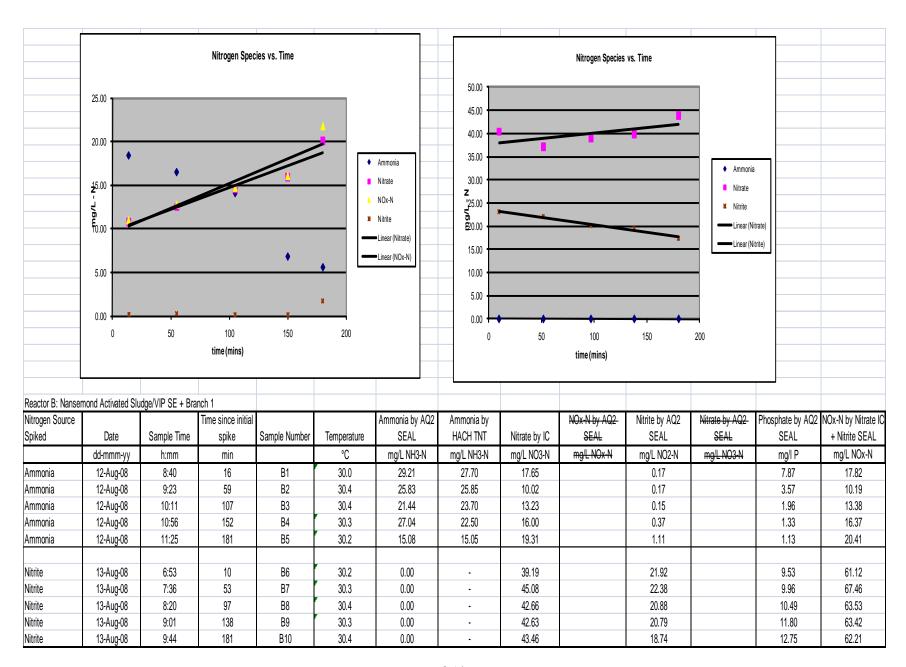


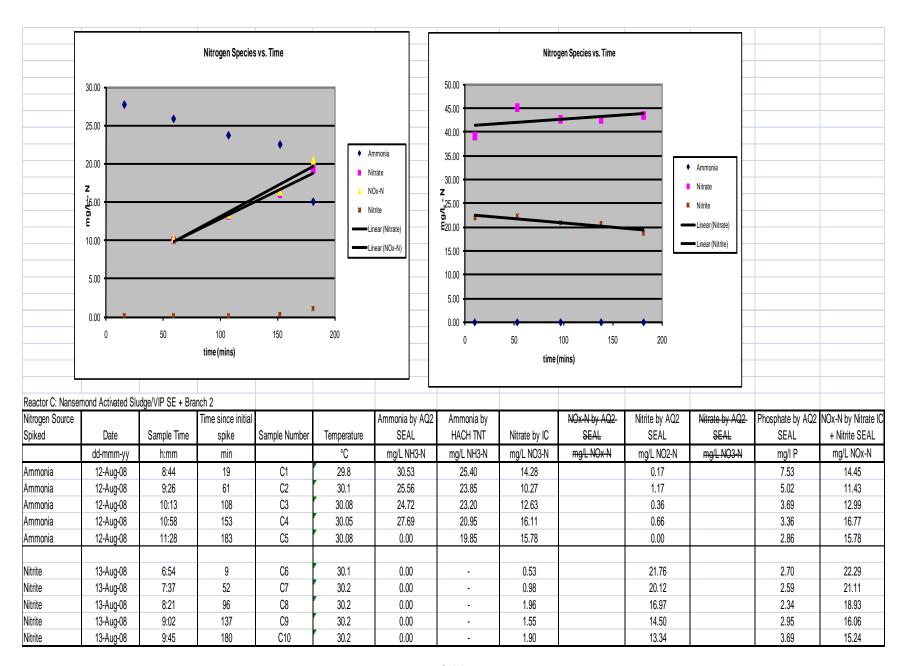


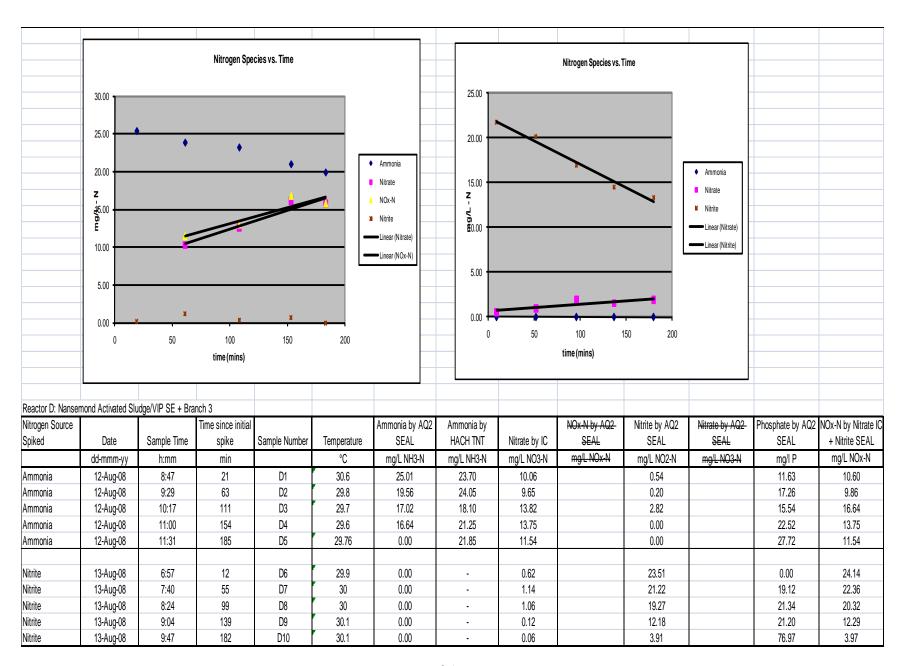


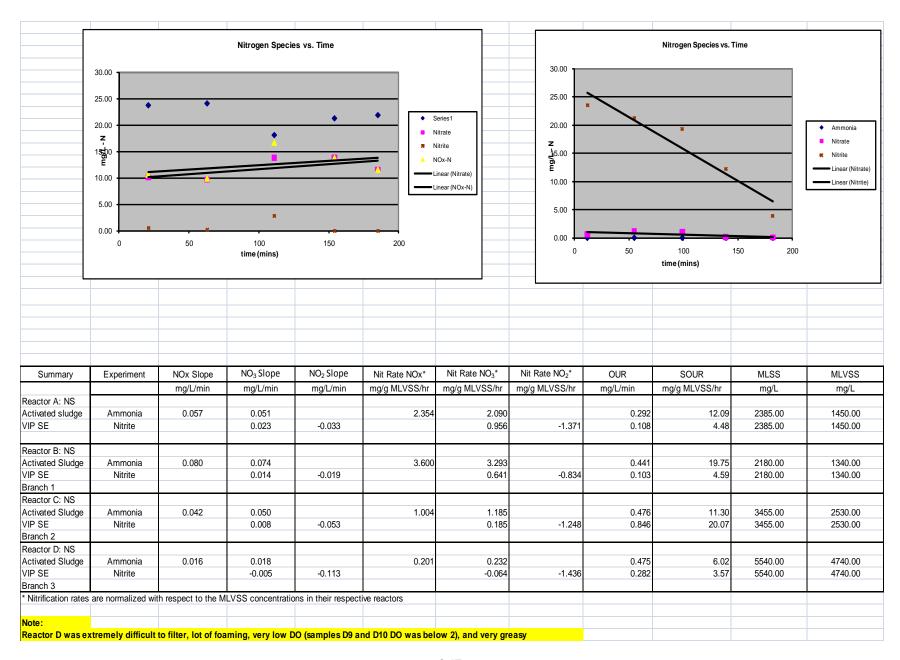
Week 7 – Branches of Collection System Day 1 AOB/NOB

Week / — Sample Data F			ion Inhibition Stu		1 AOD/I	NOD							
טמוווטוט טמומ ר	VEHOLLIOLINGLISE	THORU MUUUCAU		luy									
AOR & NOR FY	perimentation:												
I. AOB	perimentation.												
	r 3L reactors with	20 mg/L NH ₄ .											
•	or is running conti	· .	tir bars.										
	liluent source is ac												
				eks experiments t	he biomass was co	ncentrated from 201	L to 9 L.						
d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 9 L. e. Constant DO and pH were monitored and logged throughout the experiment.													
f. Oxygen/ai	ir blend was sparge	ed into the reactor	rs to maintain DO le	evels through sole	noid valves.								
g. LabView so	oftware was used	to manage DO and	d pH recording as w	vell as program and	d implement the s	olenoid valves.							
h. 5 samples	were collected ov	era period of 3 ho	ours through sampl	ling ports on the re	actors and filtered	l into sample tubes t	hrough millipore 0.	45 μm filters					
i. Reactors B	, C, and D in the di	luent source incor	rporated 1L of VIP:	SE and 1L of the as	sociated Branch lir	nes for each reactor.							
II. NOB													
		J		O . 3		ked through the use	of HACH TNT 831 A	ımmonia					
1			ın overnight to con	isume all present a	ammonia.								
	or is running conti												
			ed throughout the	•									
			s to maintain DO le										
			pH recording as w					611					
			0 1	01		l into sample tubes t	0 1	45 μm filters.					
g. Reactors B	, C, and D in the d	lluent source incor	rporated 1L of VIP	SE and 1L of the as	sociated Branch III	nes for each reactor.							
Reactor A: Nanse	mond Activated SI	Idne/VIP SF											
todotoi 7t. 14diloc	Thoras Notivated Of	augo, vii OL											
Nitrogen Source	Data	0I- T	Time since initial	O la Niilian	T	Ammonia by AQ2	Ammonia by	NEGOTA IN IO	NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2		NOx-N by Nitrate IC
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL NUO N	HACH TNT	Nitrate by IC	SEAL 	SEAL	SEAL ***** NOON	SEAL	+ Nitrite SEAL
\	dd-mmm-yy	h:mm	min	A.4	°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N
Ammonia	12-Aug-08	8:37 9:18	14 55	A1 A2	29.9 30.8	25.00 20.57	18.40	10.76 12.51		0.17		2.01 2.27	10.93 12.73
Ammonia Ammonia	12-Aug-08 12-Aug-08	10:08	105	A2 A3	30.8	19.48	16.50 14.10	12.51		0.23 0.13		2.27	12.73
Ammonia	12-Aug-08 12-Aug-08	10:53	150	A4	30.4	13.56	6.85	15.90		0.13		2.41	16.04
Ammonia	12-Aug-08	11:23	180	A5	30.2	15.14	5.63	20.07		1.70		3.07	21.77
							2.00					2.0.	
Vitrite	13-Aug-08	6:52	10	A6	30.1	0.00		40.37		23.11		14.52	63.48
Vitrite	13-Aug-08	7:34	52	A7	30.25	0.00	-	37.07		22.12		12.02	59.19
Vitrite	13-Aug-08	8:19	97	A8	30.225	0.00	•	38.95		20.25		12.45	59.20
Nitrite	13-Aug-08	9:00	138	A9	30.3	0.00	•	39.91		19.43		16.59	59.34
Nitrite	13-Aug-08	9:42	180	A10	30.3	0.00		43.89	1	17.40	1	17.08	61.29

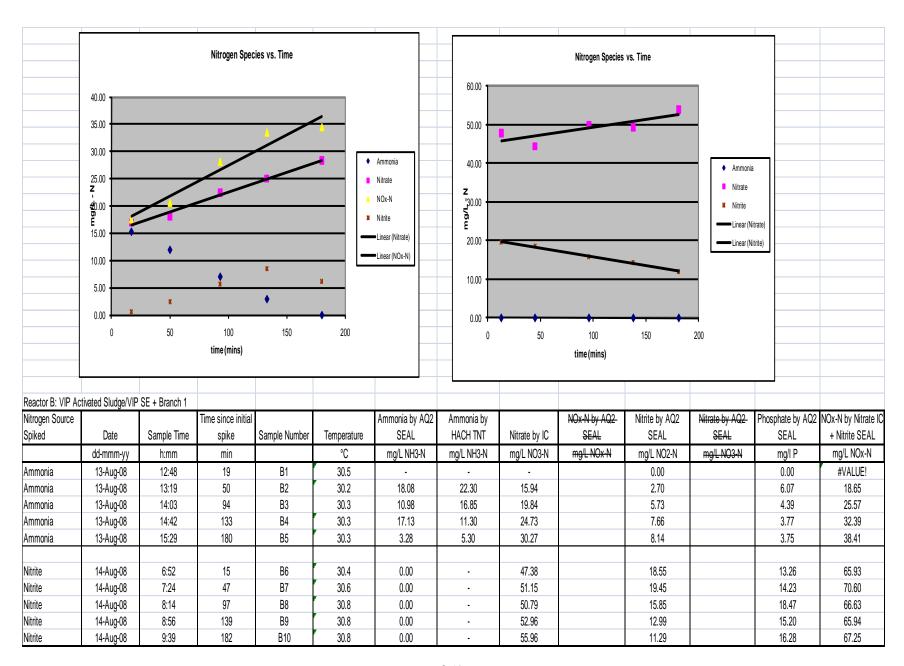


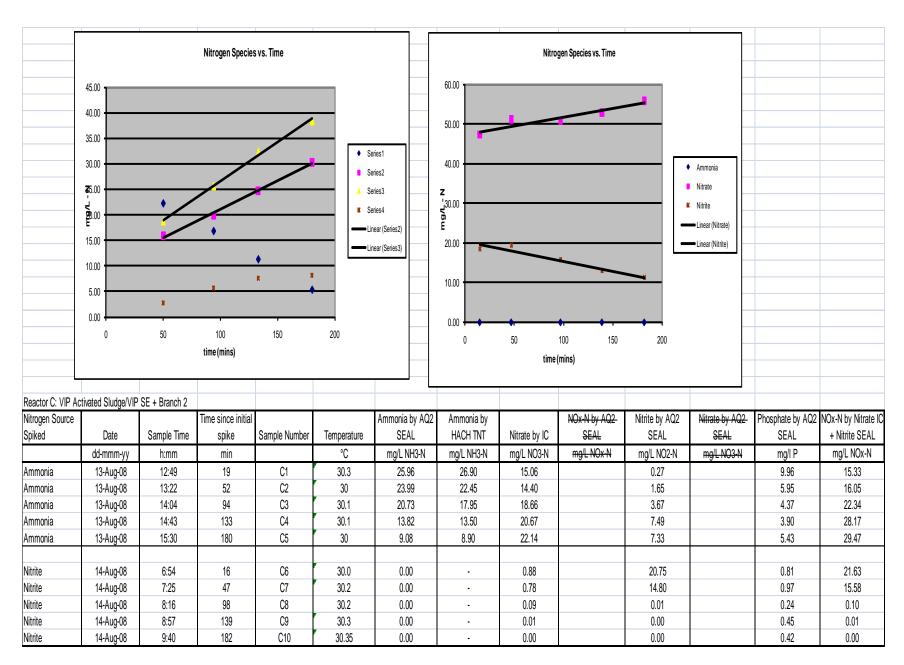


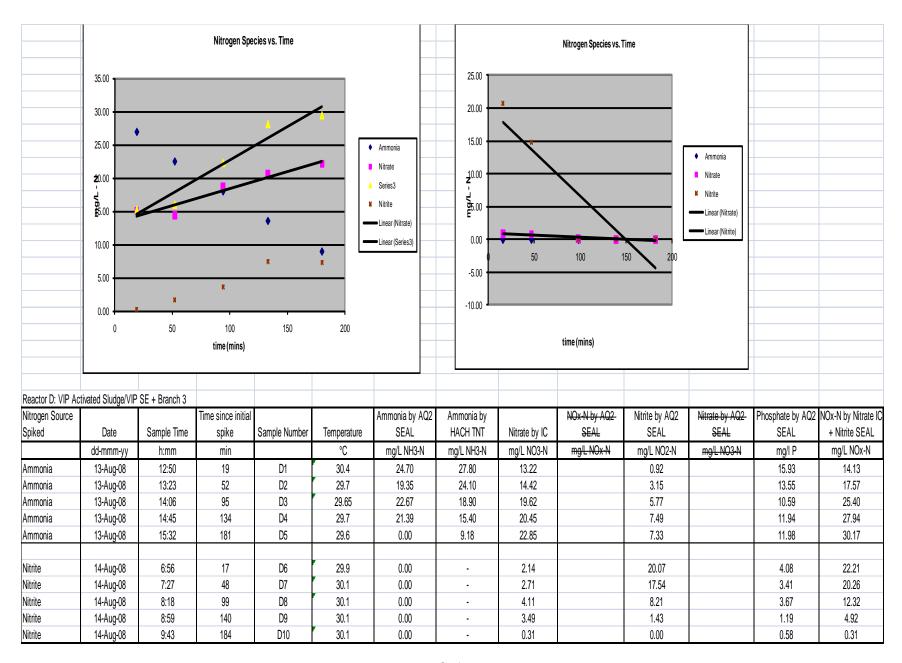


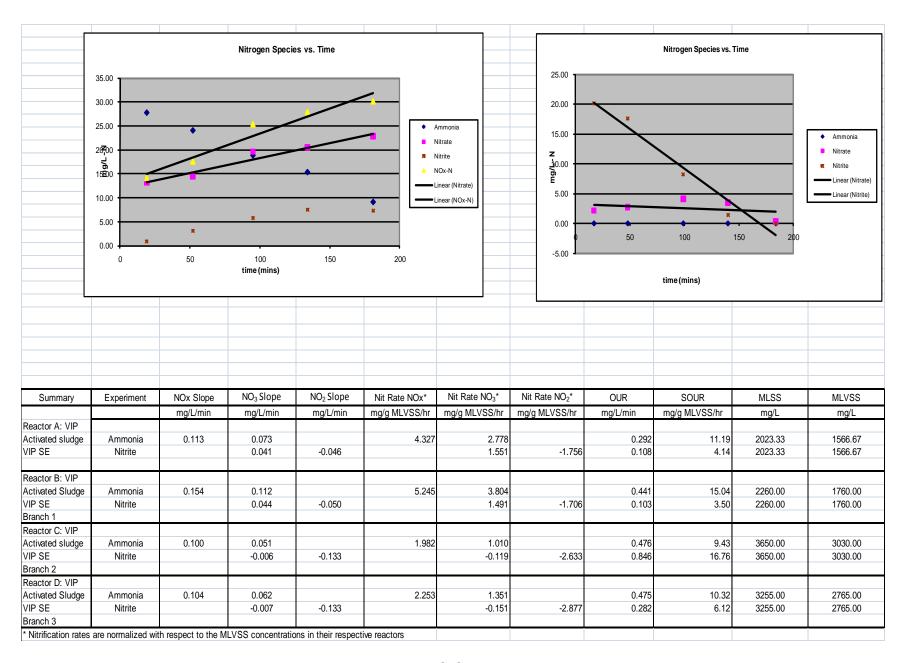


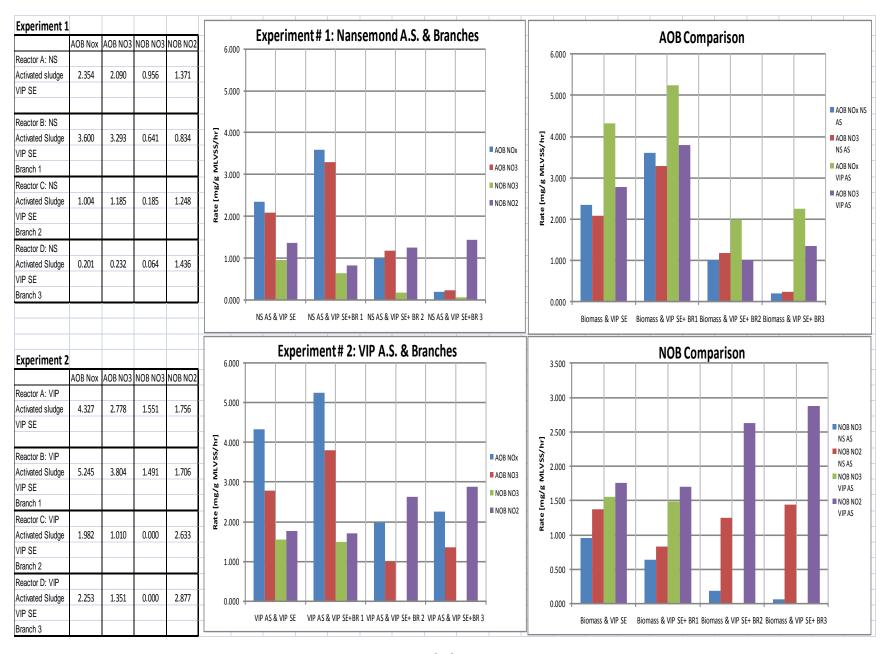
									I	I	ı		
Sample Data R	Report for Nanse	emond Nitrificat	ion Inhibition Stu	dy									
AOB & NOB Exp	perimentation:												
I. AOB													
a. Spiked fou	r 3L reactors with	20 mg/L NH ₄ .											
b. Each reacto	or is running conti	inously by use of s	tir bars.										
c. 1900 mL of	the diluent sourc	e is added to the	reactors.										
d. 1 L of conc	d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 9 L. e. Constant DO and pH were monitored and logged throughout the experiment. f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.												
e. Constant D													
f. Oxygen/ai													
			d pH recording as w			olenoid valves.							
						d into sample tubes t	hrough millipore 0	45 um filters					
						nes for each reactor.							
1	, ,		,										
II. NOB													
	r3L reactors with	25 mg/L NO ₃ - afte	erammonia levels v	vere<1mg/I NH	N which was check	ked through the use	of HACH TNT 831 A	Ammonia					
			un overnight to con			ned timodgirtine doe	011111011111111111111111111111111111111						
	or is running conti		-	sume un presente	iiiiiioiiia.								
			ed throughout the	experiment									
			rs to maintain DO le		noidvalves								
			d pH recording as w			olenoid valves							
						d into sample tubes t	hrough millingre ()	15 um filtors					
		•				nes for each reactor.		45 μπτιιτοτό.					
g. Nedetors b,	, c, and D in the di	iluciit source ilico	ipolated it of vii .	or and it of the as	30clatea Brancii III	ics for cacifficactor.							
Donator A. VID A.	ctivated Sludge/VIF	0.05											
REACTOL A. VIP AC	stivated Sludge/ vir	35											
i													
Nitrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N
Ammonia	13-Aug-08	12:45	17	A1	30.8	12.87	15.20	16.95		0.62		4.99	17.57
Ammonia	13-Aug-08	13:18	50	A2	30.7	15.69	11.90	18.14		2.47		5.04	20.61
Ammonia	13-Aug-08	14:01	93	A3	30.5	7.87	7.00	22.36		5.72		5.19	28.09
Ammonia	13-Aug-08	14:41	133	A4	30.4	1.4133ELL	2.93	25.00		8.44		5.89	33.44
Ammonia	13-Aug-08	15:28	180	A5	30.4	4.45	0.00	28.31		6.12		6.50	34.43
Nitrite	14-Aug-08	6:50	13	A6	29.9	0.00	-	47.93		19.74		15.78	67.67
Nitrite	14-Aug-08	7:22	45	A7	30.2	0.00	•	44.40		18.49		16.88	62.89
Nitrite	14-Aug-08	8:13	96	A8	30.5	0.00	•	49.99		15.92		17.71	65.91
Nitrite	14-Aug-08	8:55	138	A9	30.7	0.00	-	49.29		14.33		14.33	63.62
Nitrite	14-Aug-08	9:38	181	A10	30.7	0.00	•	53.84		12.01		18.53	65.85





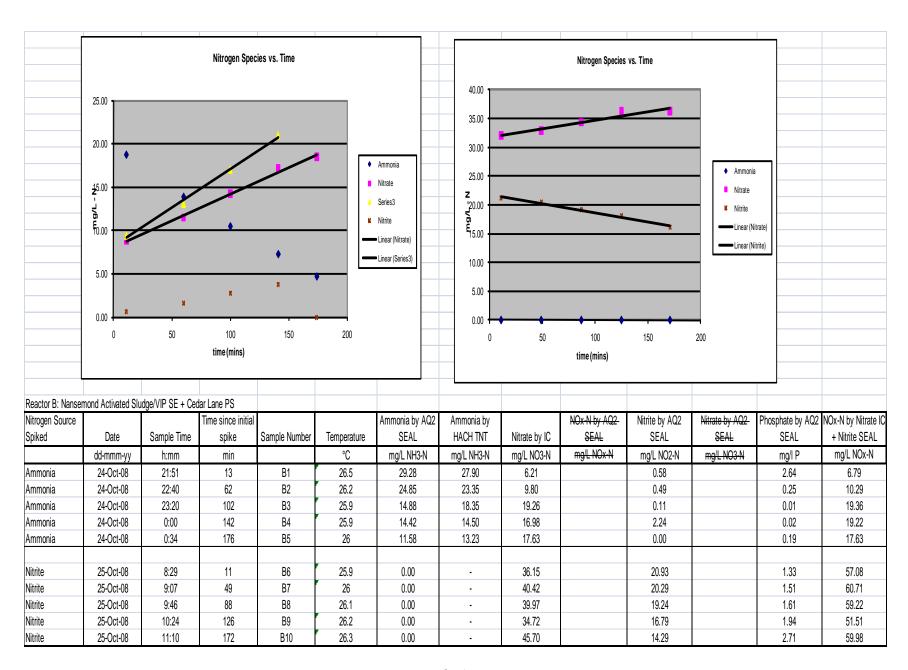


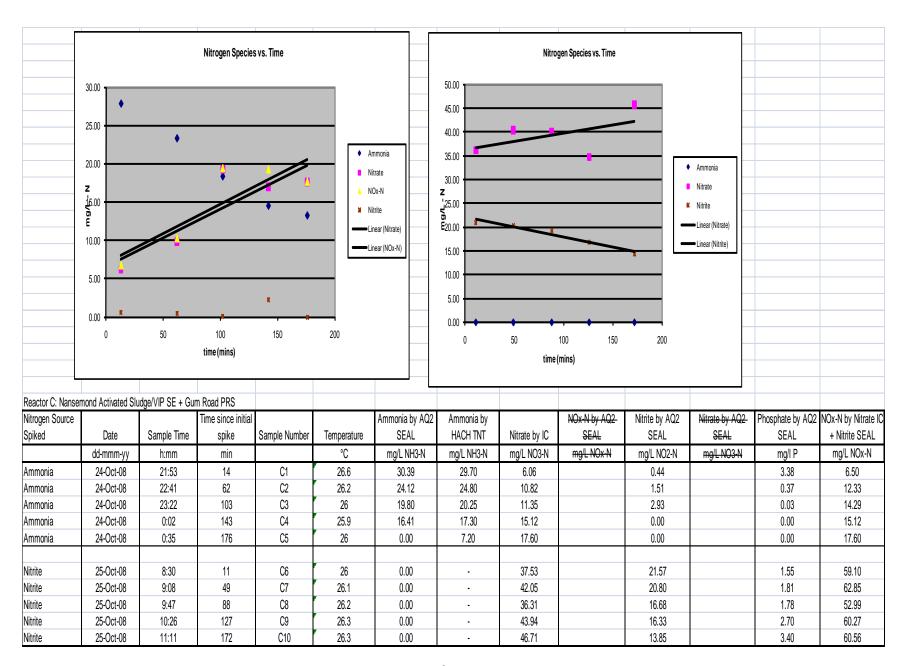


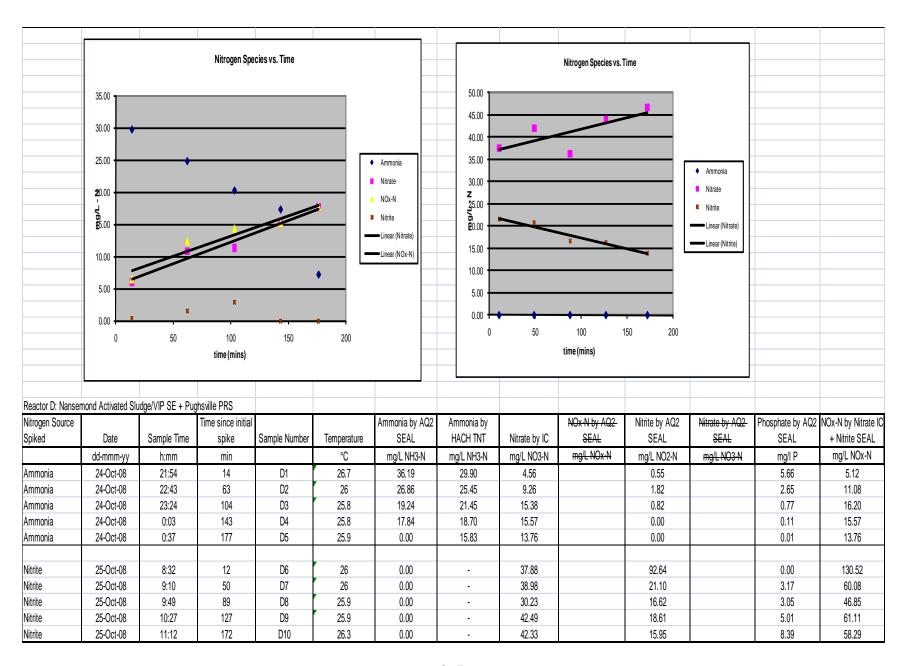


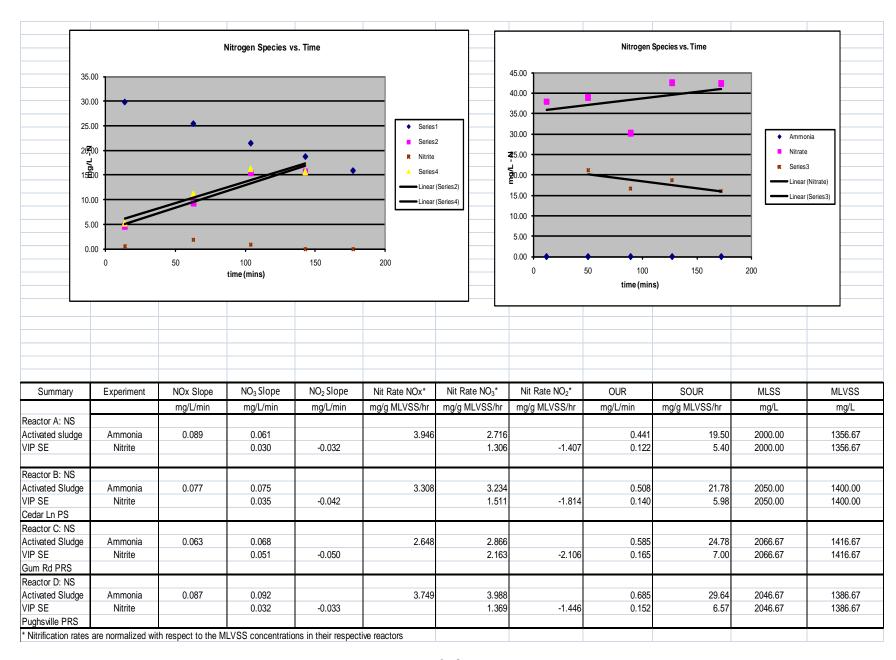
Week 8 – Pump/Pressure Reducing Stations Day 1 AOB/NOB

Sample Data r	Report for Nanse	emond Nitrificati	on Inhibition Stu	ıdy									
AOB & NOB Ex	perimentation:												
I. AOB													
a. Spiked fou	r 3L reactors with	20 mg/L NH ₄ .											
b. Each react	or is running conti	inously by use of st	tir bars.										
c. 2 L of the d	liluent source is ac	dded to the reacto	ors.										
d. 1L of conc	entrated biomass	is added to the re	actors. For this we	eks experiments t	he biomass was co	ncentrated from 201	L to 9 L.						
d. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 9 L. e. Constant DO and pH were monitored and logged throughout the experiment. f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.													
					d implement the so	olenoid valves.							
-		-			•	into sample tubes t	hrough millipore 0.	45 um filters					
						or lines for each read							
	, ,,		,		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,								
II. NOB													
	r31 reactors with	25 mg/I NO ₂ - afta	rammonia levels v	were<1mg/INH.	N which was chack	ed through the use	of HACH TNT 821 A	mmonia					
		were allowed to ru		0 . 3		ica tili ougii tile use	OTTIACIT INT 031 A	immoma					
		inously by use of st		isuille all present	aiiiiioiiia.								
				evneriment									
					noidvalvos								
c. Constant DO and pH were monitored and logged throughout the experiment. d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.													
	e. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.												
e. LabView so	oftware was used	to manage DO and	pH recording as w	vell as program and	d implement the so		٥ معمد مثالة معمل معرب معمل	AF filtore					
e. LabView so f. 5 samples	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered			45 μm filters.					
e. LabView so f. 5 samples	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView sc f. 5 samples g. Reactors B	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView sc f. 5 samples g. Reactors B	oftware was used were collected ov	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView sc f. 5 samples g. Reactors B	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through samp	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabViewsc f. 5 samples g. Reactors B,	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t or lines for each rea	ctor.	45 μm filters.	NOv.N by AO2	Nitrite by AO2	Nitrate by AO2	Phoenhate by ΔΩ2	NOv.N by Nitrate IC
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as	d implement the so eactors and filtered essociated intercept	into sample tubes to r lines for each read	Ammonia by		NOx N by AQ2-	Nitrite by AQ2	Nitrate by AQ2-	' '	,
e. LabView sc f. 5 samples g. Reactors B	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor udge/VIP SE Sample Time	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the re	d implement the so eactors and filtered essociated intercept Temperature	into sample tubes t or lines for each read Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	SEÁL	SEÁL	SEAL	SEAL	NOx-N by Nitrate IC + Nitrite SEAL
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse Nitrogen Source Spiked	oftware was used were collected ov , C, and D in the d mond Activated SI Date dd-mmm-yy	to manage DO and era period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as Sample Number	d implement the so eactors and filtered essociated intercept Temperature	into sample tubes to or lines for each read Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC	,	SEAL mg/L NO2-N	,	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse Witrogen Source Spiked	oftware was used were collected ov , C, and D in the d , and D in the d , and Activated SI Date dd-mmm-yy 24-Oct-08	to manage DO and era period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as Sample Number	d implement the signature of the sectors and filtered sectors and filter	Ammonia by AQ2 SEAL mg/L NH3-N 16.78	Ammonia by HACH TNT mg/L NH3-N 18.70	Nitrate by IC mg/L NO3-N 8.82	SEÁL	SEAL mg/L NO2-N 0.63	SEAL	SEAL mg/I P 0.27	+ Nitrite SEAL mg/L NOx-N 9.45
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse Witrogen Source Spiked	oftware was used were collected ov , C, and D in the d mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the research and 1L of the as Sample Number A1 A2	Temperature °C 27.5 26.9	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85	Nitrate by IC mg/L NO3-N 8.82 11.50	SEÁL	SEAL mg/L NO2-N 0.63 1.60	SEAL	SEAL mg/l P 0.27 0.38	+ Nitrite SEAL mg/L NOx-N 9.45 13.10
e. LabView sc f. 5 samples g. Reactors B, Reactor A: Nanse Alitrogen Source Spiked	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the research and 1L of the asset	Temperature °C 27.5 26.9 26.3	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77	SEAL	SEAL mg/l P 0.27 0.38 0.51	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03
e. LabView sc f. 5 samples g. Reactors B, Reactor A: Nanse Nitrogen Source Spiked Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18 23:59	pH recording as w urs through sample porated 1L of VIP:	vell as program and ling ports on the research and 1L of the assert and	Temperature °C 27.5 26.9 26.3 26.2	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98 5.66	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45 7.25	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26 17.17	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77 3.80	SEAL	SEAL mg/I P 0.27 0.38 0.51 0.78	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03 20.97
e. LabView sc f. 5 samples g. Reactors B, Reactor A: Nanse Alitrogen Source Spiked Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the research and 1L of the asset	Temperature °C 27.5 26.9 26.3	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77	SEAL	SEAL mg/l P 0.27 0.38 0.51	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03
e. LabView sc f. 5 samples g. Reactors B, deactor A: Nanse ditrogen Source ipiked dimmonia dimmonia dimmonia dimmonia	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08	udge/VIP SE Sample Time h:mm 21:49 22:38 23:18 23:59 0:32	pH recording as w urs through sample porated 1L of VIP:	Sample Number A1 A2 A3 A4 A5	Temperature °C 27.5 26.9 26.3 26.2	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98 5.66 2.64	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45 7.25	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26 17.17 18.46	SEÁL	SEÁL mg/L NO2-N 0.63 1.60 2.77 3.80 0.00	SEAL	SEAL mg/l P 0.27 0.38 0.51 0.78 0.97	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03 20.97 18.46
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse ditrogen Source Spiked	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18 23:59 0:32 8:28	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 11 60 100 141 174	Sample Number A1 A2 A3 A4 A5	Temperature °C 27.5 26.9 26.3 26.2 26.1	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98 5.66 2.64	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45 7.25 4.68	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26 17.17 18.46	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77 3.80 0.00	SEAL	SEAL mg/l P 0.27 0.38 0.51 0.78 0.97	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03 20.97 18.46
e. LabViewsc f. 5 samples g. Reactors B, Reactor A: Nanse Nitrogen Source Spiked Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08 25-Oct-08 25-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18 23:59 0:32 8:28 9:06	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 11 60 100 141 174	Sample Number A1 A2 A3 A4 A5 A6 A7	Temperature °C 27.5 26.9 26.2 26.1	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98 5.66 2.64	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45 7.25 4.68	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26 17.17 18.46	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77 3.80 0.00 21.23 20.48	SEAL	SEAL mg/l P 0.27 0.38 0.51 0.78 0.97	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03 20.97 18.46 53.33 53.39
e. LabViewsc f. 5 samples g. Reactors B,	mond Activated SI Date dd-mmm-yy 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08 24-Oct-08	to manage DO and er a period of 3 ho iluent source incor udge/VIP SE Sample Time h:mm 21:49 22:38 23:18 23:59 0:32 8:28	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 11 60 100 141 174	Sample Number A1 A2 A3 A4 A5	Temperature °C 27.5 26.9 26.3 26.2 26.1	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 13.04 8.98 5.66 2.64	Ammonia by HACH TNT mg/L NH3-N 18.70 13.85 10.45 7.25 4.68	Nitrate by IC mg/L NO3-N 8.82 11.50 14.26 17.17 18.46	SEÁL	SEAL mg/L NO2-N 0.63 1.60 2.77 3.80 0.00	SEAL	SEAL mg/l P 0.27 0.38 0.51 0.78 0.97	+ Nitrite SEAL mg/L NOx-N 9.45 13.10 17.03 20.97 18.46



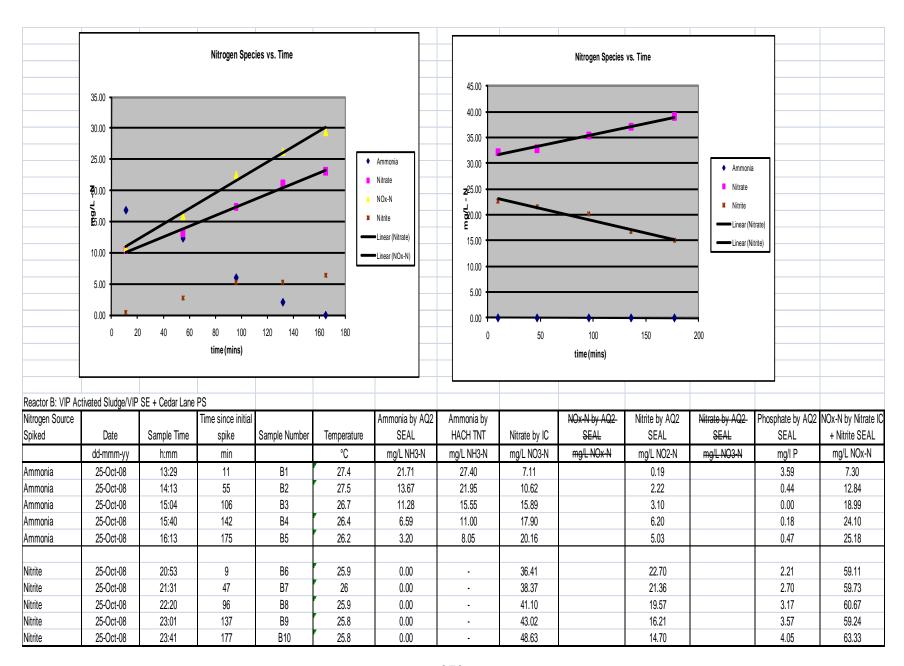


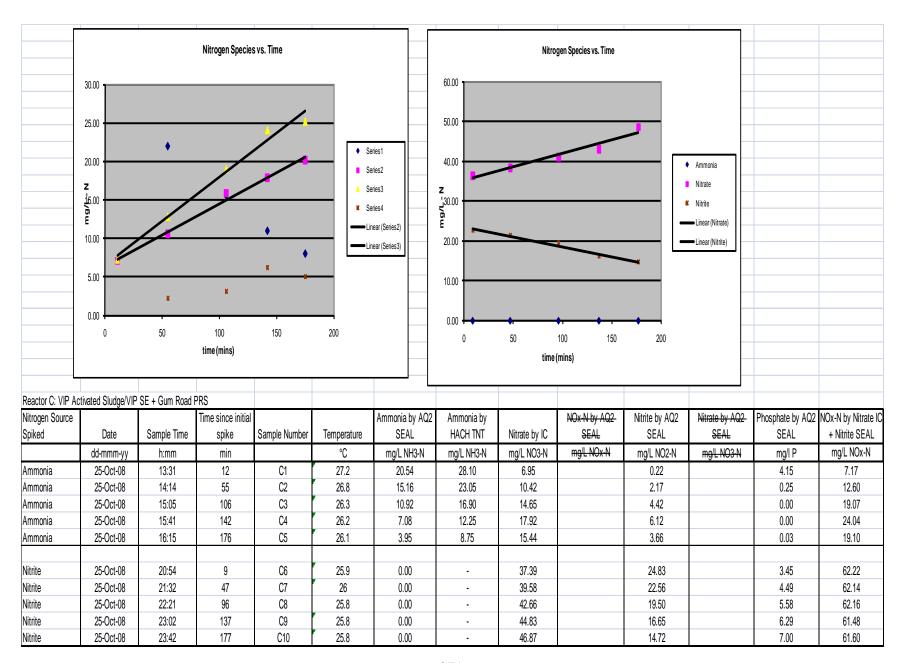


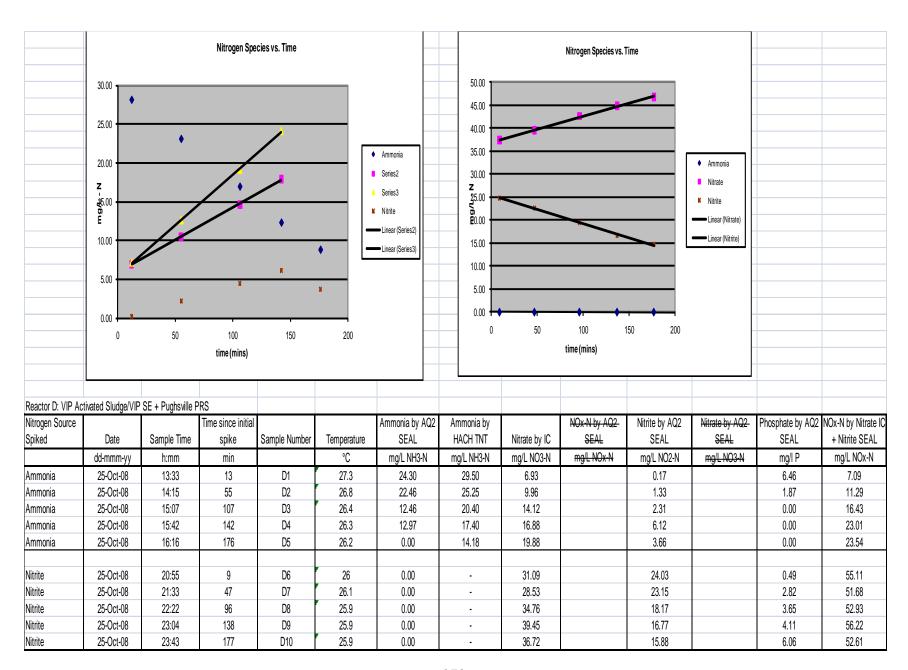


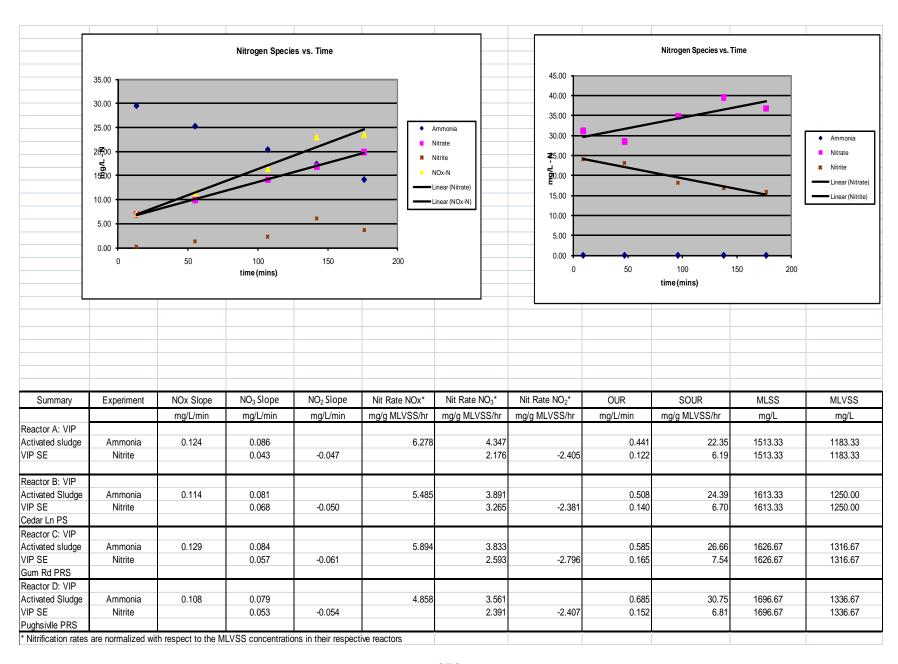
Week 8 – Pump/Pressure Reducing Stations Day 2 AOB/NOB

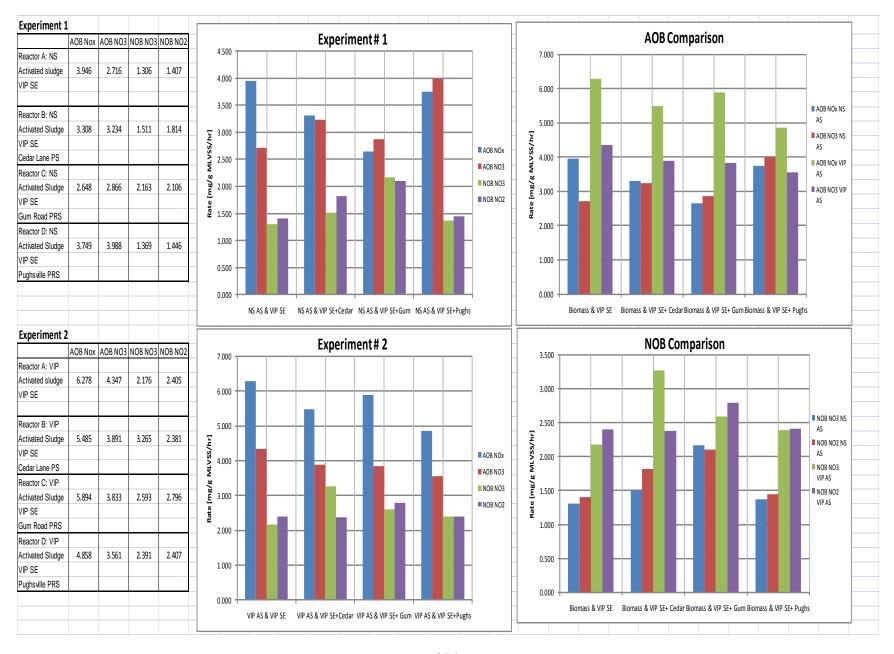
	Report for Nanse	emond Nitrificati	on Inhibition Stu	ıdy									
AOB & NOB Exp	perimentation:												
I. AOB													
	r 3L reactors with	20 mg/L NH₄.											
		inously by use of st	tir bars.										
	-	e is added to the											
				eks experiments t	he biomass was co	ncentrated from 20	L to 9 L.						
d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 9 L. e. Constant DO and pH were monitored and logged throughout the experiment.													
	f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.												
					d implement the so	olenoid valves.							
-		-			•	d into sample tubes t	through millinore 0	45 um filters					
						tor lines for each rea		io più intero					
ii Nedetors D,	, 0, 4114 5 111 1116 41	iluciic source ilicor	porateu 12 or viii .	JE und 12 or the di	oodatea iiitereept	tor micoror cucinicu	ctori						
II. NOB													
	r31 reactors with	25 mg/L NO afte	rammonia levels	wara < 1 mg/l NH .	.N which was charl	ked through the use	of HACH TNT 921 /	mmonia					
		0	in overnight to con	O . 3		keu tili ougii tile use	OTTACIT INT 031 A	illillollia					
		inously by use of st		isuille dii presenti	allillollia.								
			ed throughout the	evneriment									
			s to maintain DO le		noidvalvos								
					noid valves. d implement the so	alamaidualuaa							
			i pri recording as w	veli as programan	u impiement me st	olellolu valves.							
1 f F						والمتعاربة والمتعادية والمتعاراة		45 £!k					
						d into sample tubes t		45 μm filters.					
						d into sample tubes t tor lines for each rea		45 μm filters.					
								45 μm filters.					
								45 μm filters.					
								45 μm filters.					
								45 μm filters.					
g. Reactors B,		iluent source incor						45 μm filters.					
g. Reactors B,	, C, and D in the d	iluent source incor						45 μm filters.					
g. Reactors B,	, C, and D in the d	iluent source incor	porated 1L of VIP:			tor lines for each rea	ctor.	45 μm filters.	NO. No. 100	Nicio La Acc	Nive had 600	Physical In Account	
g. Reactors B, Reactor A: VIP Ac	tivated Sludge/VIF	iluent source incor	porated 1L of VIP:	SE and 1L of the as	ssociated intercept	tor lines for each rea	Ammonia by		NOX-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	' '	,
g. Reactors B, Reactor A: VIP Ac	ctivated Sludge/VIF	iluent source incor	porated 1L of VIP:		ssociated intercept Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	SEÁL	SEÁL	SEAL	SEAL	NOx-N by Nitrate IC + Nitrite SEAL
g. Reactors B, Reactor A: VIP Ac Vitrogen Source Spiked	ctivated Sludge/VIF	iluent source incor	porated 1L of VIP: Time since initial spike min	SE and 1L of the as	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC	,	SEAL mg/L NO2-N	,	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
g. Reactors B, Reactor A: VIP Ac Vitrogen Source Spiked	ctivated Sludge/VIF Date dd-mmm-yy 25-Oct-08	Sample Time h:mm 13:28	Time since initial spike min 11	SE and 1L of the as	Temperature °C 28.2	Ammonia by AQ2 SEAL mg/L NH3-N 11.44	Ammonia by HACH TNT mg/L NH3-N 16.80	Nitrate by IC mg/L NO3-N 10.46	SEÁL	SEAL mg/L NO2-N 0.41	SEAL	SEAL mg/I P 1.53	+ Nitrite SEAL mg/L NOx-N 10.87
g. Reactors B, Reactor A: VIP Ac Vitrogen Source Spiked Ammonia	Ctivated Sludge/VIF Date dd-mmm-yy 25-Oct-08 25-Oct-08	Sample Time h:mm 13:28 14:12	Time since initial spike min 11 55	Sample Number A1 A2	Temperature °C 28.2 28.5	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30	Nitrate by IC mg/L NO3-N 10.46 13.23	SEÁL	SEAL mg/L NO2-N 0.41 2.73	SEAL	SEAL mg/I P 1.53 1.53	+ Nitrite SEAL mg/L NOx-N 10.87 15.97
g. Reactors B, Reactor A: VIP Ac Vitrogen Source Spiked Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08	Sample Time h:mm 13:28 14:12 15:03	Time since initial spike min 11 55 96	Sample Number A1 A2 A3	Temperature °C 28.2 28.5 27.2	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25	SEAL	SEAL mg/l P 1.53 1.53 1.92	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57
g. Reactors B, Reactor A: VIP Ac Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time h:mm 13:28 14:12 15:03 15:39	Time since initial spike min 11 55 96 132	Sample Number A1 A2 A3 A4	Temperature °C 28.2 28.5 27.2 26.6	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29
g. Reactors B, Reactor A: VIP Ac Alitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08	Sample Time h:mm 13:28 14:12 15:03	Time since initial spike min 11 55 96	Sample Number A1 A2 A3	Temperature °C 28.2 28.5 27.2	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25	SEAL	SEAL mg/l P 1.53 1.53 1.92	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57
g. Reactors B, Reactor A: VIP Ac Mitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time	Time since initial spike min 11 55 96 132 165	Sample Number A1 A2 A3 A4 A5	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07 0.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01 23.09	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29 6.36	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41 2.68	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29 29.45
g. Reactors B, Reactor A: VIP Ac Ilitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time	Time since initial spike min 11 55 96 132 165	Sample Number A1 A2 A3 A4 A5	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07 0.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01 23.09	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29 6.36	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41 2.68	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29 29.45
g. Reactors B, Reactor A: VIP Ac Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time	Time since initial spike min 11 55 96 132 165 10 47	Sample Number A1 A2 A3 A4 A5	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07 0.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01 23.09 32.20 32.86	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29 6.36 22.65 21.62	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41 2.68	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29 29.45 54.85 54.48
g. Reactors B, Reactor A: VIP Ac Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Altirite Nitrite	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time h:mm 13:28 14:12 15:03 15:39 16:12 20:53 21:30 22:19	Time since initial spike min 11 55 96 132 165 10 47 96	Sample Number A1 A2 A3 A4 A5 A6 A7 A8	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07 0.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01 23.09 32.20 32.86 35.40	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29 6.36 22.65 21.62 20.16	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41 2.68 4.24 4.90 5.79	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29 29.45 54.85 54.48 55.56
g. Reactors B, Reactor A: VIP Ac litrogen Source piked mmonia mmonia mmonia mmonia mmonia litrite litrite	Date dd-mmm-yy 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08 25-Oct-08	Sample Time	Time since initial spike min 11 55 96 132 165 10 47	Sample Number A1 A2 A3 A4 A5	Temperature	Ammonia by AQ2 SEAL mg/L NH3-N 11.44 6.74 2.25 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 16.80 12.30 6.00 2.07 0.00	Nitrate by IC mg/L NO3-N 10.46 13.23 17.32 21.01 23.09 32.20 32.86	SEÁL	SEAL mg/L NO2-N 0.41 2.73 5.25 5.29 6.36 22.65 21.62	SEAL	SEAL mg/l P 1.53 1.53 1.92 2.41 2.68	+ Nitrite SEAL mg/L NOx-N 10.87 15.97 22.57 26.29 29.45 54.85 54.48





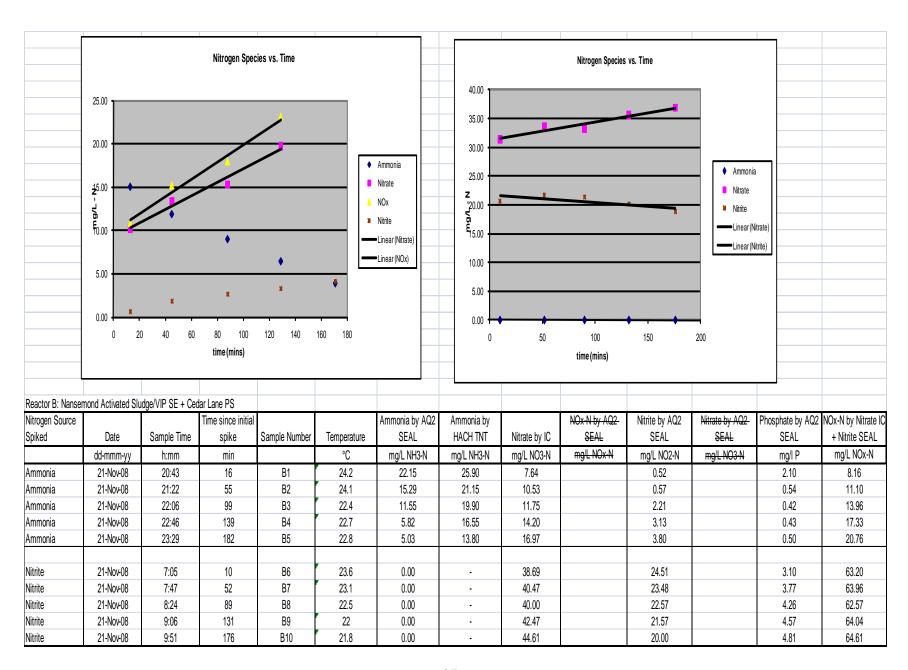


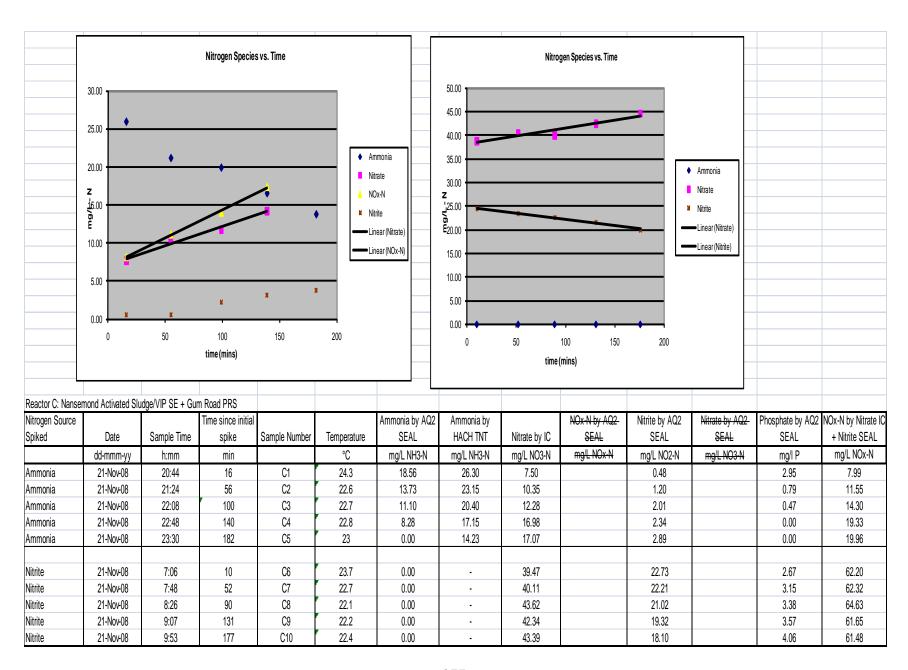


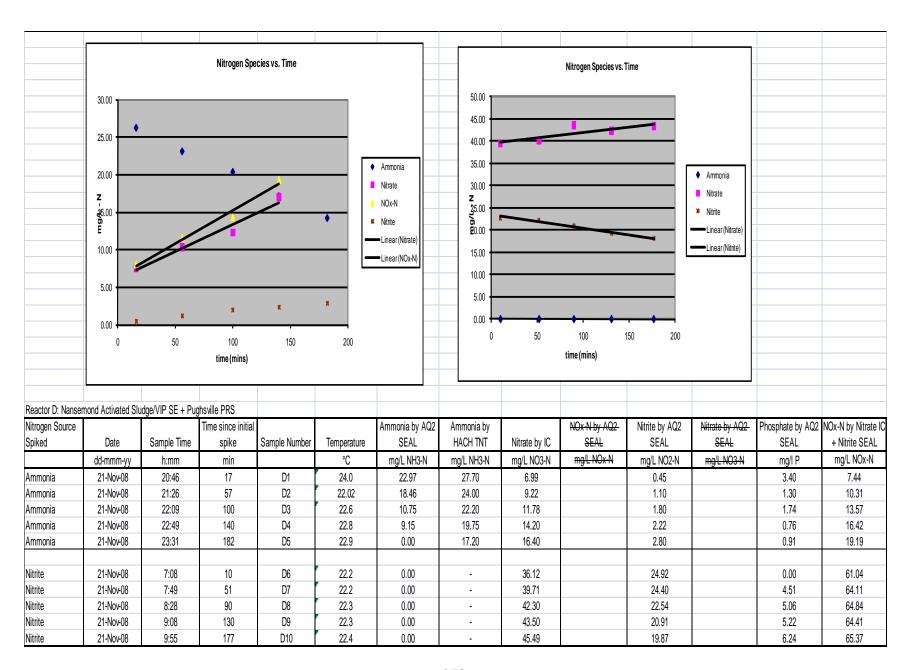


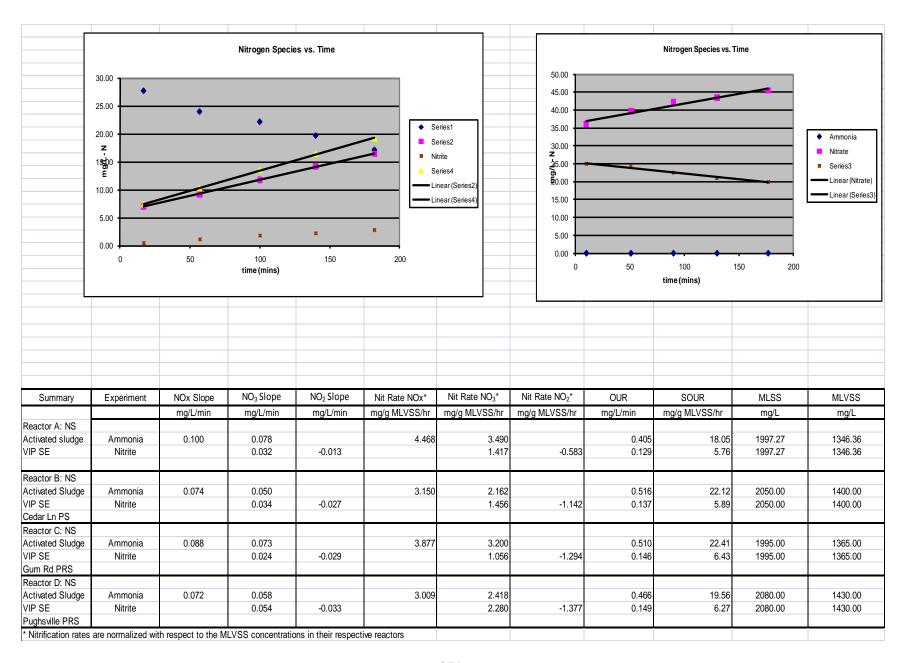
Week 9 – Pump/Pressure Reducing Stations (RERUN) Day 1 AOB/NOB

					KEKUN)	Day 1 AC	R/NOR						
Sample Data F	Report for Nanse	emond Nitrificati	ion Inhibition Stu	idy									
AOB & NOB Ex	perimentation:												
I. AOB													
	r 3L reactors with	20 mg/L NH ₄ .											
	or is running conti	· ,	tir bars										
	liluent source is ac												
				eks exneriments t	he hinmass was co	ncentrated from 201	I to 9 I						
			ed throughout the		ne biomass was co		L 10 3 L.						
			rs to maintain DO l		noid valves.								
			d pH recording as w	-		olenoid valves.							
						l into sample tubes t	hrough millinore ()	45 um filters					
						or lines for each read		ιο μιτιπιείο					
i. redectors b	, c, and b in the di	iluciti source ilicor	poruteu 11 or vii	DE una 1E or the as	ocatea intercept	or mics for eachieux	ctor.						
II. NOB													
-	r31 reactors with	25 mg/L NO - afte	rammonia levels	were < 1 mg/l NH -	N which was chac	ked through the use	of HACH TNT 921 /	mmonia					
			ın overnight to cor			cu till ough the use	OTTIACITINI 031 F	Milliollia					
	or is running conti			isuille dii present	allillollia.								
			นา มสาร. ed throughout the	ovnoriment									
			rs to maintain DO l		noidvalvos								
			s to maintain bo i			alanaidualuas							
		•			•		h o .: !!!: 0	AF £!k					
						d into sample tubes t or lines for each rea		45 μm nilers.					
g. Reactors B	, c, and D in the d	lluent source incor	porated IL of VIP	SE and IL or the as	sociated intercept	or lines for each real	ctor.						
Reactor A: Nanse	mond Activated Slu	udge/VIP SE											
Nitrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/I P	mg/L NOx-N
Ammonia	21-Nov-08	20:39	13	A1	26	5.70	15.00	10.19		0.66		0.56	10.85
Ammonia	21-Nov-08	21:21	45	A2	25.4	195.14	11.85	13.34		1.85		0.68	15.19
Ammonia	21-Nov-08	22:04	88	A3	23.5	-1.54	8.98	15.31		2.65		0.78	17.96
Ammonia	21-Nov-08	22:45	129	A4	23.2	-4.29	6.42	19.77		3.29		1.02	23.06
Ammonia	21-Nov-08	23:27	171	A5	23.1	-7.24	3.89	18.92		4.13		1.24	23.05
Nitrite	21-Nov-08	7:03	10	A6	23.4	0.00	-	31.39		20.67		3.51	52.06
Vitrite	21-Nov-08	7:45	52	A7	23.3	0.00	-	33.54		21.75		4.39	55.28
Nitrite	21-Nov-08	8:23	90	A8	22.6	0.00	-	33.22		21.38		4.61	54.60
Nitrite	21-Nov-08	9:05	132	A9	22.1	0.00	-	35.60		20.09		4.91	55.69
Nitrite	21-Nov-08	9:49	176	A10	21.9	0.00	-	36.87		18.84		5.10	55.71



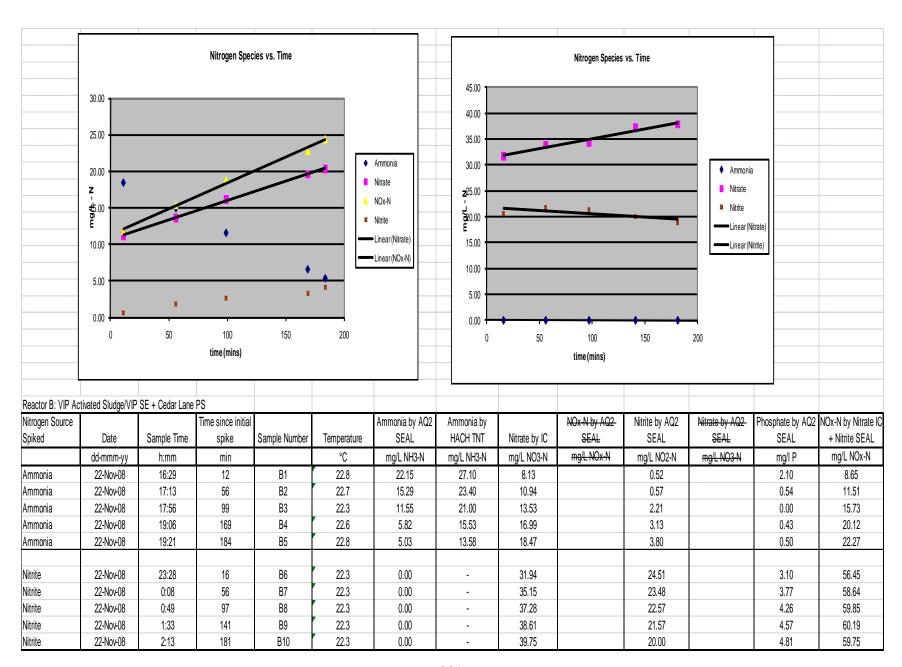


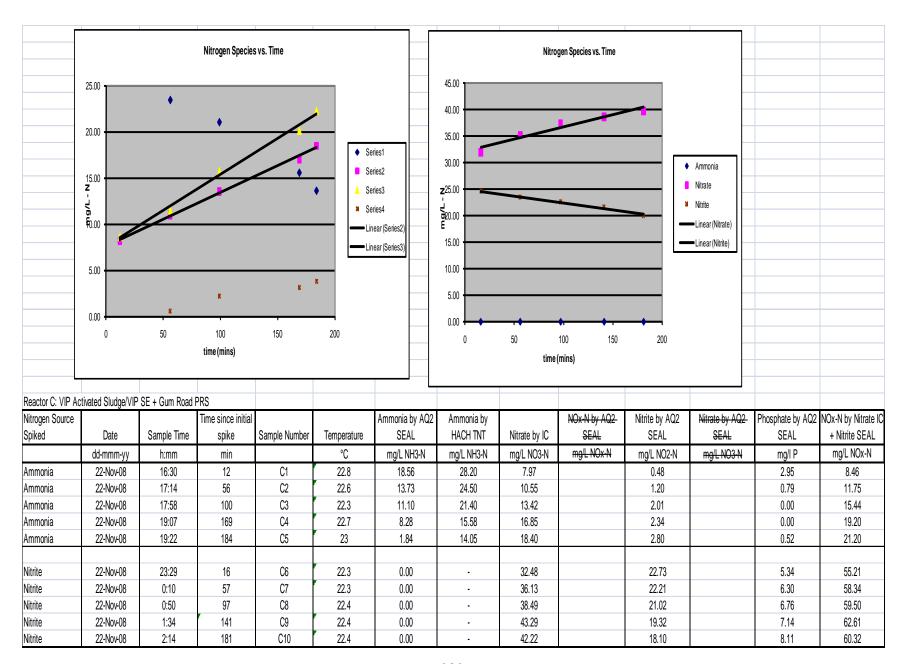


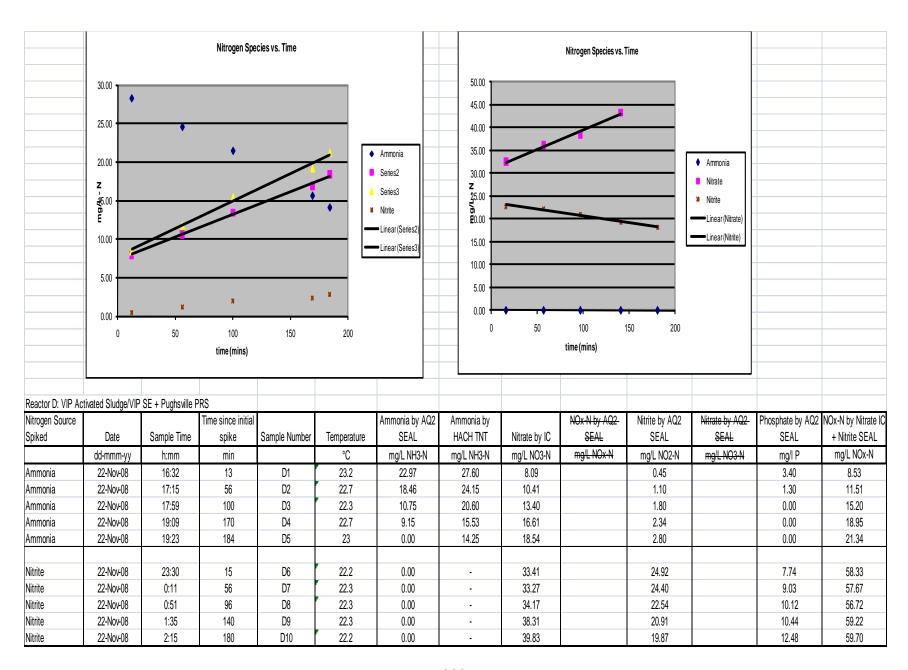


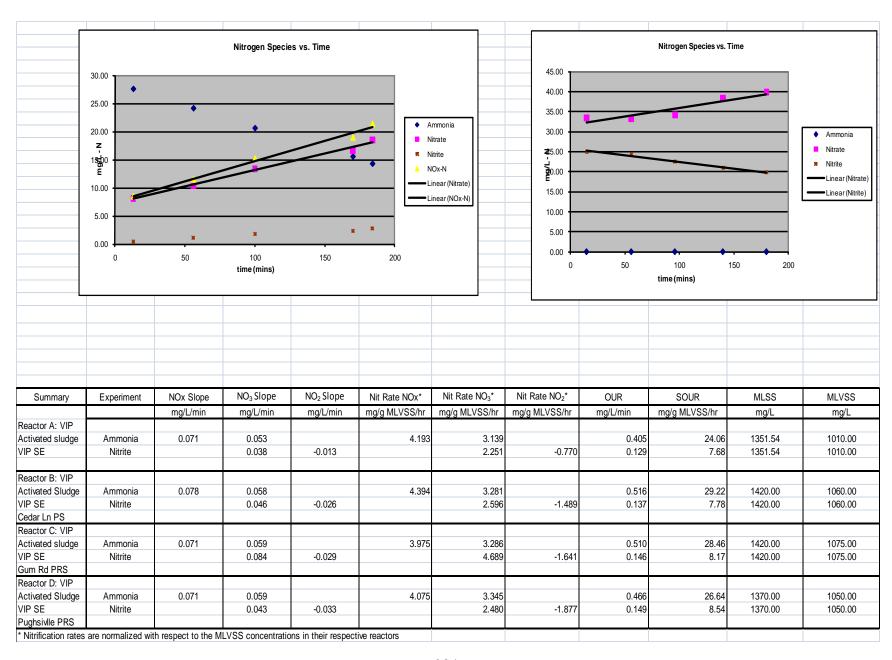
Week 9 – Day 2 AOB/NOB

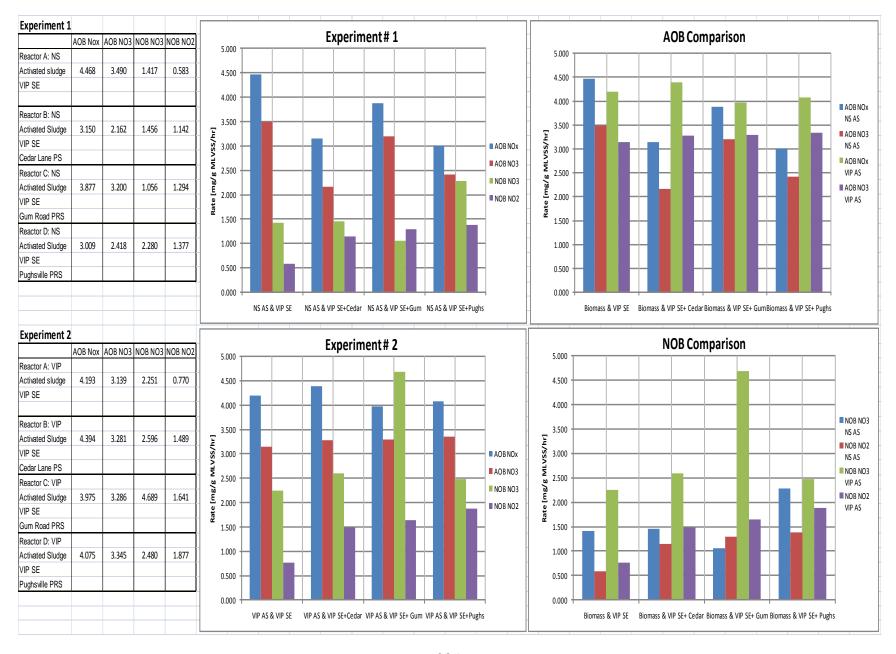
AOB & NOB Experime I. AOB a. Spiked four 3L rei b. Each reactor is ru c. 1900 mL of the di d. 1L of concentrat	nentation: eactors with 20		on Inhibition Stu	dy									
I. AOB a. Spiked four 3L rea b. Each reactor is ru c. 1900 mL of the di d. 1 L of concentrat	eactors with 20												
I. AOB a. Spiked four 3L rea b. Each reactor is ru c. 1900 mL of the di d. 1 L of concentrat	eactors with 20												
a. Spiked four 3L rea b. Each reactor is ru c. 1900 mL of the di d. 1 L of concentrat													
b. Each reactor is ru c. 1900 mL of the di d. 1 L of concentrat													
c. 1900 mL of the di d. 1 L of concentrate	unning contin	0 mg/L NH₄.											
c. 1900 mL of the di d. 1 L of concentrate	unning contin	ously by use of sti	ir bars.										
	-												
	ited biomass is	added to the rea	actors. For this we	eks experiments t ^j	he biomass was co	ncentrated from 201	to 9 L.						
e. constant DO and			ed throughout the										
f. Oxygen/airblend	nd was sparged	d into the reactors	s to maintain DO le	evels through soler	noid valves.			-					
g. LabView software						olenoid valves.		-					
h. 5 samples were c	collected over	ra period of 3 hou	urs through sampl	ing ports on the re	actors and filtered	d into sample tubes t	hrough millipore 0.	45 μm filters					
						tor lines for each reac		·					
		'			·								
II. NOB													
a. Spiked four 3L rea	eactors with 2	5 mg/L NO ₂ - after	rammonia levels v	vere <1 mg/L NH ₂ -	N which was checl	ked through the use	of HACH TNT 831 A	mmonia					
method. In addition a	all reactors w	ere allowed to rui	n overnight to con	sume all present a	ımmonia.	· ·							
b. Each reactor is ru			•										
c. Constant DO and				experiment.									
d. Oxygen/airblend	nd was sparged	d into the reactors	s to maintain DO le	evels through sole:	noid valves.								
e. LabView software						olenoid valves.							
f. 5 samples were c							hrough millipore 0.	45 μm filters.					
g. Reactors B, C, and	nd D in the dilu	uent source incorp	porated 1L of VIP S	SE and 1L of the as	sociated intercept	tor lines for each read	ctor.						
Reactor A: VIP Activated	ed Sludge/VIP	SE											
		-											
litrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2		NOx-N by Nitrate IC
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL
	d-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/I P	mg/L NOx-N
	22-Nov-08	16:27	11	A1	24.4	5.70	18.50	11.16		0.66		0.56	11.82
	22-Nov-08	17:12	56	A2	24.3	195.14	15.00	13.56		1.85		0.68	15.40
	22-Nov-08	17:55	99	A3	23.0	-1.54	11.63	16.18		2.65		0.78	18.83
	22-Nov-08	19:05	169	A4	23.1	0.00	6.64	19.56		3.29		1.02	22.85
Ammonia 22	22-Nov-08	19:20	184	A5	23.1	0.00	5.42	20.29		4.13		1.24	24.42
litrite 22	22-Nov-08	22-27	16	۸۶	22.1	0.00		21.65		20.67		2.51	52.33
inne Zi	22-Nov-08 22-Nov-08	23:27 0:07	16 56	A6 A7	22.1	0.00		31.65 33.92		20.67		3.51 4.39	52.33
itrito 21	ZZ-1NUV-U0	0.07	90	ΑI	_		•			21.70		4.39	00.07
	22-Nov-08	0.48	07	٨٥	22.2	0.00	_	3/1.28		21.20		4.61	55.66
litrite 22	22-Nov-08 22-Nov-08	0:48 1:32	97 141	A8 A9	22.2 22.3	0.00	•	34.28 37.16		21.38 20.09		4.61 4.91	55.66 57.25





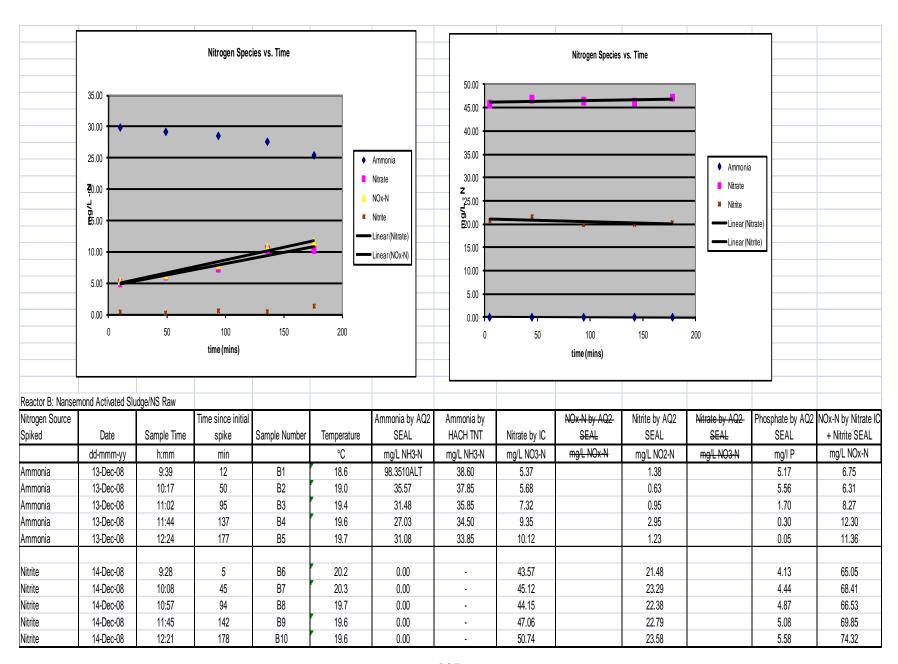


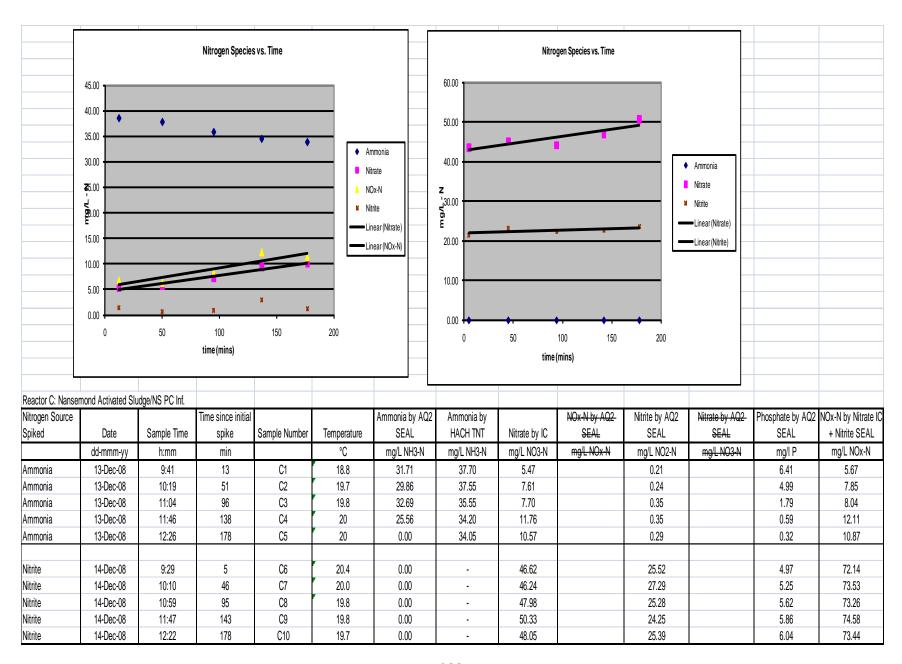


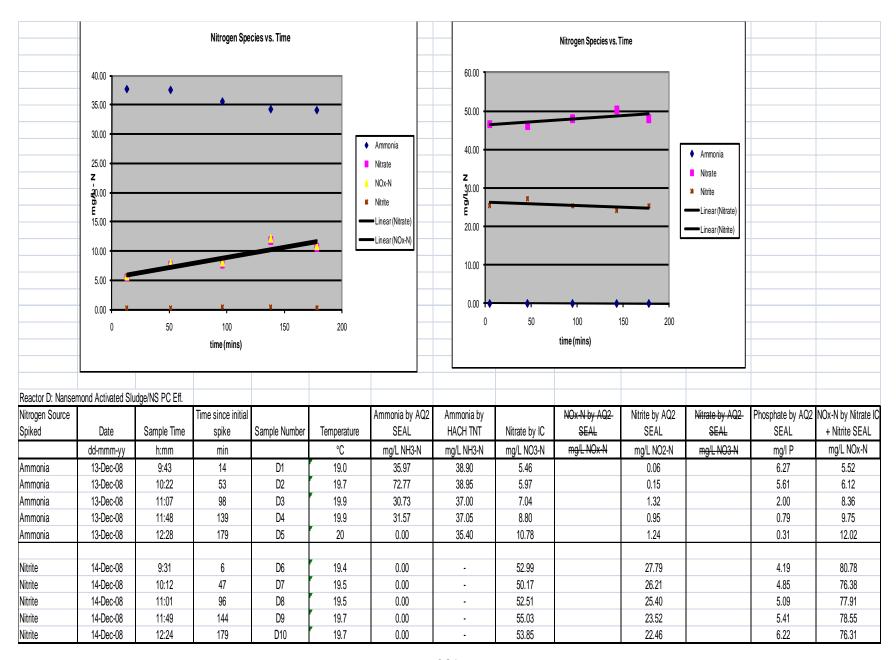


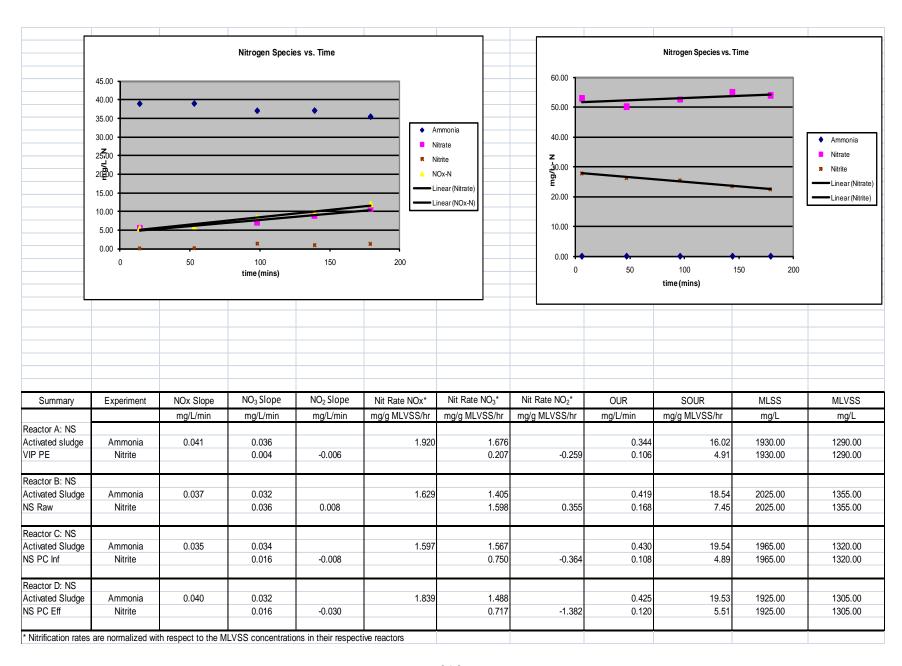
Week 10 – NS Raw, PCI, PCE Day 1 AOB/NOB

Sample Data R	Report for Nanse	emond Nitrificati	ion Inhibition Stu	ıdy									
AOB & NOB Ext	perimentation:												
I. AOB	,												
	r 3L reactors with	25 mg/L NH ₄											
		inously by use of s	tir hars										
	-	dded to the reacto											
				aks avnarimants t	ha hiamass was sa	ncentrated from 20	1 +0 0 1						
			ed throughout the		ile Dioillass was co	incentrated nom 20	L 10 9 L.						
			-	evels through sole	noidualuos								
						alamaidualuaa							
				vell as program and				45 (1)					
						d into sample tubes t		45 µm filters					
i. Keactors B,	, c, and D in the di	lluent source incor	porated 1L of VIP	SE and 1L of the as	ssociated intercept	tor lines for each rea	ctor.						
II. NOB													
1						ked through the use	of HACH TNT 831 A	mmonia					
1			•	nsume all present a	ammonia.								
		inously by use of s											
c. Constant D	O and pH were m	ionitored and loggi	ed throughout the	experiment.									
d. Oxygen/ai	ir blend was sparge	ed into the reactor	s to maintain DO l	evels through sole	noid valves.								
1.130	fr				al francisco de la compansión de la compa								
e. Labview sc	oftware was used	to manage DO and	l pH recording as v	veil as program ani	a impiement the si	olenoid valves.							
						olenoid valves. d into sample tubes t	hrough millipore 0.	45 μm filters.					
f. 5 samples	were collected ov	er a period of 3 ho	urs through samp	ling ports on the re	eactors and filtered			45 μm filters.					
f. 5 samples	were collected ov	er a period of 3 ho	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples	were collected ov	er a period of 3 ho	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples	were collected ov	er a period of 3 ho	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples	were collected ov	er a period of 3 ho	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d	er a period of 3 hc iluent source incor	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples of g. Reactors B,	were collected ov	er a period of 3 hc iluent source incor	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d	er a period of 3 hc iluent source incor	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.					
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d	er a period of 3 hc iluent source incor	urs through samp	ling ports on the re	eactors and filtered	d into sample tubes t		45 μm filters.	NOx-N-by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d	er a period of 3 hc iluent source incor	urs through samp porated 1L of VIP	ling ports on the re	eactors and filtered	d into sample tubes to rlines for each rea	ctor.		NOX-N-by AQ2- SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2- SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d mond Activated SI	era period of 3 ho iluent source incor udge/VIP PE	urs through samp porated 1L of VIP	ling ports on the re SE and 1L of the as	eactors and filtered essociated intercept	d into sample tubes to rlines for each rea	Ammonia by	45 μm filters. Nitrate by IC mg/L NO3-N		,	,		
f. 5 samples g. Reactors B,	were collected ov , C, and D in the d mond Activated SI	era period of 3 ho iluent source incor udge/VIP PE Sample Time	urs through samp porated 1L of VIP	ling ports on the re SE and 1L of the as	eactors and filtered sociated intercept	d into sample tubes tor lines for each rea Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	SEAL	SEÁL	SEÁL	SEAL	+ Nitrite SEAL
f. 5 samples g. Reactors B, g. Reactor A: Nansel Nitrogen Source Spiked	were collected ov , C, and D in the d mond Activated SI Date dd-mmm-yy	era period of 3 ho iluent source incor udge/VIP PE Sample Time h:mm	urs through samp porated 1L of VIP Time since initial spike min	ling ports on the re SE and 1L of the as Sample Number	eactors and filtered sociated intercept Temperature	d into sample tubes tor lines for each rea Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC	SEAL	SEAL mg/L NO2-N	SEÁL	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
f. 5 samples g. Reactors B, Reactor A: Nansei Nitrogen Source Spiked	mond Activated SI Date dd-mmm-yy 13-Dec-08	era period of 3 ho iluent source incor udge/VIP PE Sample Time h:mm 9:36	Time since initial spike min	ling ports on the re SE and 1L of the as Sample Number	Temperature °C 20.9	Ammonia by AQ2 SEAL mg/L NH3-N 25.97	Ammonia by HACH TNT mg/L NH3-N 29.90	Nitrate by IC mg/L NO3-N 4.97	SEAL	SEAL mg/L NO2-N 0.33	SEÁL	SEAL mg/l P 2.03	+ Nitrite SEAL mg/L NOx-N 5.31
f. 5 samples g. Reactors B, Reactor A: Nansel Nitrogen Source Spiked Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08	era period of 3 ho iluent source incor udge/VIP PE Sample Time h:mm 9:36 10:15	Time since initial spike min 10 49	Sample Number A1 A2	Temperature °C 20.9 20.7	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20	Nitrate by IC mg/L NO3-N 4.97 6.05	SEAL	SEAL mg/L NO2-N 0.33 0.23	SEÁL	SEAL mg/l P 2.03 0.65	+ Nitrite SEAL mg/L NOx-N 5.31 6.28
f. 5 samples g. Reactors B, Reactor A: Nansel Nitrogen Source Spiked Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08	udge/VIP PE Sample Time h:mm 9:36 10:15 11:00	Time since initial spike min 10 49 94	Sample Number A1 A2 A3	Temperature °C 20.9 20.7 20.8	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62	SEÁL	SEAL mg/l P 2.03 0.65 0.21	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91
f. 5 samples g. Reactors B, Reactor A: Nansel Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08	udge/VIP PE Sample Time h:mm 9:36 10:15 11:00 11:42	Time since initial spike min 10 49 94 136	Sample Number A1 A2 A3 A4	Temperature °C 20.9 20.7 20.8 20.4	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT 22.56	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55 27.60	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29 10.33	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62 0.44	SEÁL	SEAL mg/l P 2.03 0.65 0.21 0.27	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91 10.77
f. 5 samples g. Reactors B, g. Reactor A: Nansel Altrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08	udge/VIP PE Sample Time h:mm 9:36 10:15 11:00 11:42	Time since initial spike min 10 49 94 136	Sample Number A1 A2 A3 A4	Temperature °C 20.9 20.7 20.8 20.4	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT 22.56	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55 27.60	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29 10.33	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62 0.44	SEÁL	SEAL mg/l P 2.03 0.65 0.21 0.27	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91 10.77
f. 5 samples g. Reactors B, Reactor A: Nansel Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08	udge/VIP PE Sample Time h:mm 9:36 10:15 11:00 11:42 12:22	Time since initial spike min 10 49 94 136 176	Sample Number A1 A2 A3 A4 A5	Temperature °C 20.9 20.7 20.8 20.4 20.2	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT 22.56 22.35	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55 27.60 25.40	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29 10.33 10.36	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62 0.44 1.33	SEÁL	SEAL mg/l P 2.03 0.65 0.21 0.27 0.29	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91 10.777 11.69
f. 5 samples g. Reactors B, Reactor A: Nansel Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08	era period of 3 ho iluent source incor udge/VIP PE Sample Time h:mm 9:36 10:15 11:00 11:42 12:22	Time since initial spike min 10 49 94 136 176	Sample Number A1 A2 A3 A4 A5	Temperature °C 20.9 20.7 20.8 20.4 20.2	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT 22.56 22.35	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55 27.60 25.40	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29 10.33 10.36	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62 0.44 1.33	SEÁL	SEAL mg/l P 2.03 0.65 0.21 0.27 0.29 5.40	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91 10.77 11.69
f. 5 samples g. Reactor A: Nanser Reactor A: Nanser Nitrogen Source Spiked Ammonia Ammonia Ammonia Ammonia Ammonia Ammonia Mitrite	mond Activated SI Date dd-mmm-yy 13-Dec-08 13-Dec-08 13-Dec-08 13-Dec-08 14-Dec-08 14-Dec-08	udge/VIP PE Sample Time h:mm 9:36 10:15 11:00 11:42 12:22 9:26 10:06	Time since initial spike min 10 49 94 136 176	Sample Number A1 A2 A3 A4 A5 A6 A7	Temperature °C 20.9 20.7 20.8 20.4 20.2 20.2 20.3	Ammonia by AQ2 SEAL mg/L NH3-N 25.97 23.85 92.8506ALT 22.56 22.35	Ammonia by HACH TNT mg/L NH3-N 29.90 29.20 28.55 27.60 25.40	Nitrate by IC mg/L NO3-N 4.97 6.05 7.29 10.33 10.36 45.79 46.83	SEAL	SEAL mg/L NO2-N 0.33 0.23 0.62 0.44 1.33 20.69 21.60	SEÁL	SEAL mg/l P 2.03 0.65 0.21 0.27 0.29 5.40 5.76	+ Nitrite SEAL mg/L NOx-N 5.31 6.28 7.91 10.77 11.69 66.47 68.42



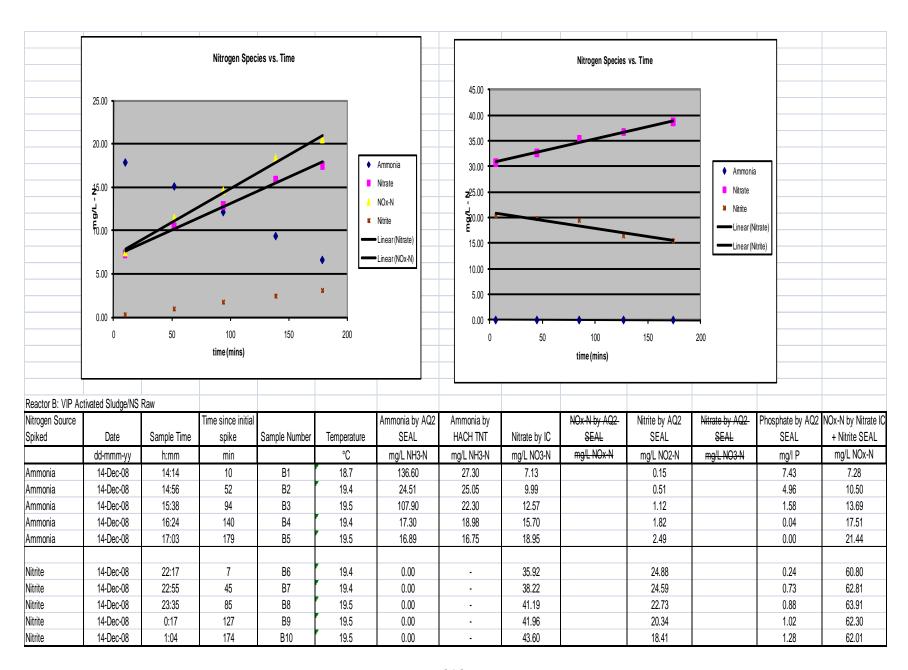


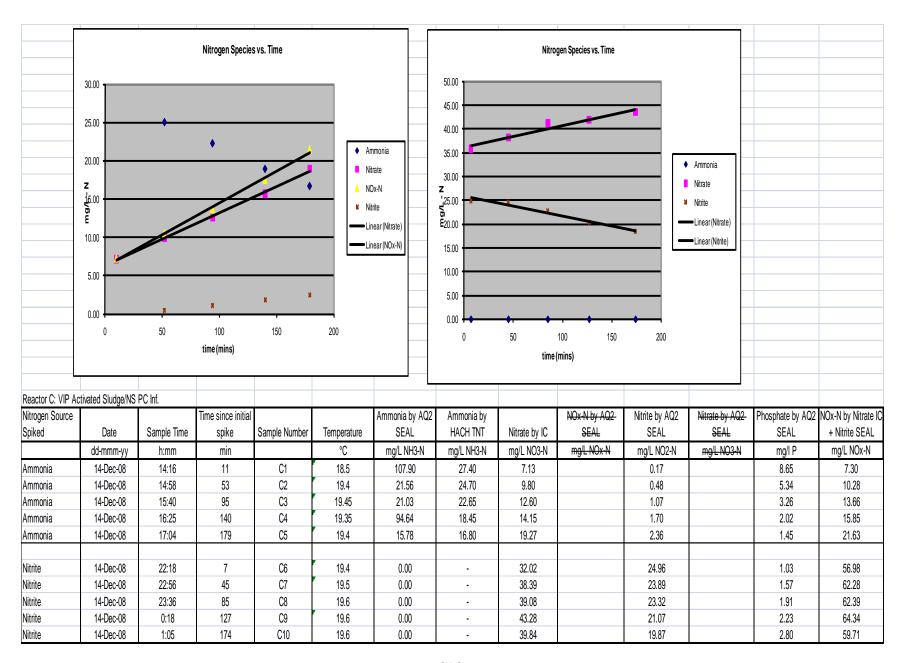


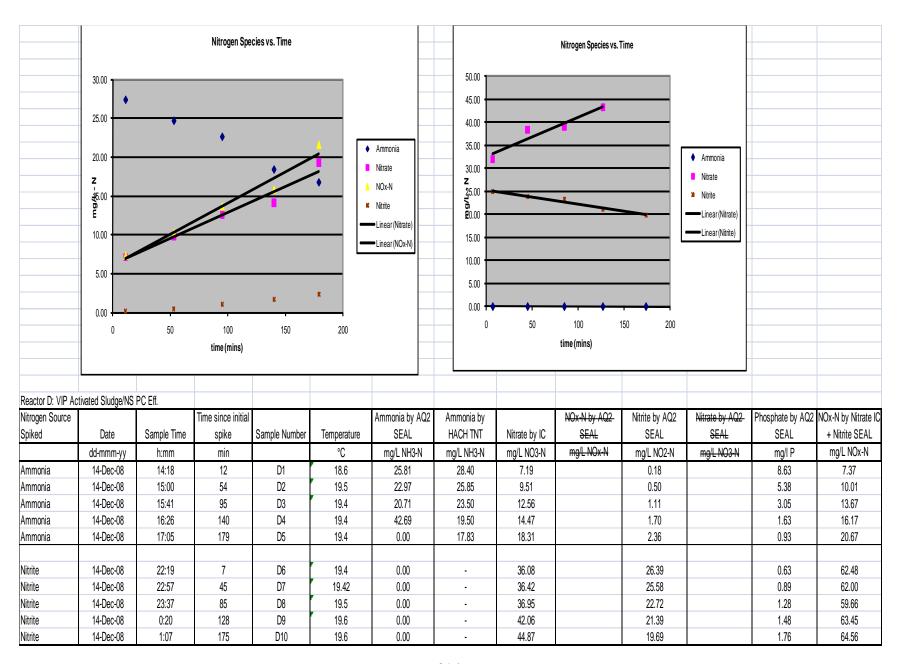


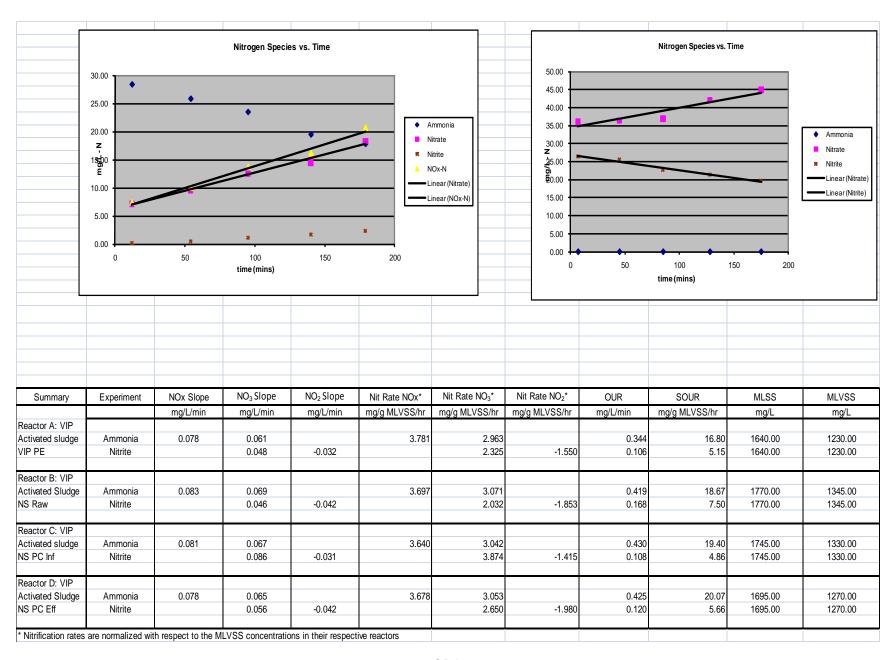
Week 10 – Day 2 AOB/NOB

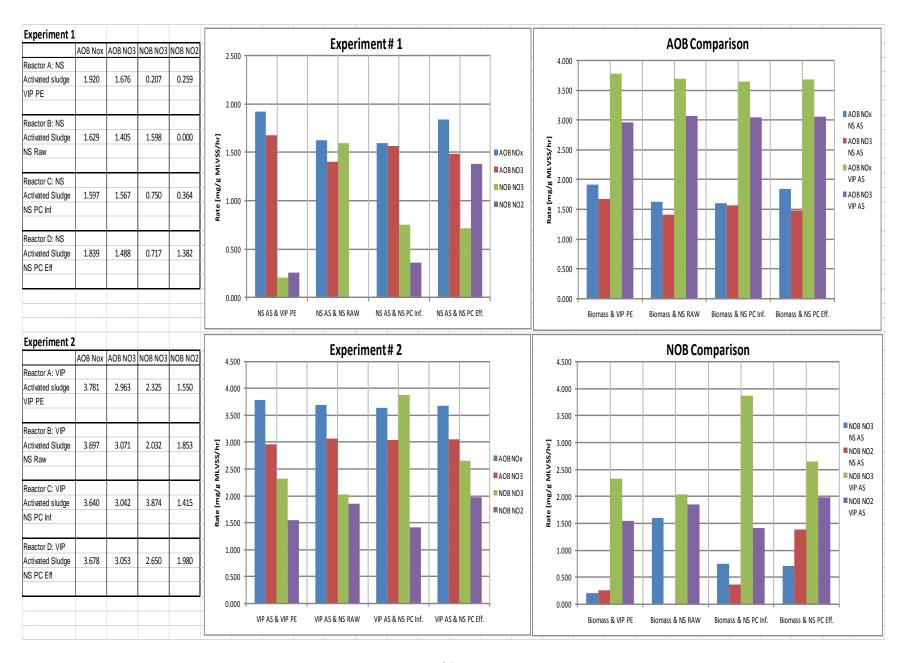
- ampio Daia N	kepoπ for ivanse	emond Nitrificati	on Inhibition Stu	ıdy									
AOB & NOB Exp	perimentation:												
I. AOB													
a. Spiked four	r 3L reactors with	10 mg/L NH₄.											
		inously by use of st	tir bars.										
	-	ce is added to the											
d. 1L of conce	entrated biomass	is added to the re	actors. For this we	eks experiments t	he biomass was co	ncentrated from 201	L to 9 L.						
e. Constant D	O and pH were m	onitored and logg	ed throughout the	experiment.									
	•	ed into the reactor	-		noid valves.								
		to manage DO and				olenoid valves.							
						into sample tubes t	hrough millipore 0.	45 um filters					
						or lines for each read							
	,		, == =										
II. NOB													
	r 3L reactors with	25 mg/L NO ₂ - afte	rammonia levels v	were < 1 mg/L NH ₂ -	N which was check	ed through the use	of HACH TNT 831 A	mmonia					
		were allowed to ru											
		inously by use of st		isame an present									
		onitored and logge		experiment.									
		ed into the reactor	•	•	noid valves								
g. Oxygen/air													
						olenoid valves.							
e. LabView so	oftware was used	to manage DO and	pH recording as w	vell as program and	d implement the so		hrough millinore 0	45 um filters					
e. LabView so f. 5 samples v	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered			45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v	oftware was used were collected ov	to manage DO and er a period of 3 ho	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t		45 μm filters.					
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t or lines for each rea	ctor.	45 μm filters.	NOx N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov , C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as wurs through sampl	vell as program and ling ports on the re	d implement the so eactors and filtered	into sample tubes t			NOx-N-by AQ2- SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2- SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate II + Nitrite SEAL
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov . C, and D in the d bitwated Sludge/VIF	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as	d implement the so eactors and filtered sociated intercept	into sample tubes to r lines for each read	Ammonia by	45 μm filters. Nitrate by IC mg/L NO3-N	,	,	,		-
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac litrogen Source piked	oftware was used were collected ov . C, and D in the d	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as Sample Number	d implement the so eactors and filtered esociated intercept Temperature	into sample tubes to or lines for each read Ammonia by AQ2 SEAL	Ammonia by HACH TNT mg/L NH3-N	Nitrate by IC	SEÁL	SEAL	SEÁL	SEAL	+ Nitrite SEAL
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac itrogen Source piked	oftware was used were collected ov C, and D in the d clivated Sludge/VIF Date dd-mmm-yy	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as	d implement the so eactors and filtered sociated intercept sociated intercept Temperature	into sample tubes to or lines for each read Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT	Nitrate by IC	SEÁL	SEAL mg/L NO2-N	SEÁL	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac litrogen Source piked mmonia mmonia	oftware was used were collected ov C, and D in the d clivated Sludge/VIF Date dd-mmm-yy 14-Dec-08	to manage DO and era period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as programani ling ports on the re SE and 1L of the as Sample Number	d implement the so eactors and filtered sociated intercept sociated intercept Temperature °C 20.8	Ammonia by AQ2 SEAL mg/L NH3-N 16.78	Ammonia by HACH TNT mg/L NH3-N 17.80	Nitrate by IC mg/L NO3-N 7.26	SEÁL	SEAL mg/L NO2-N 0.23	SEÁL	SEAL mg/I P 3.32	+ Nitrite SEAL mg/L NOx-N 7.49
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac itrogen Source piked mmonia mmonia mmonia	oftware was used were collected ov C, and D in the d clivated Sludge/VIF Date dd-mmm-yy 14-Dec-08 14-Dec-08	to manage DO and er a period of 3 ho iluent source incor	pH recording as w urs through sampl porated 1L of VIP:	vell as program aniling ports on the re SE and 1L of the as Sample Number	Temperature °C 20.8 20.8	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05	Nitrate by IC mg/L NO3-N 7.26 10.49	SEÁL	SEAL mg/L NO2-N 0.23 0.94	SEÁL	SEAL mg/l P 3.32 1.58	+ Nitrite SEAL mg/L NOx-N 7.49 11.44
e. LabView so f. 5 samples v g. Reactors B,	oftware was used were collected ov. C, and D in the d bit	to manage DO and er a period of 3 ho iluent source incor	pH recording as w urs through sample porated 1L of VIP:	vell as program aniling ports on the re SE and 1L of the as Sample Number	Temperature °C 20.8 20.8 20.7	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87	SEÁL	SEAL mg/L NO2-N 0.23 0.94 1.70	SEÁL	SEAL mg/l P 3.32 1.58 1.11	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac itrogen Source piked mmonia mmonia mmonia mmonia	bftware was used were collected ov. C, and D in the d collecte	Sample Time h:mm 14:13 14:55 15:37 16:22	pH recording as w urs through sample porated 1L of VIP:	sell as program and ling ports on the research and 1L of the assert and	Temperature °C 20.8 20.8 20.7 20.3	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69 0.00	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10 9.35	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87 15.86	SEÁL	SEAL mg/L NO2-N 0.23 0.94 1.70 2.46	SEÁL	SEAL mg/I P 3.32 1.58 1.11 0.93	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57 18.32
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac trogen Source biked mmonia mmonia mmonia mmonia mmonia mmonia	bftware was used were collected ov. C, and D in the d collecte	Sample Time h:mm 14:13 14:55 15:37 16:22	pH recording as w urs through sample porated 1L of VIP:	sell as program and ling ports on the research and 1L of the assert and	Temperature °C 20.8 20.8 20.7 20.3	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69 0.00	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10 9.35	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87 15.86	SEÁL	SEAL mg/L NO2-N 0.23 0.94 1.70 2.46	SEÁL	SEAL mg/I P 3.32 1.58 1.11 0.93	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57 18.32
e. LabView so f. 5 samples v g. Reactors B, eactor A: VIP Ac itrogen Source piked mmonia mmonia mmonia mmonia mmonia mmonia itrite	britware was used were collected ov. C, and D in the d collect	Sample Time h:mm 14:13 14:55 15:37 16:22 17:02	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 10 52 94 139 179	Sample Number A1 A2 A3 A4 A5	Temperature °C 20.8 20.8 20.7 20.3 19.9	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10 9.35 6.61	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87 15.86 17.48	SEÁL	SEAL mg/L NO2-N 0.23 0.94 1.70 2.46 3.03	SEÁL	SEAL mg/l P 3.32 1.58 1.11 0.93 0.96	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57 18.32 20.51
e. LabView so f. 5 samples v g. Reactors B, leactor A: VIP Ac litrogen Source piked mmonia mmonia mmonia mmonia mmonia itrite litrite	britware was used were collected ov. C, and D in the d collect	Sample Time h:mm 14:13 14:55 15:37 16:22 17:02	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 10 52 94 139 179	Sample Number A1 A2 A3 A4 A5	Temperature °C 20.8 20.8 20.7 20.3 19.9	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10 9.35 6.61	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87 15.86 17.48	SEÁL	SEAL mg/L NO2-N 0.23 0.94 1.70 2.46 3.03	SEÁL	SEAL mg/l P 3.32 1.58 1.11 0.93 0.96	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57 18.32 20.51
e. LabView so f. 5 samples v g. Reactors B, Reactor A: VIP Ac litrogen Source ipiked mmoniammoniammoniammonia	britware was used were collected ov. C, and D in the d collect	Sample Time h:mm 14:13 14:55 15:37 16:22 17:02	pH recording as w urs through sample porated 1L of VIP: Time since initial spike min 10 52 94 139 179	Sample Number A1 A2 A3 A4 A5	Temperature °C 20.8 20.8 20.7 20.3 19.9	Ammonia by AQ2 SEAL mg/L NH3-N 16.78 33.31 101.69 0.00 0.00	Ammonia by HACH TNT mg/L NH3-N 17.80 15.05 12.10 9.35 6.61	Nitrate by IC mg/L NO3-N 7.26 10.49 12.87 15.86 17.48	SEÁL	SEAL mg/L NO2·N 0.23 0.94 1.70 2.46 3.03 20.40 19.92	SEÁL	SEAL mg/l P 3.32 1.58 1.11 0.93 0.96 4.31 2.24	+ Nitrite SEAL mg/L NOx-N 7.49 11.44 14.57 18.32 20.51 51.18 52.53











Week 11 – FeCl3 Addition Day 1 AOB/NOB

Nitrite

Nitrite

6-Jan-09

6-Jan-09

6-Jan-09

18:38

19:24

20:09

89

135

180

Week II Samnle Data R			ion Inhibition Stu		ОВ									
Jampie Dala N	report for marise	THORIG MIGHICAG		uy										
AOB & NOB Exp	narimantation:													
I. AOB	Jerimentation.													
	. Al	20 ma/L NIII												
	r 3L reactors with 2	· .	eta kana											
	or is running conti													
	iluent source is ad					15 20								
					ne biomass was co	ncentrated from 201	L to 9 L.							
			ed throughout the	•										
, .				evels through sole										
				vell as program and										
		•		0.		l into sample tubes t	• .	•						
						ed on an as Fe basis	relative to the 1L B	iomass						
j. A two hour	period was allowe	d after FeCl ₃ addi	tion to monitor pH	I adjustment due t	o FeCl ₃ addition.									
II. NOB														
a. Spiked four	r 3L reactors with 2	25 mg/L NO ₂ - afte	r ammonia levels	were < 1 mg/L NH ₃ -	N which was checl	ked through the use	of HACH TNT 831 A	Ammonia						
method. In add	lition all reactors v	vere allowed to ru	ın overnight to cor	nsume all present a	mmonia.									
b. Each reacto	or is running conti	nously by use of st	tir bars.											
c. Constant DO	O and pH were mo	onitored and logge	ed throughout the	experiment.										
				evels through sole:	noid valves.									
				vell as program and		olenoid valves.								
						l into sample tubes t	hrough millipore 0	.45 um filters.						
						ed on an as Fe basis								
				H adjustment due		000000000000000000000000000000000000000		.0						
III.71CHO IIOUI	period masanom	ea arter reeizada	ncion co monicor p	i ruajustinent uue	to reeizadattoni									
loactor A: Nancor	mond Activated Slu	idao/\/ID SE												
icacioi A. Nansci	IIIOIIU ACIIVAICU SII	auge/vir 3L												
litrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC	Phosphate by
piked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL	HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
mmonia	6-Jan-09	10:06	13	A1	21.2	16.09	16.80	10.52		1.28		0.21	11.80	0.675
mmonia	6-Jan-09	10:49	56	A2	20.3	14.22	15.05	13.27		0.43		0.31	13.70	0.725
mmonia	6-Jan-09	11:32	99	A3	19.7	13.52	13.03	15.03		1.01		0.39	16.04	0.875
mmonia	6-Jan-09	12:12	139	A4	19.4	11.69	11.18	18.49		0.97		0.48	19.46	0.95
mmonia	6-Jan-09	12:53	180	A5	19.6	8.48	9.18	22.83		1.82		0.62	24.65	1.07
Vitrite	6-Jan-09	17:15	6	A6	18.6	0.00	-	36.41		29.31		1.46	65.73	1.82
litrite	6-Jan-09	17:55	46	A7	18.7	0.00	-	32.86		28.50		1.63	61.36	2.04
e												0.40		

34.31

40.75

35.26

27.88

26.62

25.30

A8

Α9

A10

18.8

18.8

0.00

0.00

0.00

62.19

67.37

60.56

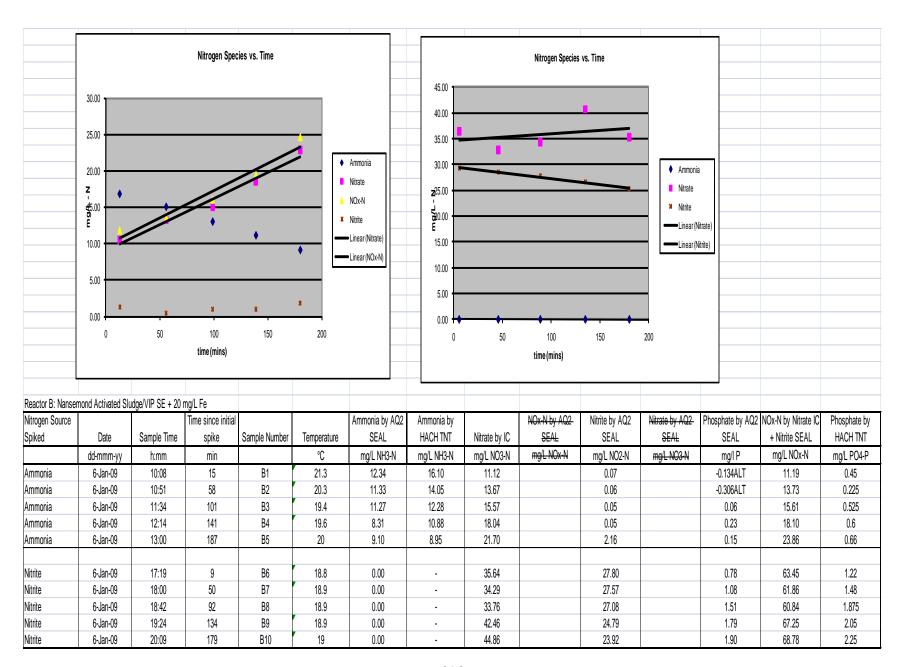
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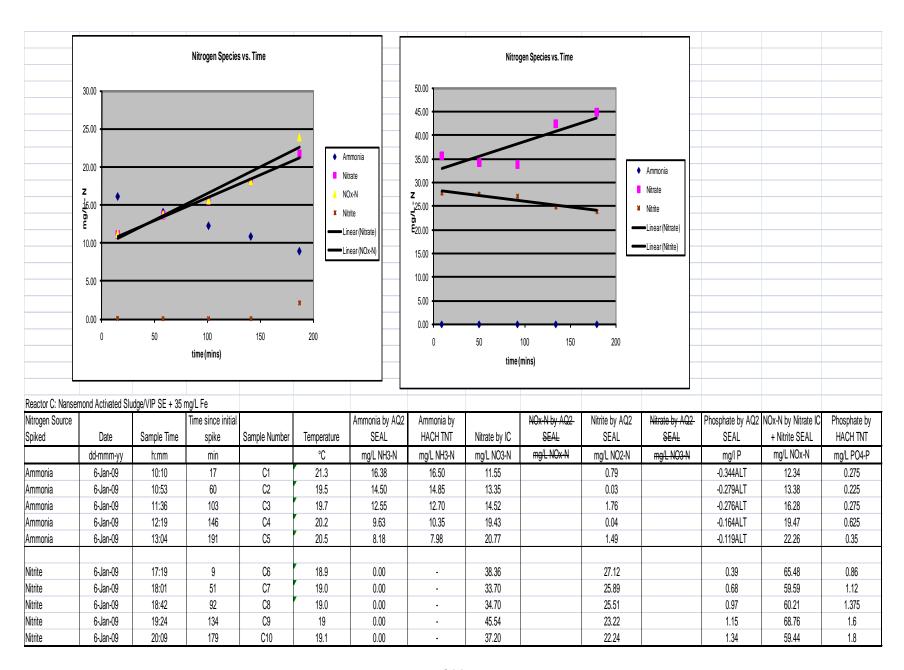
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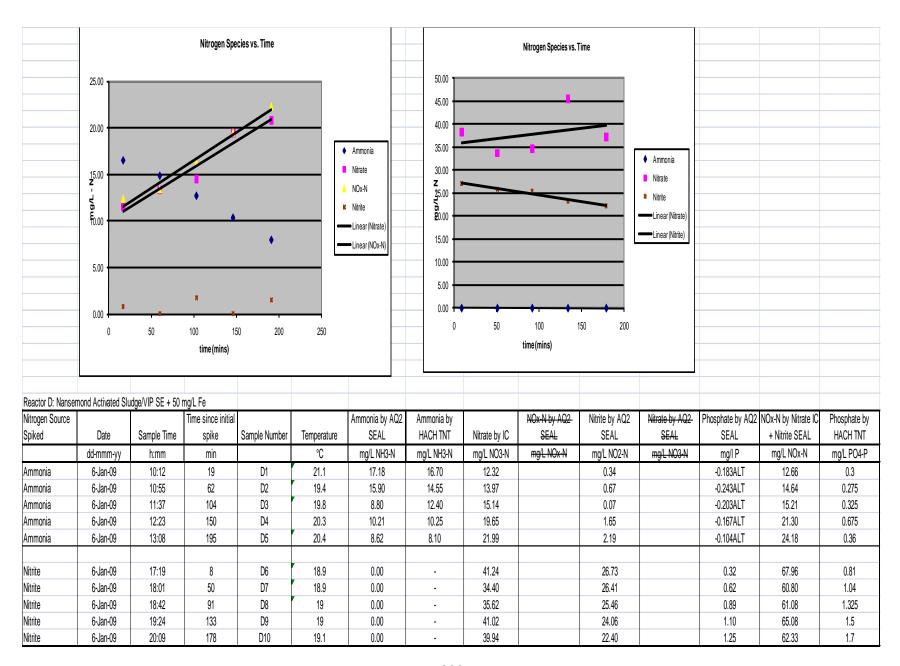
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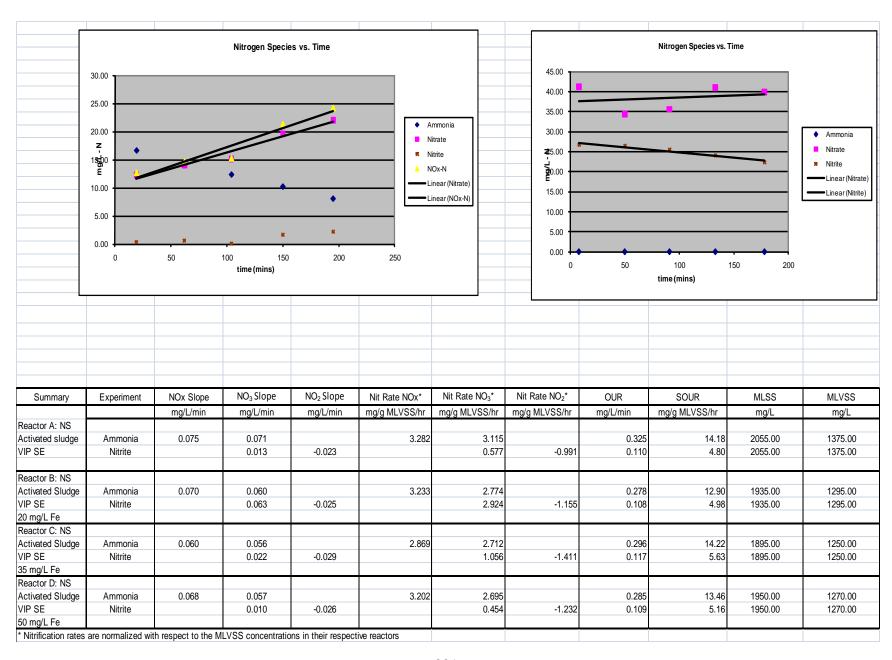
2.16

2.42









Week 11 – Day 2 AOB/NOB

Nitrite

Nitrite

Nitrite

Nitrite

14-Dec-08

14-Dec-08

14-Dec-08

14-Dec-08

22:54

23:34

0:16

1:03

45

85

127

174

Α7

A8

Α9

A10

#DIV/0!

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0.00

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		emond Nitrificati		ıdv										
AOB & NOB Exp	perimentation:													
I. AOB														
	r 3L reactors with	20 mg/L NH.												
		nously by use of st	ir hars											
	U	e is added to the r												
				eks evneriments t	ne hinmass was co	ncentrated from 20 L	to 9 I							
		onitored and logge			ic bioinuss was coi	ncentrated from 201	10 3 L			NOTE NOB EXPER	IMENT WAS NOT R	<mark>UN DURING DAY 2 E</mark>	EXPERIMENTATION	
	•		•	evels through soler	oovley bion									
				vell as program and		alenoid valves								
-		-			•	into sample tubes tl	h rough millinara ()	15 um filtors						
						ed on an as Fe basis r								
						eu oil dii da re basis i	elative to the 1F p	IUIIId55						
J. A two nour	period was allowe	ed after FeCi ₃ addit	tion to monitor ph	l adjustment due to	reci ₃ addition.									
II NOD														
II. NOB		"				1.1	· · · · · · · · · · · · · · · · · · ·							
		O		o . 3		ked through the use	of HACH TNT 831 A	Ammonia						
				nsume all present a	mmonia.									
	-	nously by use of st												
	•	onitored and logge	-											
d. Oxygen/aii	r blend was sparge	ed into the reactor	s to maintain DO le	evels through soler	noid valves.									
e. LabView so	oftware was used t	to manage DO and	pH recording as w	vell as program and	l implement the so	olenoid valves.								
f. 5 samples v	were collected ove	er a period of 3 ho	urs through samp	ling ports on the re	actors and filtered	into sample tubes t	hrough millipore 0.	.45 μm filters.						
g. Reactors B,	, C, and D in the di	iluent source incor	porated varying co	oncentrations of Fe	Cl ₃ calculated base	ed on an as Fe basis i	relative to the 1L B	iomass						
h. A two hour	r period was allow	ed after FeCl ₃ add	ition to monitor pl	H adjustment due 1	to FeCl ₃ addition									
					•									
Reactor A: VIP Ac	ctivated Sludge/VIP	SE												
														i
Nitrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AO2	NOx-N by Nitrate IC	Phosphate by
Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL	HACH TNT
opou	dd-mmm-yy	h:mm	min	Campio Hamboi	°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	7-Jan-09	19:12	9	A1	19.2	19.10	18.60	15.21		0.16	IIIg E 110011	0.88	15.37	1.08
Ammonia	7-Jan-09	19:55	52	A2	19.1	17.96	16.60	16.79		0.46		0.96	17.25	1.15
Ammonia	7-Jan-09	20:39	96	A3	19.0	15.22	14.70	17.64		0.46		0.95	18.30	1.25
Ammonia	7-Jan-09	21:22	139	A4	19.0	13.39	12.93	17.11		0.79		0.99	17.91	1.25
Ammonia	7-Jan-09	22:04	181	A5	18.9	12.05	11.73	20.31		0.73		1.04	21.25	1.225
VIIIU	7 0011 03	££.07	101	/10	10.0	12.00	11.70	20.01		0.01		1.07	21.20	1.220
Vitrite	14-Dec-08	22:15	6	A6	#DIV/0!	0.00		0.00		0.00		0.00	0.00	0
The tr	14 000-00	22.10	45	110	#DIV/U:	0.00	-	0.00		0.00		0.00	0.00	0

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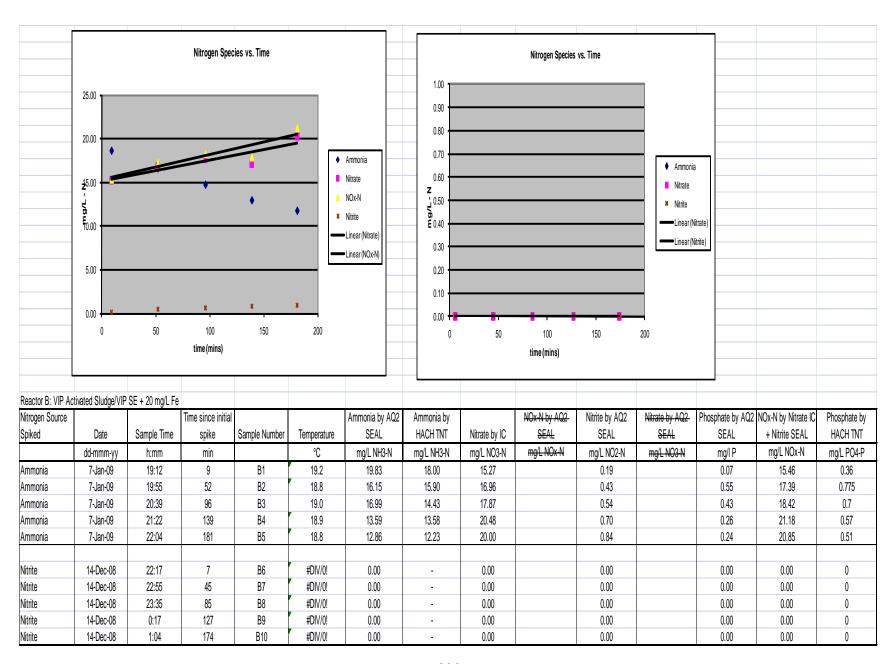
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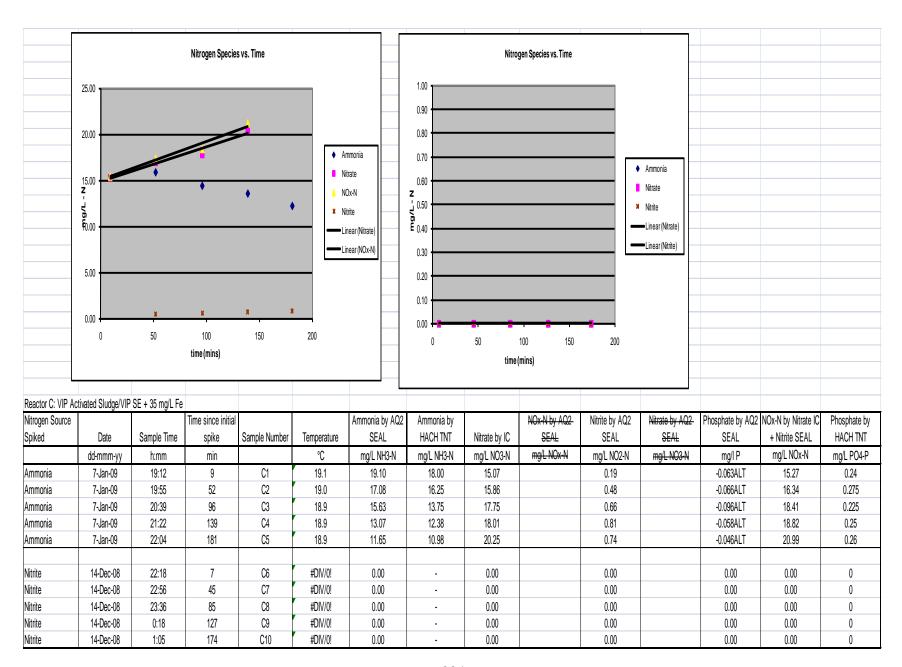
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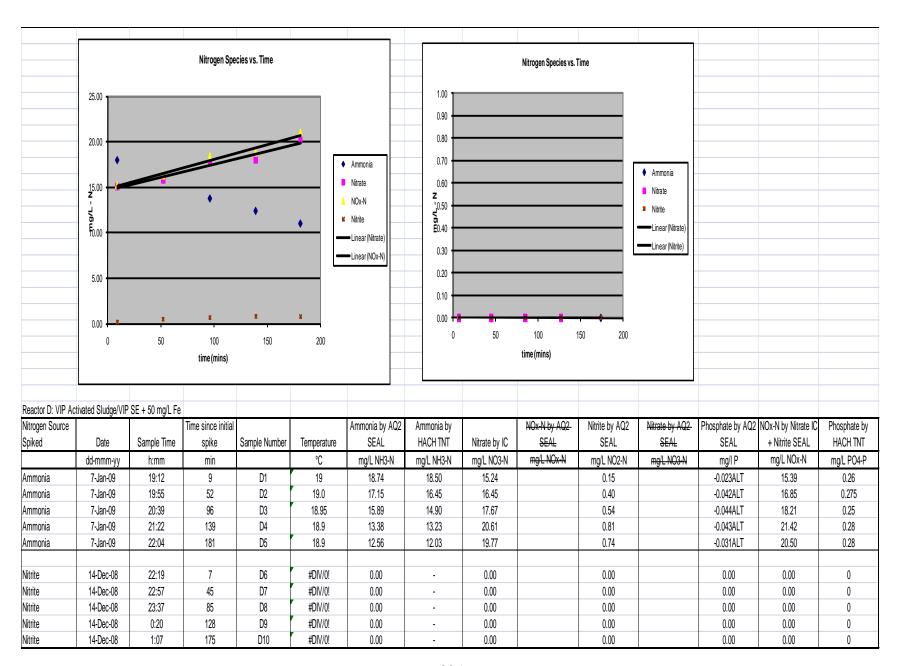
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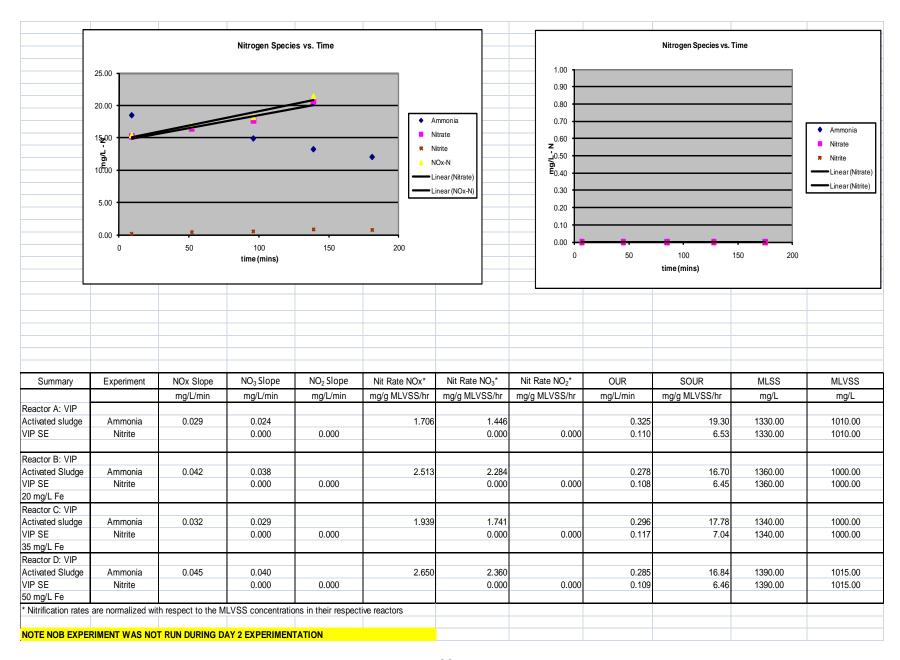
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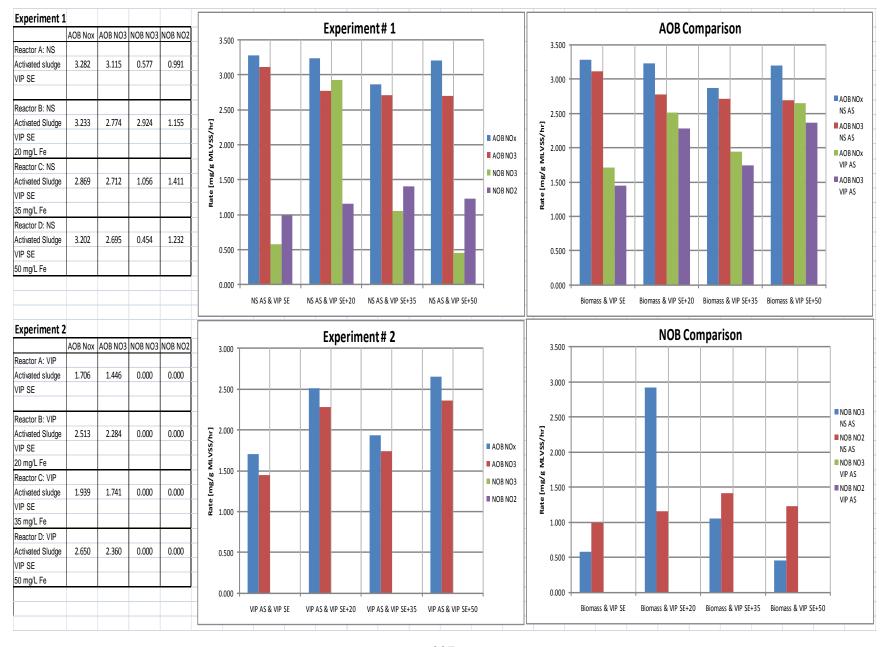
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Week 12 – FeCl3 Addition RERUN Day 1 AOB/NOB

13-Jan-09

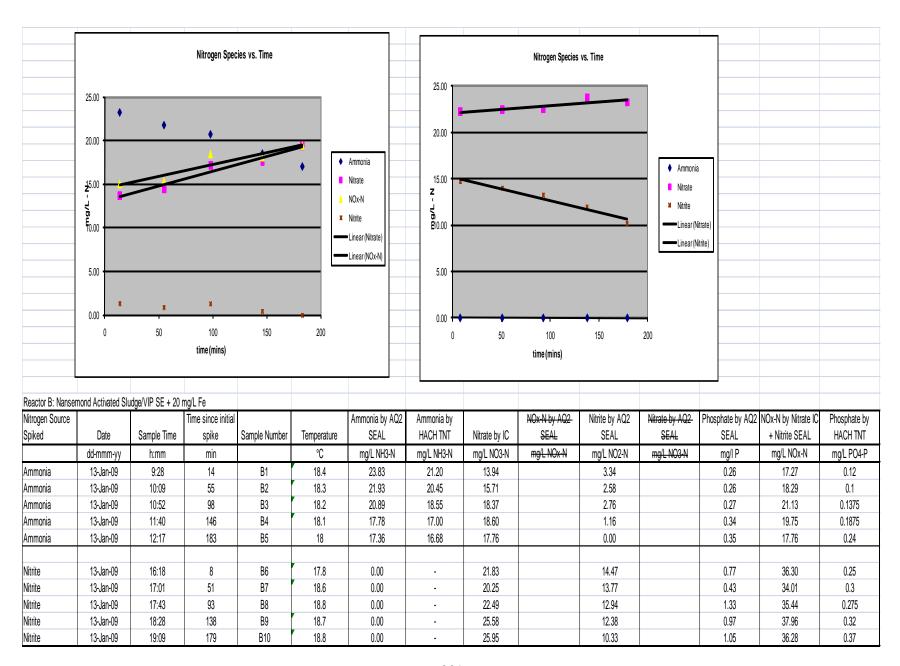
nple Data Report for Nansemond Nitrification Inhibition Study							
OB & NOB Experimentation:							
AOB							
a. Spiked four 3L reactors with 20 mg/L NH _a .							
b. Each reactor is running continously by use of stir bars .							
c. 2 L of the diluent source is added to the reactors.							
d. 1L of concentrated biomass is added to the reactors. For this weeks experiment	s the biomass was o	oncentrated from	201 to 91.				
e. Constant DO and pH were monitored and logged throughout the experiment.	5 tire 5 to 11 do 5						
f. Oxygen/air blend was sparged into the reactors to maintain DO levels through so	lenoid valves.						
g. LabView software was used to manage DO and pH recording as well as program a		solenoid valves.					
h. 5 samples were collected over a period of 3 hours through sampling ports on the			es th rough millipore	0.45 µm filters			
i. Reactors B, C, and D in the diluent source incorporated varying concentrations of							
. A two hour period was allowed after FeCl $_{\!3}$ addition to monitor pH adjustment due							
NOB							
a. Spiked four 3L reactors with 25 mg/L NO ₂ ⁻ after ammonia levels were <1 mg/L NI		cked through the u	ise of HACH TNT 831	Ammonia			
ethod. In addition all reactors were allowed to run overnight to consume all preser	t ammonia.						
b. Each reactor is running continously by use of stir bars.							
c. Constant DO and pH were monitored and logged throughout the experiment.							
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through so							
e. LabView software was used to manage DO and pH recording as well as program a	•						
f. 5 samples were collected over a period of 3 hours through sampling ports on the							
g. Reactors B, C, and D in the diluent source incorporated varying concentrations of	J.		sis relative to the 1L	Biomass			
h. A two hour period was allowed after FeCl ₃ addition to monitor pH adjustment du	e to FeCl ₃ addition.						

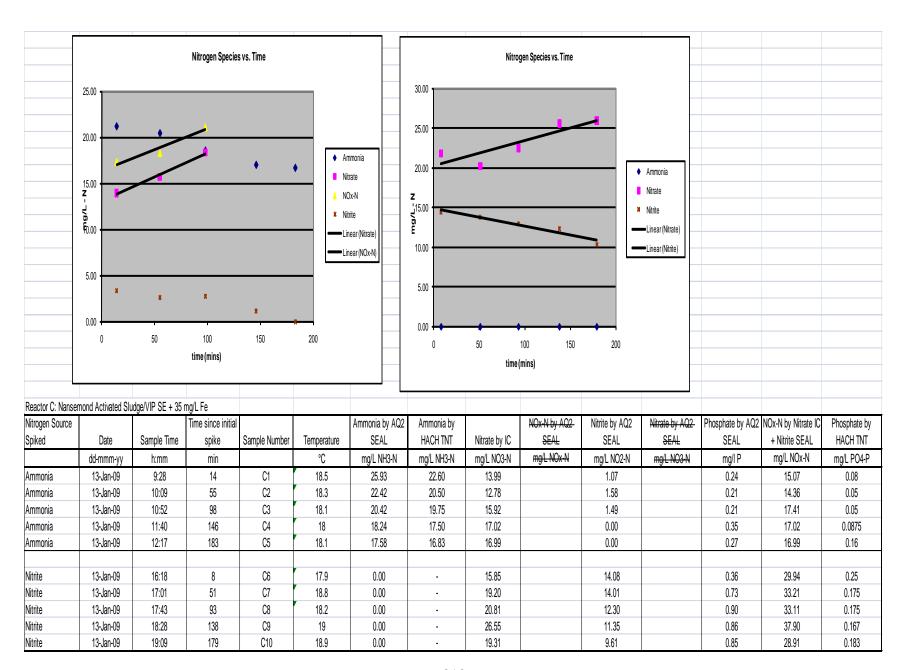
Reactor A: Nanser	mond Activated Slu	udge/VIP SE												
Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOx-N by AQ2 SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2 SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	13-Jan-09	9:28	14	A1	18.4	26.19	23.20	13.67		1.30		0.78	14.97	0.68
Ammonia	13-Jan-09	10:09	55	A2	18.2	23.44	21.75	14.39		0.89		0.79	15.28	0.6
Ammonia	13-Jan-09	10:52	98	A3	18.2	28.11	20.70	17.13		1.28		0.87	18.41	0.7625
Ammonia	13-Jan-09	11:40	146	A4	18	19.12	18.50	17.61		0.45		0.97	18.06	0.95
Ammonia	13-Jan-09	12:17	183	A5	18	17.83	17.03	19.37		0.00		1.09	19.37	1.05
Nitrite	13-Jan-09	16:18	8	A6	17.7	0.00	-	22.26		14.69		1.39	36.95	1.35
Nitrite	13-Jan-09	17:01	51	A7	18.375	0.00	-	22.44		13.98		1.96	36.42	1.5
Nitrite	13-Jan-09	17:43	93	A8	18.7	0.00	-	22.51		13.24		2.26	35.75	1.65
Nitrite	13-Jan-09	18:28	138	A9	18.6	0.00	-	23.71		11.93		2.17	35.64	1.8

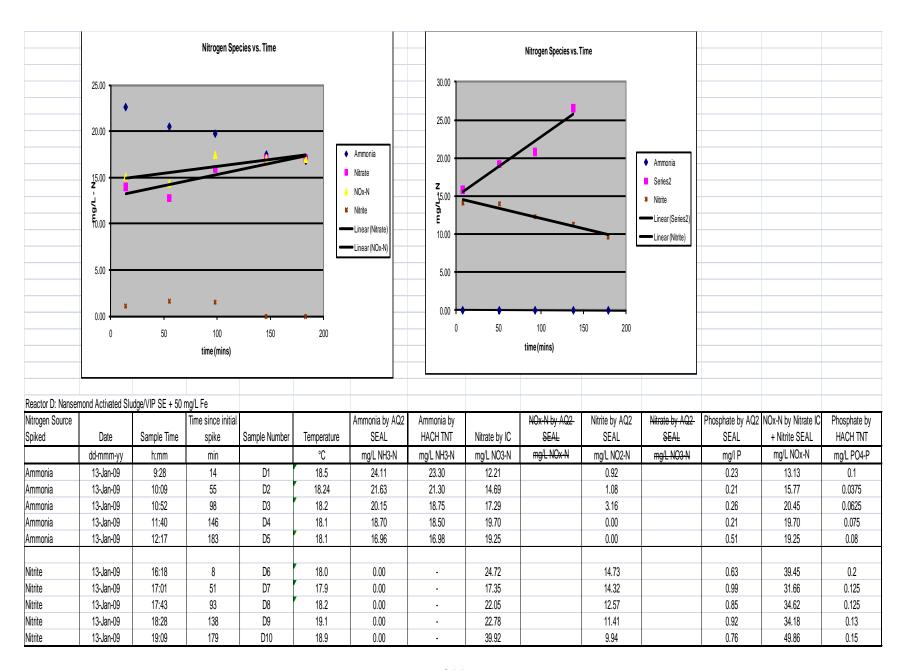
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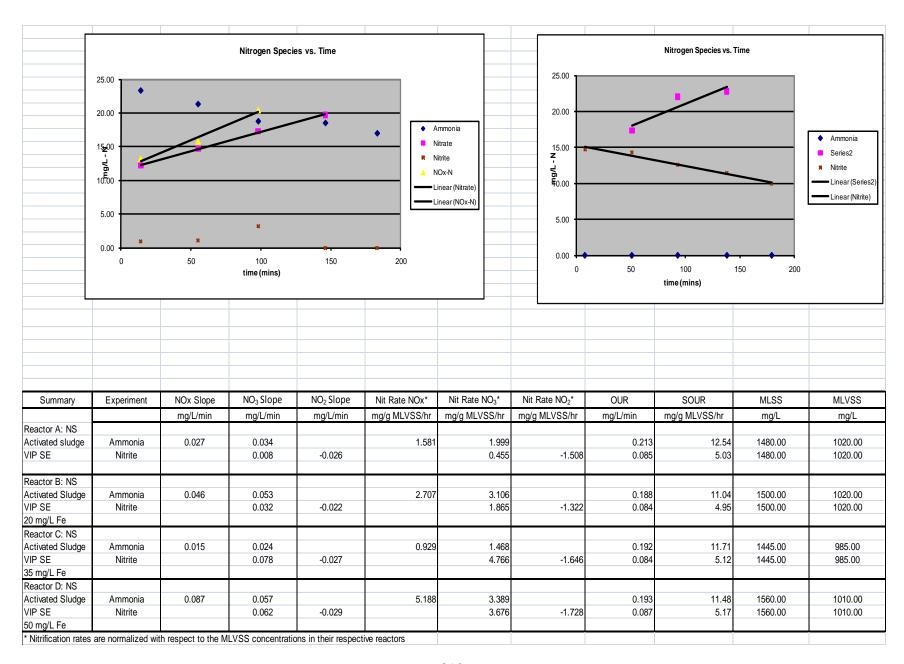
1.9

0.00



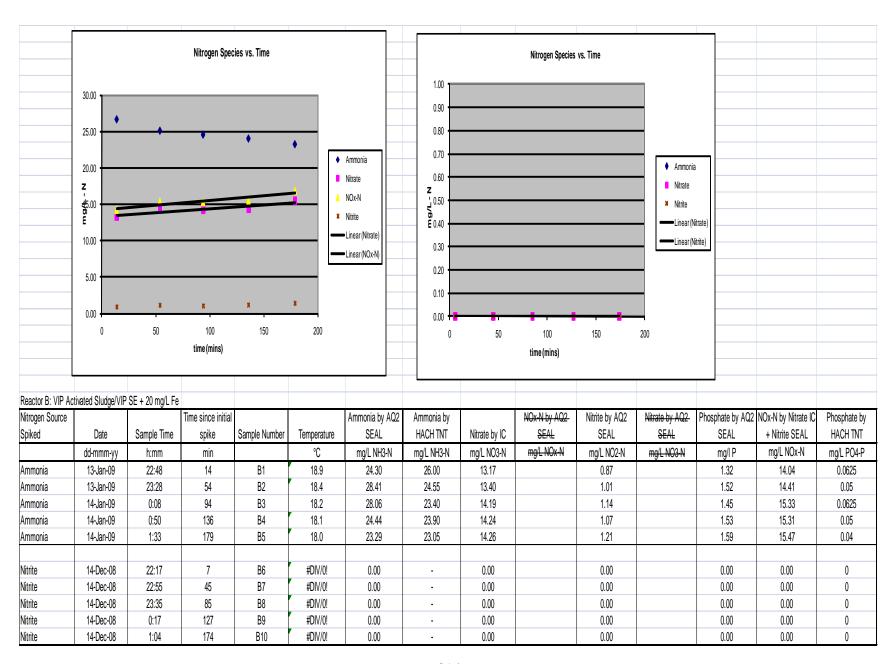


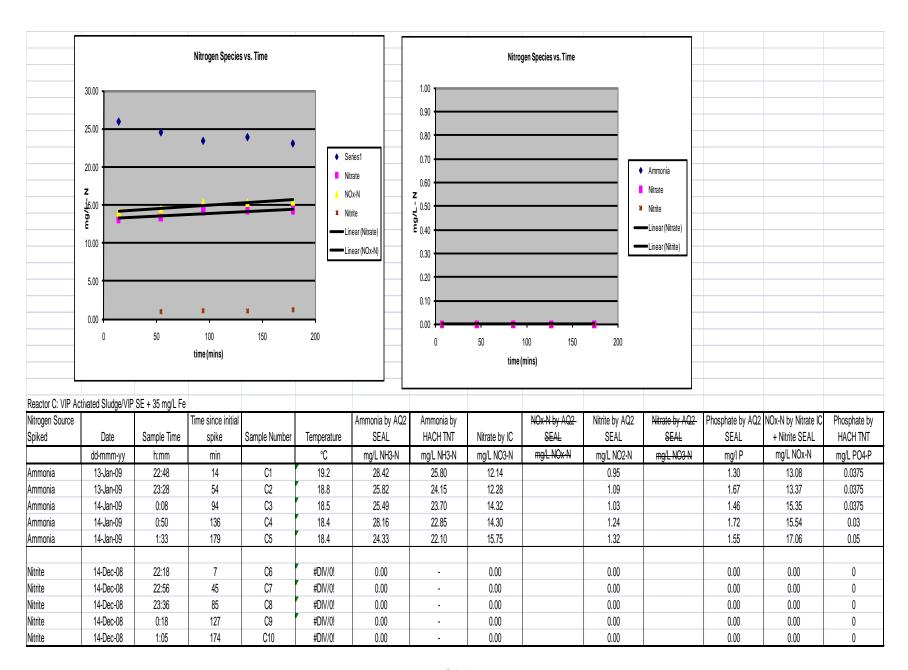


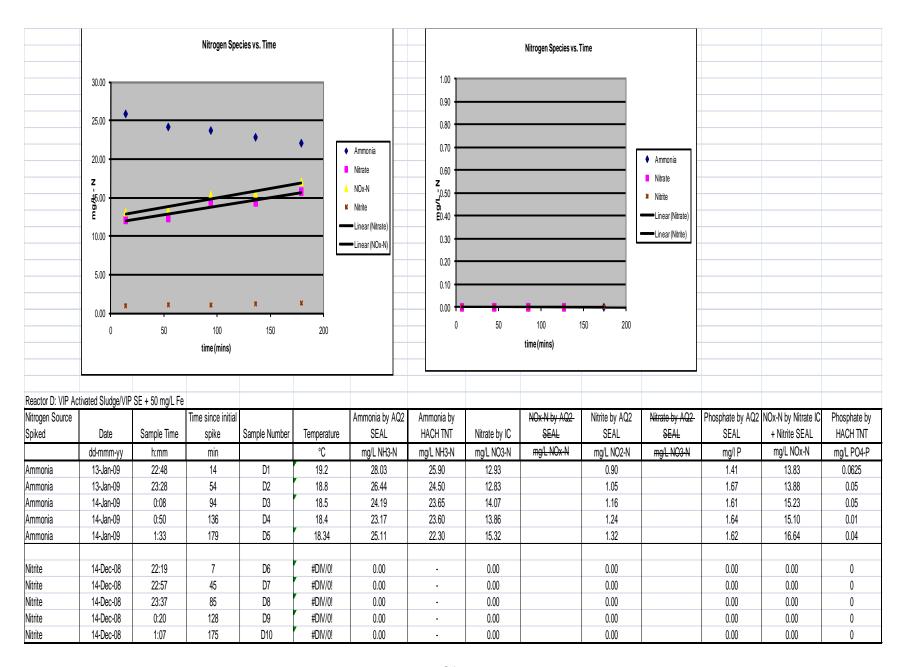


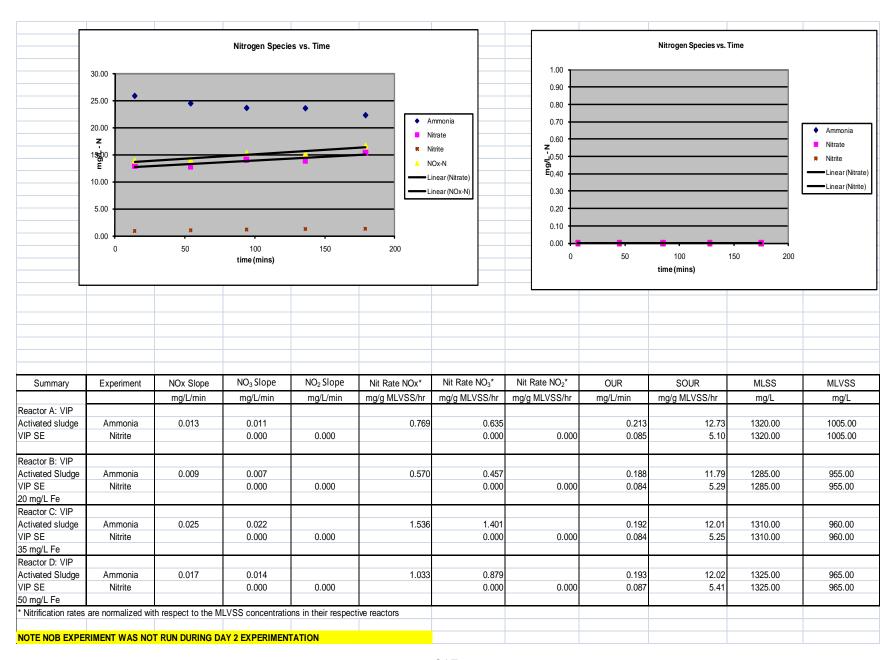
Week 12 – Day 2 AOB/NOB

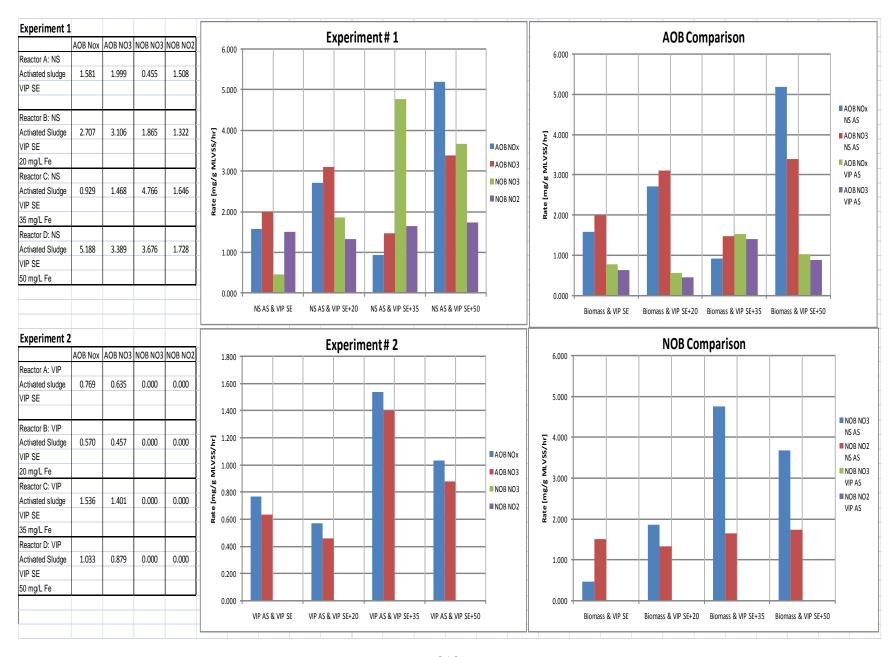
Sample Data F		TARGET OF	1.											
	Report for Nanse	emond Nitrificati	on Inhibition Stu	ldy										
AOB & NOB Ex	perimentation:													
I. AOB	•													
a. Spiked fou	ır 3L reactors with :	20 mg/L NH ₄ .												
'	or is running conti	0, 4	ir bars.											
	f the diluent sourc													
d. 1 L of cond	centrated biomass	is added to the rea	actors. For this we	eks experiments th	ne biomass was co	ncentrated from 20 L	to 9 L.			NOTE NOD EVDED	IMPUT WAS NOT D	UNI DUDINO DAVO E	VDEDIMENTATION	
e. Constant [OO and pH were m	onitored and logge	ed throughout the	experiment.						NOTE NOR EXPER	IMENI WAS NOT R	UN DURING DAY 2 E	EXPERIMENTATION	
				evels through soler	oid valves.									
				ell as program and		olenoid valves.								
						into sample tubes tl	h rough millipore 0.	.45 µm filters						
						ed on an as Fe basis r		•						
				adjustment due to										
•		J	r	,	,									
II. NOB														
a. Spiked fou	r 3L reactors with	25 mg/L NO ₂ - afte	rammonia levels v	were < 1 mg/L NH ₂ -	N which was check	ed through the use	of HACH TNT 831 A	Ammonia						
				sume all present a		Ü								
	or is running conti		· ·											
	O and pH were m	, ,		experiment.										
	•		•	evels through soler	oid valves.									
				ell as program and		olenoid valves.								
		era period ot 3 no	urs through sampl	ling ports on the re-	actors and filtered	linto sample tubes t	hrough millipore 0.	.45 um filters.						
g. Reactors B						l into sample tubes t ed on an as Fe basis (•						
	, C, and D in the di	iluent source incor	porated varying co	oncentrations of Fe	Cl _{3.} calculated base	l into sample tubes t ed on an as Fe basis I		•						
	, C, and D in the di	iluent source incor	porated varying co		Cl _{3.} calculated base			•						
	, C, and D in the di	iluent source incor	porated varying co	oncentrations of Fe	Cl _{3.} calculated base			•						
h. A two hou	, C, and D in the di r period was allow	iluent source incor ed after FeCl ₃ add	porated varying co	oncentrations of Fe	Cl _{3.} calculated base			•						
h. A two hou	, C, and D in the di	iluent source incor ed after FeCl ₃ add	porated varying co	oncentrations of Fe	Cl _{3.} calculated base			•						
h. A two hou	, C, and D in the di r period was allow	iluent source incor ed after FeCl ₃ add	porated varying co	oncentrations of Fe	Cl _{3.} calculated base			•						
h. A two hou	, C, and D in the di r period was allow	iluent source incor ed after FeCl ₃ add	porated varying co	oncentrations of Fe	Cl _{3.} calculated base			•	NOx N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2	NOx-N by Nitrate IC	Phosphate by
h. A two hou	, C, and D in the di r period was allow	iluent source incor ed after FeCl ₃ add	porated varying co ition to monitor pl	oncentrations of Fe	Cl _{3.} calculated base	ed on an as Fe basis (relative to the 1L B	•	NOx-N by AQ2- SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2- SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL	Phosphate by HACH TNT
h. A two hou Reactor A: VIP A	r, C, and D in the di r period was allow ctivated Sludge/VIF	iluent source incor ed after FeCl ₃ add	porated varying co ition to monitor pl	oncentrations of Fe H adjustment due t	Cl ₃ calculated bass o FeCl ₃ addition	ed on an as Fe basis of Ammonia by AQ2	relative to the 1L B Ammonia by	iomass			,			
h. A two hou Reactor A: VIP Ar litrogen Source Spiked	, C, and D in the di r period was allow ctivated Sludge/VIF	iluent source incor ed after FeCl ₃ add SE Sample Time	porated varying co ition to monitor pl Time since initial spike	oncentrations of Fe H adjustment due t	Cl ₃ calculated base o FeCl ₃ addition Temperature	Ammonia by AQ2	relative to the 1L B Ammonia by HACH TNT	iomass Nitrate by IC	SEAL	SEAL	SEÁL	SEAL	+ Nitrite SEAL	HACH TNT
h. A two hou Reactor A: VIP Ar Ilitrogen Source ipiked	, C, and D in the di r period was allow ctivated Sludge/VIF Date dd-mmm-yy	iluent source incor ed after FeCl ₃ add SE Sample Time h:mm	porated varying co ition to monitor pl Time since initial spike min	oncentrations of Fe H adjustment due t Sample Number	Cl ₃ calculated base o FeCl ₃ addition Temperature °C	Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT	Nitrate by IC	SEAL	SEAL mg/L NO2-N	SEÁL	SEAL mg/l P	+ Nitrite SEAL mg/L NOx-N	HACH TNT mg/L PO4-P
h. A two hou Reactor A: VIP Ar Relationary American Source Repiked	, C, and D in the di r period was allow ctivated Sludge/VIF Date dd-mmm-yy 13-Jan-09	iluent source incor ed after FeCl ₃ add SEE Sample Time h:mm 22:48	porated varying co ition to monitor pl Time since initial spike min 14	oncentrations of Fe H adjustment due t Sample Number	Cl ₃ calculated base o FeCl ₃ addition Temperature °C 18.9	Ammonia by AQ2 SEAL mg/L NH3-N 27.86	Ammonia by HACH TNT mg/L NH3-N 26.70	Nitrate by IC mg/L NO3-N 13.31	SEAL	SEAL mg/L NO2-N 0.94	SEÁL	SEAL mg/l P 1.41	+ Nitrite SEAL mg/L NOx-N 14.25	HACH TNT mg/L PO4-P 0.075
h. A two hou Reactor A: VIP Ar Ilitrogen Source Epiked Immonia Immonia Immonia	, C, and D in the di r period was allow ctivated Sludge/VIF Date dd-mmm-yy 13-Jan-09 13-Jan-09	sample Time h:mm 22:48 23:28	porated varying co ition to monitor ph Time since initial spike min 14 54	Sample Number A1 A2	Cl ₂ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3	Ammonia by AQ2 SEAL mg/L NH3-N 27.86 27.90	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10	Nitrate by IC mg/L NO3-N 13.31 14.25	SEAL	SEAL mg/L NO2-N 0.94 1.09	SEÁL	SEAL mg/l P 1.41 1.80	+ Nitrite SEAL mg/L NOx-N 14.25 15.33	MACH TNT mg/L PO4-P 0.075 0.075
h. A two hou Reactor A: VIP Ar Ilitrogen Source Epiked Immonia Immonia Immonia Immonia Immonia	, C, and D in the di r period was allow ctivated Sludge/VIP Date dd-mmm-yy 13-Jan-09 14-Jan-09	sample Time h:mm 22:48 23:28 0:08	porated varying co ition to monitor ph Time since initial spike min 14 54 94	Sample Number A1 A2 A3	Cl ₂ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3	Ammonia by AQ2 SEAL mg/L NH3-N 27.86 27.90 25.19	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10 24.55	Nitrate by IC mg/L NO3-N 13.31 14.25 14.25	SEAL	SEAL mg/L NO2-N 0.94 1.09 1.03	SEÁL	SEAL mg/l P 1.41 1.80 1.58	+ Nitrite SEAL mg/L NOx-N 14.25 15.33 15.28	HACH TNT mg/L PO4-P 0.075 0.075 0.075
h. A two hou teactor A: VIP Ar litrogen Source piked mmonia mmonia mmonia mmonia	, c, and D in the di r period was allow ctivated Sludge/VIP Date dd-mmm-yy 13-Jan-09 14-Jan-09 14-Jan-09	sample Time h:mm 22:48 23:28 0:08 0:50	porated varying co ition to monitor ph Time since initial spike min 14 54 94 136	Sample Number A1 A2 A3 A4	Cl ₂ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3 18.1	Ammonia by AQ2 SEAL mg/L NH3-N 27.86 27.90 25.19 23.57	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10 24.55 24.00	Nitrate by IC mg/L NO3-N 13.31 14.25 14.25 14.35	SEAL	SEAL mg/L NO2-N 0.94 1.09 1.03 1.18	SEÁL	SEAL mg/l P 1.41 1.80 1.58 1.71	+ Nitrite SEAL mg/L NOx-N 14.25 15.33 15.28 15.54	HACH TNT mg/L PO4-P 0.075 0.075 0.075 0.100
h. A two hou leactor A: VIP Ar litrogen Source piked mmonia mmonia mmonia mmonia mmonia mmonia	, c, and D in the di r period was allow ctivated Sludge/VIP Date dd-mmm-yy 13-Jan-09 14-Jan-09 14-Jan-09	sample Time h:mm 22:48 23:28 0:08 0:50	porated varying co ition to monitor pl Time since initial spike min 14 54 94 136 179	Sample Number A1 A2 A3 A4 A5	Cl ₂ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3 18.1	Ammonia by AQ2 SEAL mg/L NH3-N 27.86 27.90 25.19 23.57	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10 24.55 24.00	Nitrate by IC mg/L NO3-N 13.31 14.25 14.25 14.35	SEAL	SEAL mg/L NO2-N 0.94 1.09 1.03 1.18	SEÁL	SEAL mg/l P 1.41 1.80 1.58 1.71	+ Nitrite SEAL mg/L NOx-N 14.25 15.33 15.28 15.54	HACH TNT mg/L PO4-P 0.075 0.075 0.075 0.100
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h. A two hou Reactor A: VIP Ar ditrogen Source spiked Ammonia Ammonia Ammonia Ammonia Altirite ditrite	, C, and D in the di r period was allow ctivated Sludge/VIP Date dd-mmm-yy 13-Jan-09 14-Jan-09 14-Jan-09 14-Jan-09 14-Dec-08 14-Dec-08 14-Dec-08	Sample Time	porated varying co ition to monitor pl Time since initial spike min 14 54 94 136 179 6 45	Sample Number A1 A2 A3 A4 A5 A6 A7 A8	Cl ₃ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3 18.1 18.1 18.0 #DIV/0! #DIV/0! #DIV/0!	Ammonia by AQ2 SEAL mg/L NH3-N 27.90 25.19 23.57 25.84	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10 24.55 24.00	Nitrate by IC mg/L NO3-N 13.31 14.25 14.25 14.35 15.44 0.00 0.00 0.00	SEAL	SEAL mg/L NO2-N 0.94 1.09 1.03 1.18 1.35 0.00 0.00 0.00	SEÁL	SEAL mg/l P 1.41 1.80 1.58 1.71 1.54 0.00 0.00 0.00	+ Nitrite SEAL mg/L NOx-N 14.25 15.33 15.28 15.54 16.79 0.00 0.00 0.00	HACH TNT mg/L PO4-P 0.075 0.075 0.075 0.075 0.100 0.063
h. A two hou Reactor A: VIP A: ditrogen Source tipiked Ammonia Ammonia Ammonia Ammonia Altirite ditrite	, C, and D in the di r period was allow ctivated Sludge/VIP Date dd-mmm-yy 13-Jan-09 14-Jan-09 14-Jan-09 14-Jan-09 14-Jan-09 14-Dec-08	Sample Time	Time since initial spike min 14 54 94 136 179 6 45	Sample Number A1 A2 A3 A4 A5 A6 A7	Cl ₃ calculated bass o FeCl ₃ addition Temperature °C 18.9 18.3 18.1 18.1 18.0 #DIV/0!	Ammonia by AQ2 SEAL mg/L NH3-N 27-90 25-19 23-57 25-84	Ammonia by HACH TNT mg/L NH3-N 26.70 25.10 24.55 24.00	Nitrate by IC mg/L NO3-N 13.31 14.25 14.35 15.44 0.00 0.00	SEAL	SEAL mg/L NO2-N 0.94 1.09 1.03 1.18 1.35 0.00 0.00	SEÁL	SEAL mg/l P 1.41 1.80 1.58 1.71 1.54 0.00 0.00	+ Nitrite SEAL mg/L NOx-N 14.25 15.33 15.28 15.54 16.79 0.00 0.00	HACH TNT mg/L PO4-P 0.075 0.075 0.075 0.075 0.100 0.063







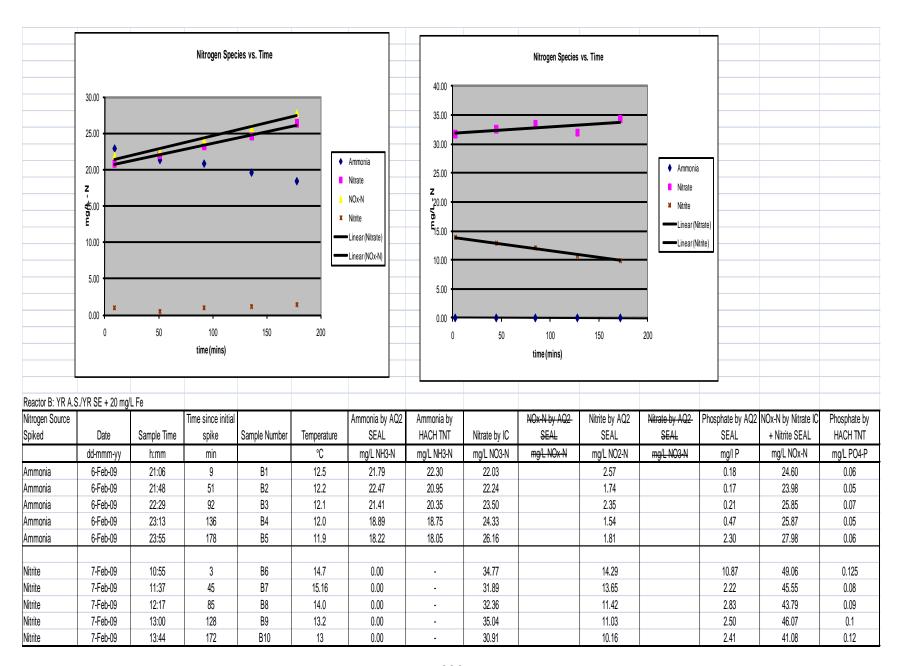


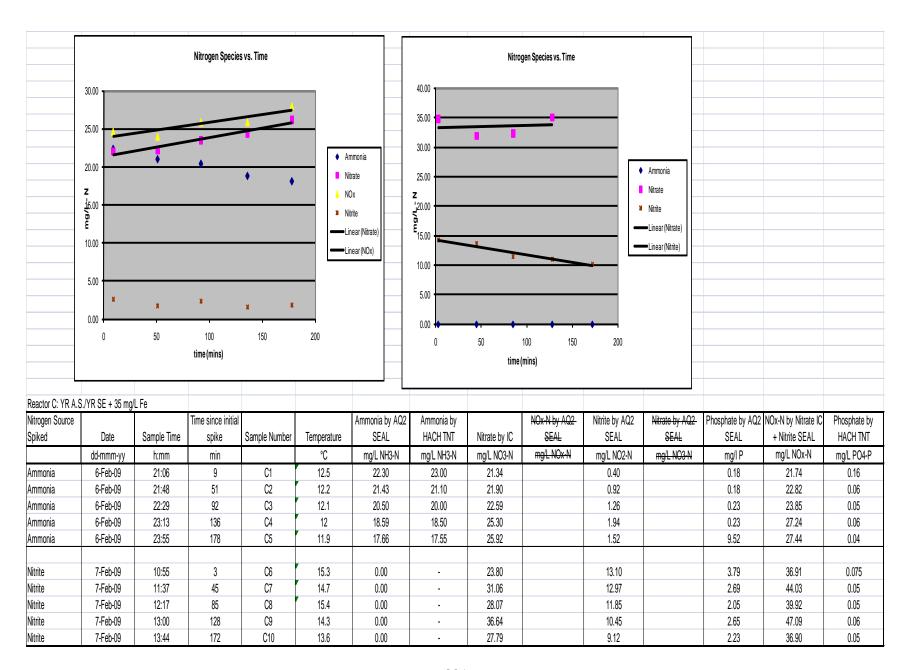


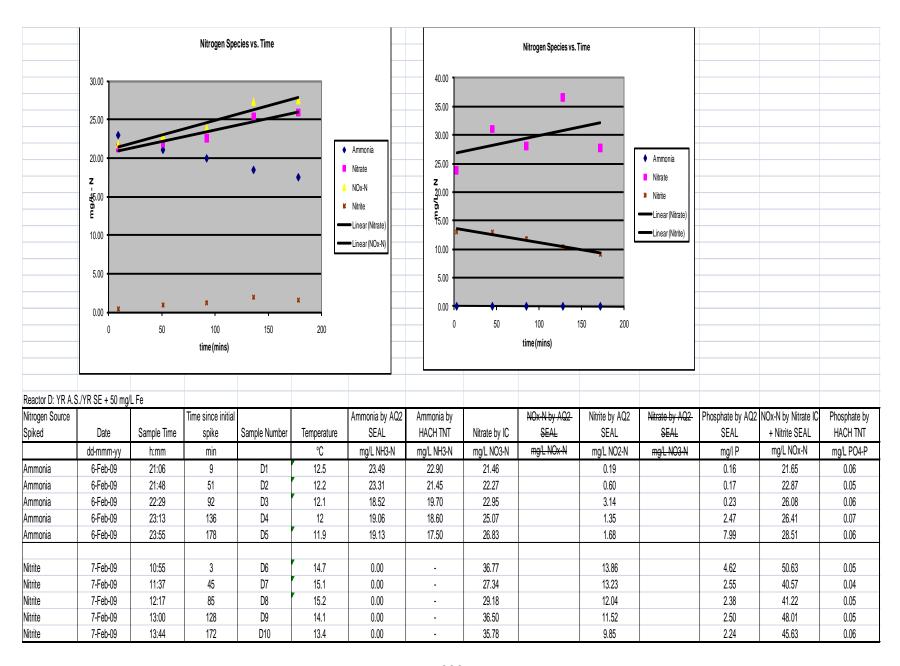
Week 13 – FeCl3 Addition with York River Biomass Day 1 AOB/NOB

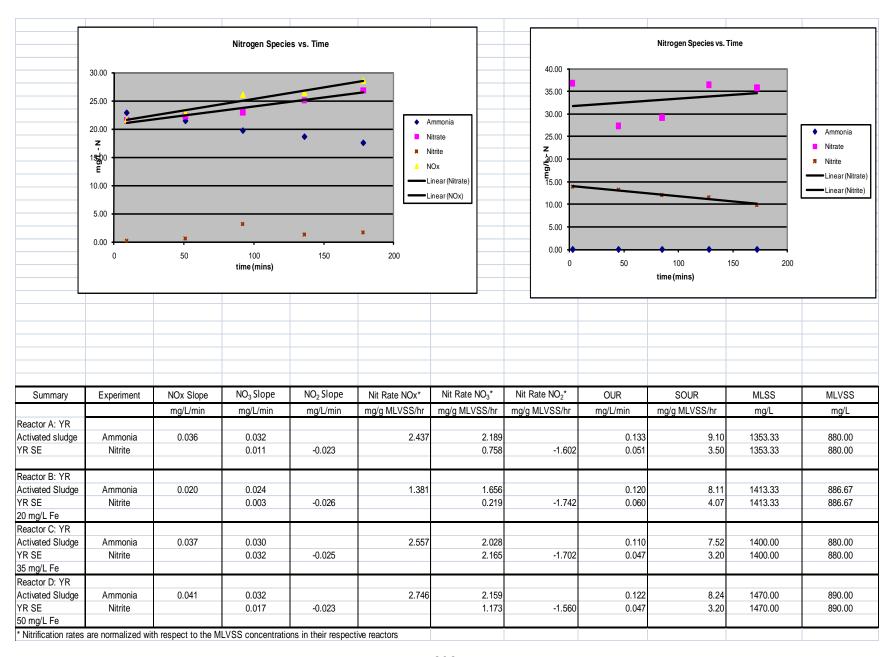
ample Data Report for Nansemond Nitrific	ation Inhibition Stu	dy							
AOB & NOB Experimentation: 1. AOB a. Spiked four 3L reactors with 25 mg/L NH ₄ . b. Each reactor is running continously by use o c. 2 L of the diluent source is added to the rea d. 1 L of concentrated biomass is added to the e. Constant DO and pH were monitored and lo f. Oxygen/air blend was sparged into the reac g. LabView software was used to manage DO a h. 5 samples were collected over a period of 3 i. Reactors B, C, and D in the diluent source in j. A two hour period was allowed after FeCl ₃ ac k. Alkalinity was added to each reactor accord	tors. eactors. For this we ged throughout the ors to maintain DO le hd pH recording as w nours through sampl orporated varying codition to monitor pH	experiment. evels through sol rell as program a ing ports on the oncentrations of adjustment due	enoid valves. nd implement the sectors and filtere FeCl ₃ calculated bar to FeCl ₃ addition.	olenoid valves. d into sample tubes sed on an as Fe basis	through millipore 0	•			
II. NOB a. Spiked four 3L reactors with 15 mg/L NO ₂ · at a paddition all reactors were allowed to run over b. Each reactor is running continously by use o c. Constant DO and pH were monitored and loid. Oxygen/air blend was sparged into the react e. LabView software was used to manage DO a f. 5 samples were collected over a period of 3 g. Reactors B, C, and D in the diluent source inch. A two hour period was allowed after FeCl ₃ a i. Alkalinity was added to each reactor according	night to consume all stir bars. ged throughout the ors to maintain DO le and pH recording as w nours through sampl orporated varying co Idition to monitor pl	present ammon experiment. evels through sol rell as program a ing ports on the oncentrations of H adjustment du	enoid valves. Ind implement the sereactors and filtere FeCl ₃ calculated bare to FeCl ₃ addition.	olenoid valves. d into sample tubes sed on an as Fe basis	through millipore 0	45 μm filters.			
actor A: YR A.S. / YR SE									

Reactor A: YR A.	S. / YR SE													
Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOx-N by AQ2 SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2 SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L-NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	6-Feb-09	21:06	9	A1	12.7	23.46	22.90	20.84		1.01		0.27	21.85	0.18
Ammonia	6-Feb-09	21:48	51	A2	12.5	21.75	21.30	21.99		0.54		0.24	22.53	0.14
Ammonia	6-Feb-09	22:29	92	A3	12.3	20.66	20.80	23.21		0.95		0.24	24.16	0.09
Ammonia	6-Feb-09	23:13	136	A4	12.2	20.06	19.55	24.54		1.21		0.28	25.75	0.11
Ammonia	6-Feb-09	23:55	178	A5	12.1	16.84	18.35	26.36		1.44		10.17	27.80	0.13
Nitrite	7-Feb-09	10:55	3	A6	14.9	0.00	-	31.75		13.85		10.47	45.60	0.175
Nitrite	7-Feb-09	11:37	45	A7	15.35	0.00	-	32.56		12.87		2.00	45.42	0.15
Nitrite	7-Feb-09	12:17	85	A8	14.1	0.00	-	33.39		12.13		2.55	45.51	0.14
Nitrite	7-Feb-09	13:00	128	A9	13.3	0.00	-	31.98		10.77		2.29	42.75	0.14
Nitrite	7-Feb-09	13:44	172	A10	13.2	0.00	-	34.37		9.96		2.51	44.32	0.14







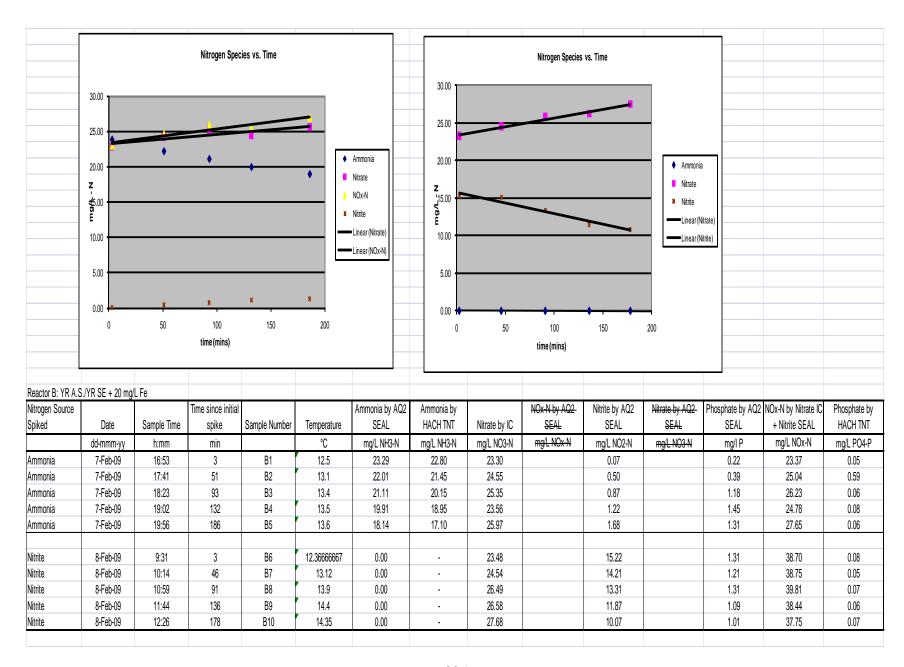


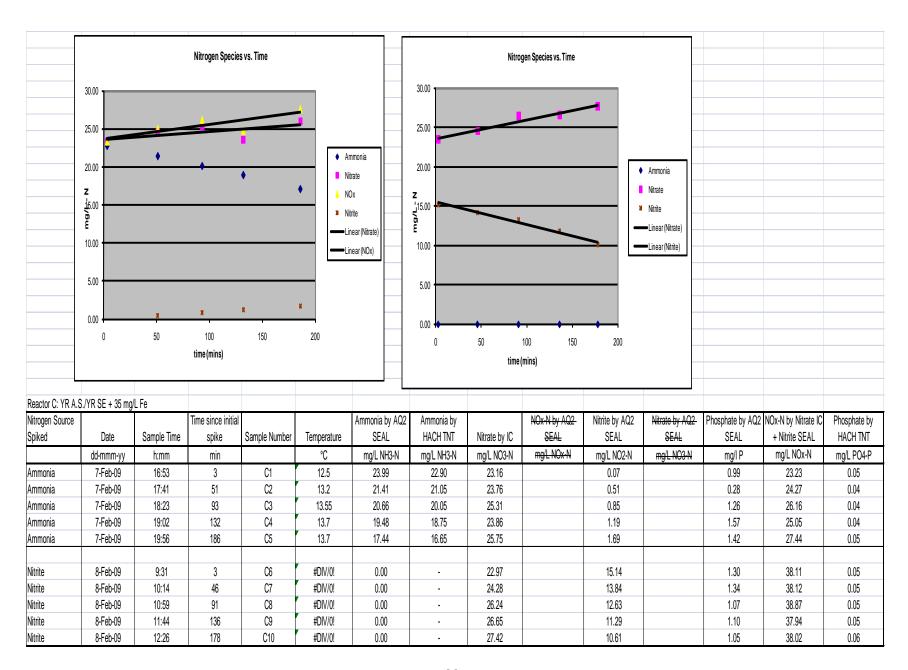
Week 13 – Day 2 AOB/NOB

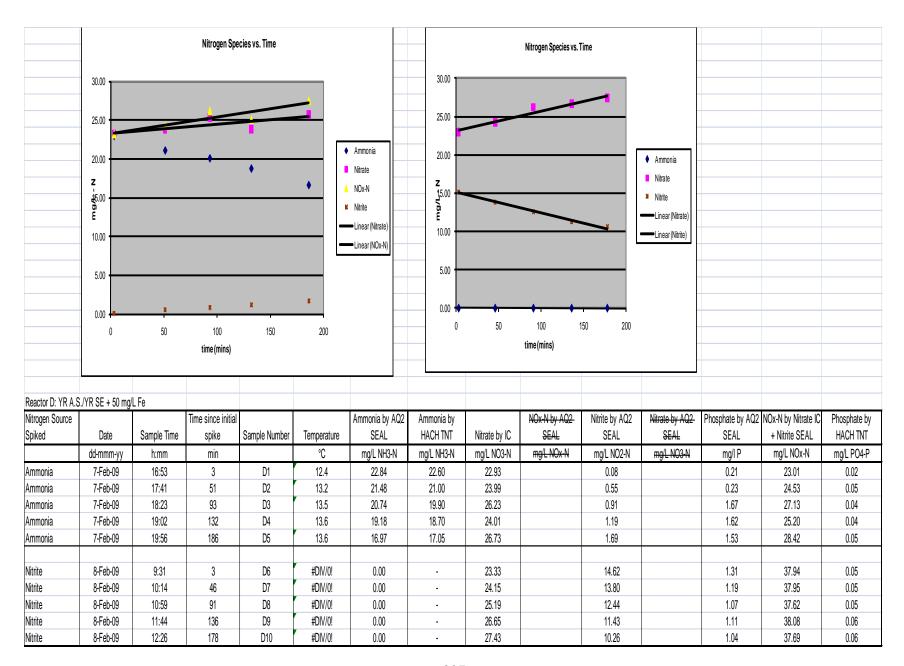
Week 13 - Day 2 AOB/NOB Sample Data Report for Nansemond Nitrification Inhibition Study			
ample Data Report of Natisethora Nithingation Study			
AOB & NOB Experimentation:			
I.AOB			
a. Spiked four 3L reactors with 25 mg/L NH ₄ .			
b. Each reactor is running continously by use of stir bars .			
c. 2 L of the diluent source is added to the reactors.			
d.1Lofconcentratedbiomassisaddedtothereactors.Forthisweeksexperimentsthebiomasswasconcentratedfrom20Lto9L			
e. Constant DO and pH were monitored and logged throughout the experiment.			
f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.			
g. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.			
h. 5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 μ m filters			
i. Reactors B, C, and D in the diluent source incorporated varying concentrations of FeCl ₃ calculated based on an as Fe basis relative to the Biomass			
j. A two hour period was allowed after $FeCl_3$ addition to monitor pH adjustment due to $FeCl_3$ addition.			
k. Alkalinity was added to each reactor according to the Fe dose added to keep the pH from dropping to low.			
II.NOB			
a. Spiked four 3L reactors with 15 mg/L NO ₂ after ammonia levels were <1 mg/L NH ₃ -N which was checked through the use of HACH TNT 831 Ammonia method.			
In addition all reactors were allowed to run overnight to consume all present ammonia.			
b. Each reactor is running continously by use of stir bars.			
c. Constant DO and pH were monitored and logged throughout the experiment.			
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.			
e. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.			
f. 5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 μm filters. g. Reactors B, C, and D in the diluent source incorporated varying concentrations of FeCl ₃ calculated based on an as Fe basis relative to thet. Biomass			
b. A two hour period was allowed after FeCl ₃ addition to monitor pH adjustment due to FeCl ₃ addition.			
i. Alkalinity was added to each reactor according to the Fe dose added to keep the pH from dropping to low.			
i. Aikallility was added to each reactor according to the re dose added to keep the printfull dropping to low.			

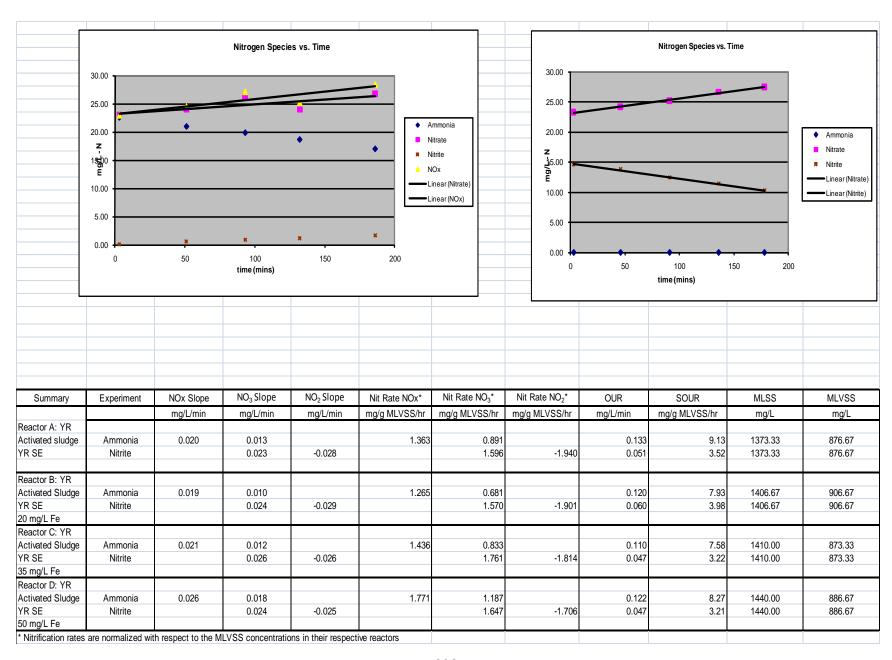
Reactor	Δ.	ΥR	Δ	ς	/YR	SF

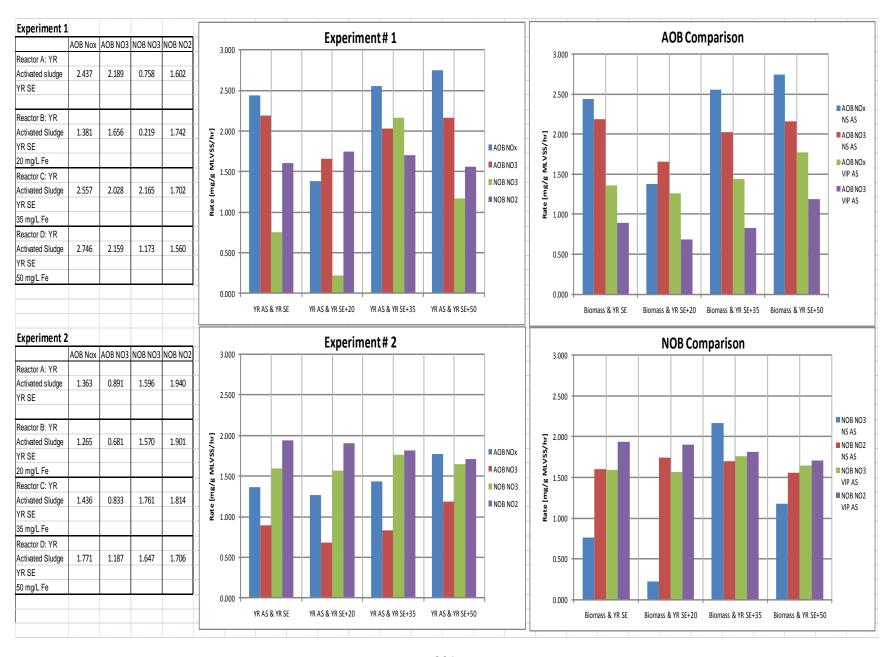
NEAUTOL A. TIN A.C)./ TIN OL													
Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOx-N by AQ2 SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2 SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L-NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	7-Feb-09	16:53	3	A1	12.5	24.52	23.90	22.90		0.07		0.31	22.96	0.16
Ammonia	7-Feb-09	17:41	51	A2	13.1	22.25	22.25	24.19		0.44		3.72	24.63	0.13
Ammonia	7-Feb-09	18:23	93	A3	13.3	22.25	21.15	25.15		0.78		0.30	25.92	0.15
Ammonia	7-Feb-09	19:02	132	A4	13.4	20.24	20.00	24.49		1.08		1.43	25.57	0.15
Ammonia	7-Feb-09	19:56	186	A5	13.5	20.87	19.00	25.62		1.30		1.43	26.91	0.14
Nitrite	8-Feb-09	9:31	3	A6	12.4	0.00		23.27		15.30		1.29	38.57	0.13
Nitrite	8-Feb-09	10:14	46	A7	12.9	0.00	-	24.53		15.12		1.27	39.65	0.16
Nitrite	8-Feb-09	10:59	91	A8	14.0	0.00	-	25.87		13.39		1.27	39.26	0.15
Nitrite	8-Feb-09	11:44	136	A9	14.4	0.00	-	26.27		11.50		1.25	37.78	0.16
Nitrite	8-Feb-09	12:26	178	A10	14.4	0.00	-	27.53		10.88		1.05	38.41	0.15



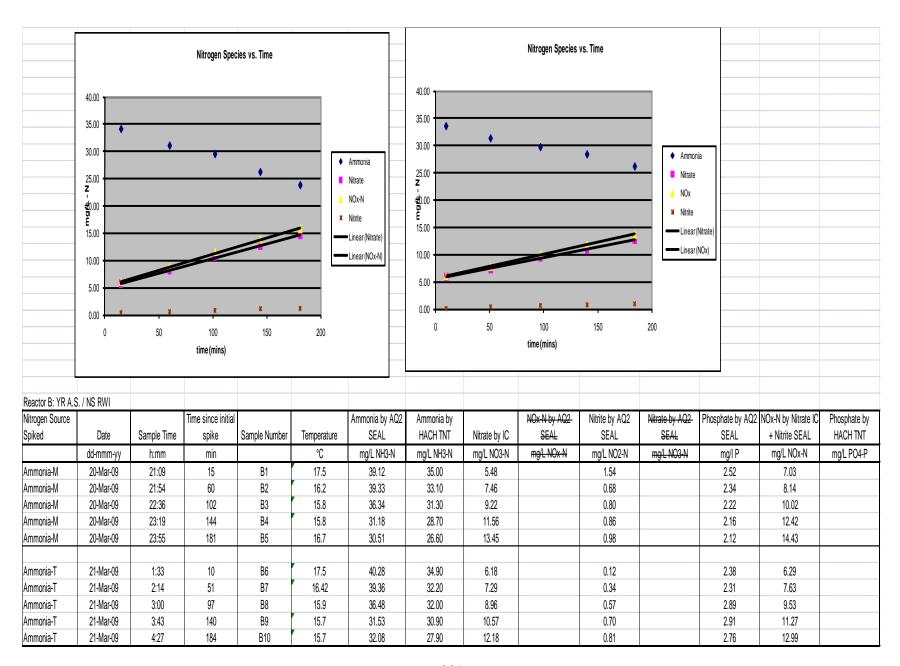


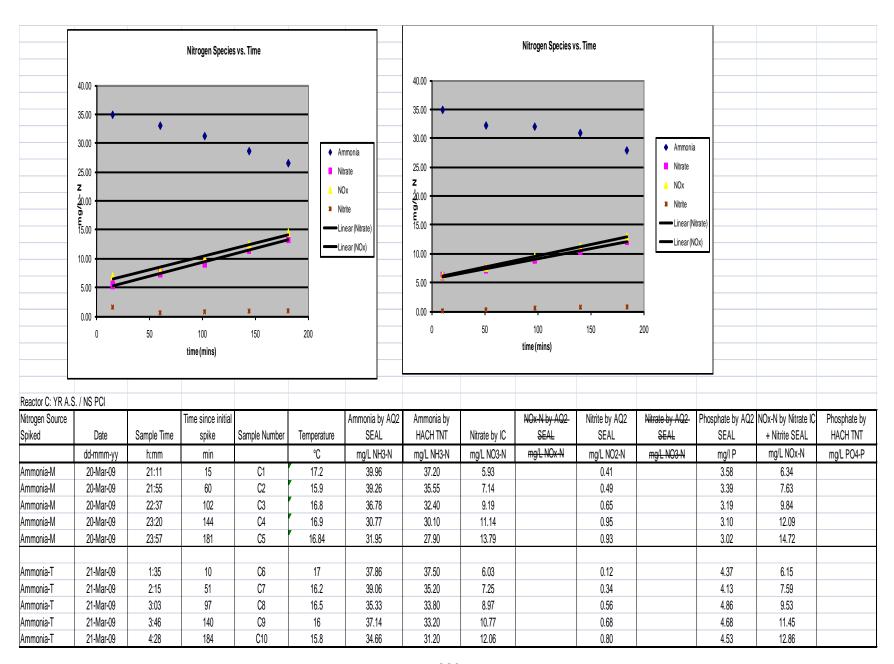


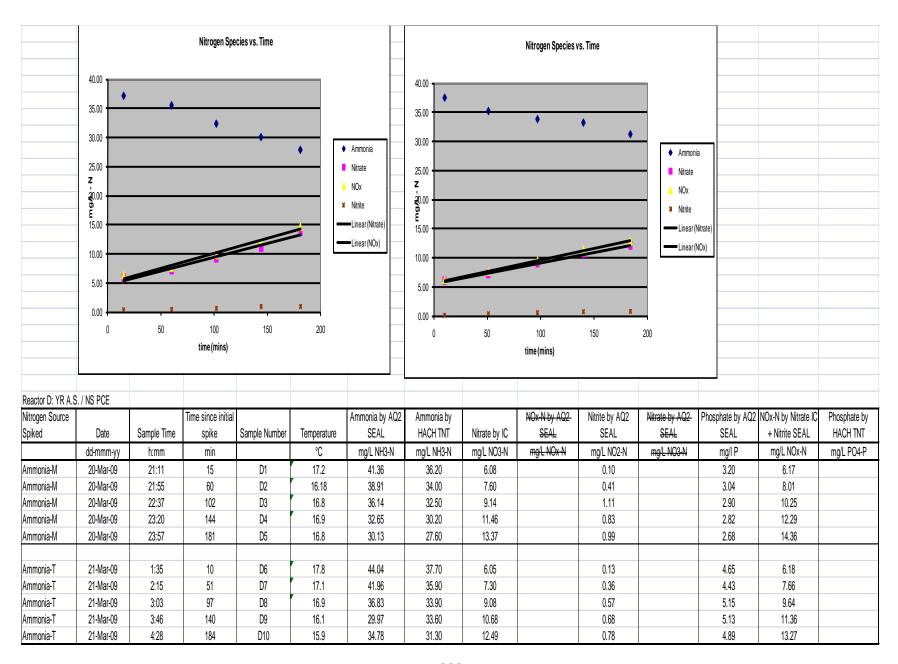


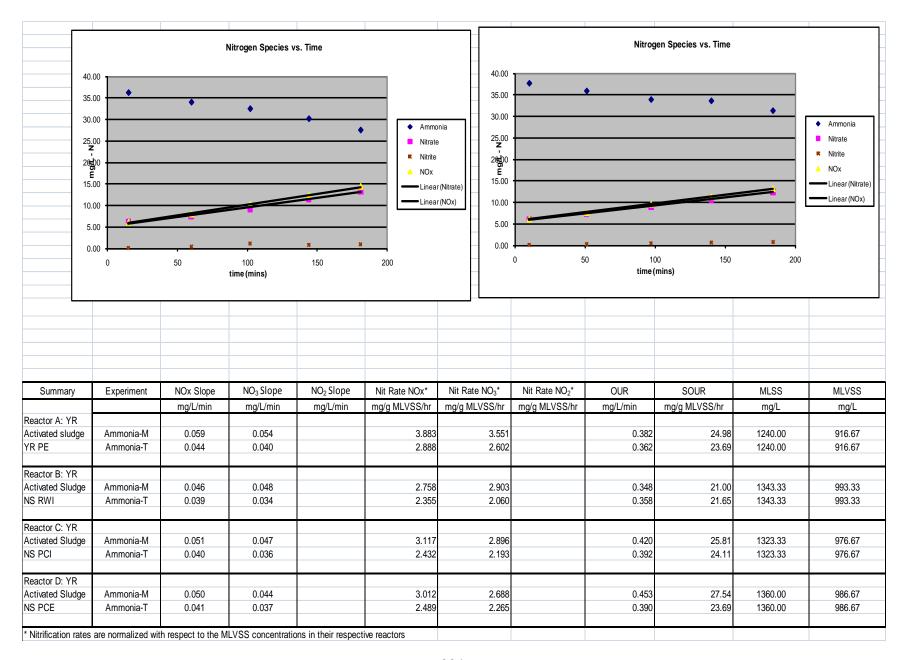


Date Sample Time Spike Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL HACH TNT	Week 14	-NTP	Raw, PC	I, PCE (Composit	te Sampl	es Day 1	AOB							
1.00 Specific or 1.20 Specific or 2.20	Sample Data R	Report for Nanse	emond Nitrificati	on Inhibition Stu	ıdy	_									
1.008															
3.5)Red for SI. resultives with 25 mig/L New, 1	AOB Experime	ntation:													
b. Each restor's naming continously lyoue of sir bars. 2.1 of the fallations used sadded to the meastors. For this week segariments the bismass was concentrated from 201 to 9 L. e. Constant DD and pit were monitored and logged throughout the experiment. D. Oxygen/air belled was supped into the restories to maintain DD week through scleracid valves. g. Labifives with swell was some and the restories to maintain DD wheek through scleracid valves. g. Labifives with swell was some and the restories to maintain DD with suppersional valves. g. Labifives with swell was some incorporated 22 of the associated composite samples taken for each day respectively. I. Reactors B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE	I. AOB														
b. Each restor's naming continously lyoue of sir bars. 2.1 of the fallations used sadded to the meastors. For this week segariments the bismass was concentrated from 201 to 9 L. e. Constant DD and pit were monitored and logged throughout the experiment. D. Oxygen/air belled was supped into the restories to maintain DD week through scleracid valves. g. Labifives with swell was some and the restories to maintain DD wheek through scleracid valves. g. Labifives with swell was some and the restories to maintain DD with suppersional valves. g. Labifives with swell was some incorporated 22 of the associated composite samples taken for each day respectively. I. Reactors B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE Reactor A, YR A, S. / YR PE Reactor B, C, and D in the dilutent source incorporated 22 of the associated composite samples taken for each day respectively. Reactor A, YR A, S. / YR PE	a. Spiked four	r 3L reactors with	25 mg/L NH₄.												
A 1.1 of concentrated biomas is added to the reactors. For this week superiments be biomass was concentrated from 20.1 to 9.1. Construct Op and were monitored and year emonitored and boust strongs ampling part of the reactors to maintain DOI levels through solenoid valves. 1. Ongen/artikend was spaged into the reactors to maintain DOI levels through solenoid valves. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporated 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporate 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporate 2L of the associated composite samples taken for each day respectively. 2. Exercise S. C. and D in the dissent source incorporate 2L of the associated compo				ir bars.											
L Constant XO and pit were monitored and logged throughpout the experiment. 1. Changes of 2 hours strong in a control of the sector on an air 20 May 10 May 12 May	c. 2 L of the d	iluent source is ad	lded to the reacto	rs.											
1. Oxygen/fair bend was spaged into the reactors to maintain DO loves through sale not wives. Like actions 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporated 2. of the associated composite samples taken for each day respectively. Reactors 8, C, and D in the dilutent source incorporate by ACQ. Reactors 9, Notice by ACQ. Reactors 9, No	d. 1 L of conc	entrated biomass	is added to the rea	actors. For this we	eks experiments t	he biomass was co	ncentrated from 20 l	. to 9 L.							
g LaiDives workname was used to manage CD and pit recording as well as program and implement the velocid valves. h. 5 samples were collected over a period of 3 hours through sampling port on the reactors and filtered into sample tubes through millipore 0.45 µm filters. I. Reactors 8, C, and D in the diluent source incorporated 2. of the associate of composite samples taken for each day respectively. It is a second of the composite samples provided in the composite samples taken for each day respectively. It is a second of the composite samples provided in the composite samples taken for each day respectively. It is a second of the composite samples provided in the composite samples taken for each day respectively. It is a second of the composite samples provided in the composite samples taken for each day respectively. It is a second of the composite samples provided in the composite samples taken for each day respectively. It is a second of the composite samples provided in the composition of the	e. Constant D	O and pH were m	onitored and logge	ed throughout the	experiment.										
I. Reactor 8, C, and D in the diluent source incorporated 2, of the associated composite samples taken for each day respectively. Reactor A, YR A, S, /YR PE	f. Oxygen/ai	r blend was sparge	ed into the reactor	s to maintain DO le	evels through sole	noid valves.									
Reactors B, C, and D in the dilluent source incorporated 2.0 of the associated composite samples taken for each day respectively.	g. LabView so	ftware was used t	to manage DO and	pH recording as w	ell as program and	d implement the so	olenoid valves.								
Reactor A: YR A.S. / YR PE Itingen Source Date Sample Time since initial spike Sample Number Temperature SEAL HACH TNT	h. 5 samples v	were collected ove	er a period of 3 ho	urs through sampl	ing ports on the re	actors and filtered	into sample tubes t	h rough millipore 0.	45 μm filters						
Time since initial spike Sample Time Sample Time Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL SEAL SEAL SEAL HACH TNT Nitrate by IC SEAL S	i. Reactors B,	C, and D in the di	luent source incor	porated 2L of the	associated compos	site samples taken	for each day respect	tively.							
Time since initial spike Sample Time Sample Time Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL SEAL SEAL SEAL HACH TNT Nitrate by IC SEAL S															
Time since initial spike Sample Time Sample Time Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL SEAL SEAL SEAL HACH TNT Nitrate by IC SEAL S															
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Time since initial Sample Time Sample Time Sample Number Temperature SEAL Ammonia by AQ2 Ammonia by HACH TINT Nitrate by IC SEAL															
Time since initial Sample Time Sample Time Sample Number Temperature SEAL Ammonia by AQ2 Ammonia by HACH TINT Nitrate by IC SEAL															
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Time since initial Sample Time Sample Time Sample Number Temperature SEAL HACH TINT Nitrate by IC SEAL	Reactor A: YR A S	S / YR PF													
Date Sample Time Spike Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL SEAL HACH TNT	Trouble Tri														
Date Sample Time Spike Sample Number Temperature SEAL HACH TNT Nitrate by IC SEAL SEAL SEAL SEAL SEAL HACH TNT															
dd-mmr-yy	Nitrogen Source			Time since initial			Ammonia by AQ2						Phosphate by AQ2	NOx-N by Nitrate IC	Phosphate by
Ammonia-M 20-Mar-09 21:09 15 A1 19.7 38.48 34.20 5.71 0.37 0.95 6.08 Ammonia-M 20-Mar-09 21:54 60 A2 17.75 35.30 31.15 8.13 0.61 0.93 8.74 Ammonia-M 20-Mar-09 22:36 102 A3 17.3 33.41 29.60 10.77 0.82 0.90 11.59 Ammonia-M 20-Mar-09 23:18 144 A4 16.55 26.44 26.30 12.78 1.12 0.93 13.90 Ammonia-M 20-Mar-09 23:55 181 A5 17 27.52 23.90 14.66 1.17 1.03 15.83 Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33.50 5.90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77	Spiked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL	HACH TNT
Ammonia-M 20-Mar-09 21:54 60 A2 17.75 35.30 31.15 8.13 0.61 0.93 8.74 Ammonia-M 20-Mar-09 22:36 102 A3 17.3 33.41 29.60 10.77 0.82 0.90 11.59 Ammonia-M 20-Mar-09 23:18 144 A4 16.55 26.44 26.30 12.78 1.12 0.93 13.90 Ammonia-M 20-Mar-09 23:55 181 A5 17 27.52 23.90 14.66 1.17 1.03 15.83 Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33.50 5.90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78		dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia-M 20-Mar-09 22:36 102 A3 17.3 33.41 29.60 10.77 0.82 0.90 11.59 Ammonia-M 20-Mar-09 23:18 144 A4 16.55 26.44 26.30 12.78 1.12 0.93 13.90 Ammonia-M 20-Mar-09 23:55 181 A5 17 27.52 23.90 14.66 1.17 1.03 15.83 Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33.50 5.90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78	Ammonia-M	20-Mar-09	21:09			19.7	38.48	34.20			0.37		0.95		
Ammonia-M 20-Mar-09 23:18 144 A4 16.55 26.44 26.30 12.78 1.12 0.93 13.90 Ammonia-M 20-Mar-09 23:55 181 A5 17 27.52 23.90 14.66 1.17 1.03 15.83 Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33.50 5.90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78	Ammonia-M														
Ammonia-M 20-Mar-09 23:55 181 A5 17 27:52 23:90 14:66 1.17 1.03 15:83 Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33:50 5:90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35:92 31:30 7:33 0.44 1.04 7:77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29:70 9:39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16:3 29:25 28:40 10.98 0.80 1.60 11:78	Ammonia-M					_									
Ammonia-T 21-Mar-09 1:33 10 A6 20.1 36.13 33.50 5.90 0.14 1.03 6.04 Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78						_	-								
Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78	Ammonia-M	20-Mar-09	23:55	181	A5	17	27.52	23.90	14.66		1.17		1.03	15.83	
Ammonia-T 21-Mar-09 2:14 51 A7 18.3 35.92 31.30 7.33 0.44 1.04 7.77 Ammonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Ammonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78	Ammonic T	24 M 00	4.00	40	40	20.4	20.40	22.50	E 00		0.44		4.00	6.04	
Immonia-T 21-Mar-09 3:00 97 A8 16.9 33.04 29.70 9.39 0.69 1.04 10.08 Immonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78						_									
ımmonia-T 21-Mar-09 3:43 140 A9 16.3 29.25 28.40 10.98 0.80 1.60 11.78															
						-									
	Ammonia-T				-	-									

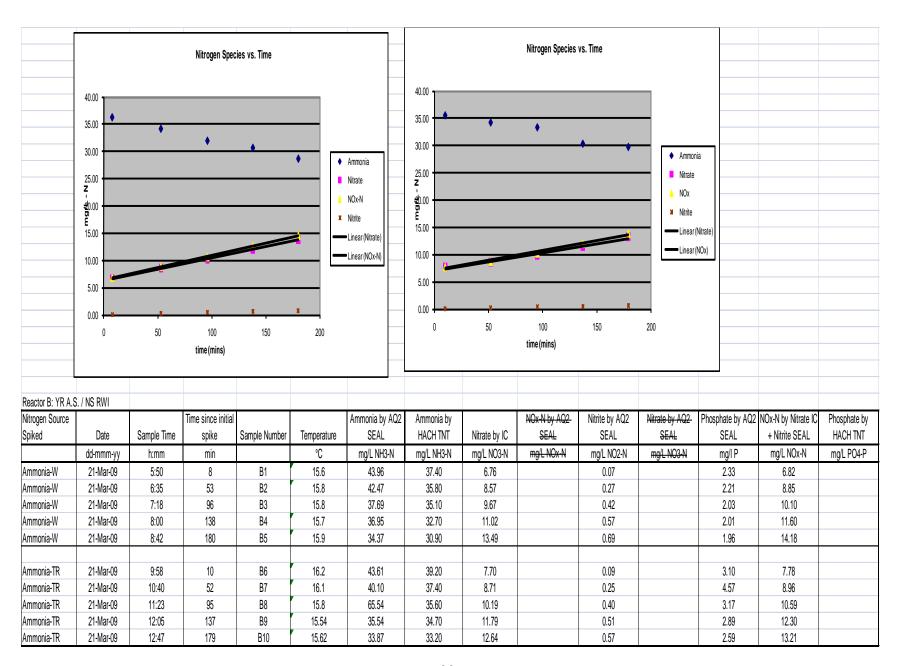


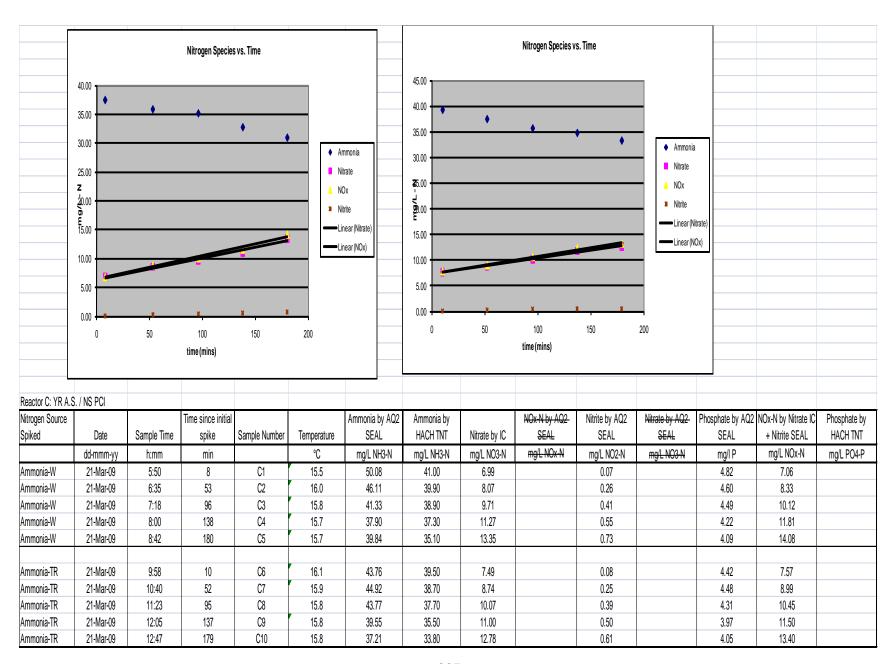


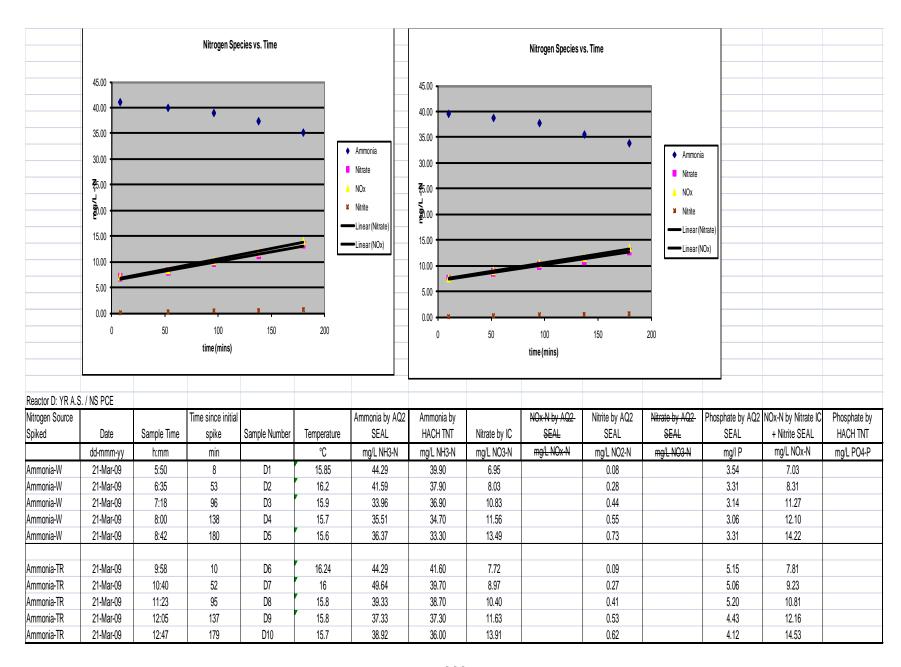


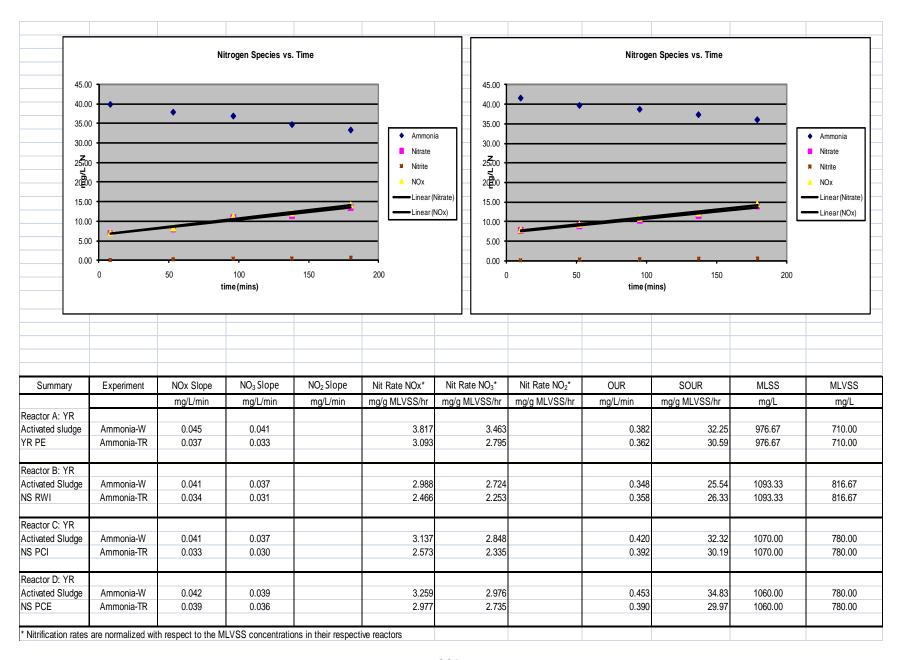


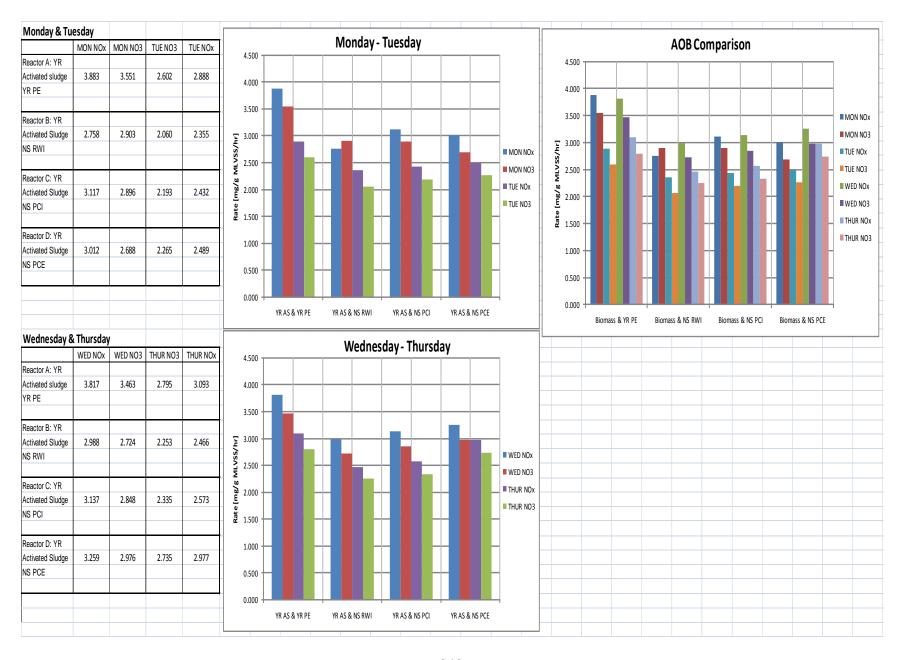
Week 14	- Day 2	2 AOB												
Sample Data R	eport for Nanse	emond Nitrificati	ion Inhibition Stu	ıdy										
400 0 NOC 5														
AOB & NOB Exp	perimentation:													
	r 3L reactors with	25 mg/I NIU												
		25 IIIg/LINH4. inously by use of s	tir harc											
		dded to the react												
				eks eyneriments t	he hinmass was co	ncentrated from 20	I to 9 I							
			ed throughout the		ne biomass was ce	meentrated from 20	L (0 J L.							
			rs to maintain DO I		noid valves.									
			d pH recording as v			olenoid valves.								
						d into sample tubes t	h rough millipore 0.	.45 μm filters						
						for each day respec								
				·	•		·							
eactor A: YR A.S	S. / YR PE													
itrogen Source			Time since initial			Ammonia by AQ2	Ammonia by		NOx-N by AQ2	Nitrite by AQ2	Nitrate by AQ2	Phosphate by AQ2		Phosphate b
piked	Date	Sample Time	spike	Sample Number	Temperature	SEAL	HACH TNT	Nitrate by IC	SEAL	SEAL	SEAL	SEAL	+ Nitrite SEAL	HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-I
mmonia-W	21-Mar-09	5:50	8	A1	17.2	43.17	36.20	6.73		0.07		3.99	6.80	
mmonia-W mmonia-W	21-Mar-09 21-Mar-09	6:35 7:18	53 96	A2 A3	17.1 16.9	37.57 36.42	34.10 31.90	8.59 10.19		0.30 0.49		0.66 0.62	8.89 10.68	
mmonia-W	21-Mar-09	8:00	138	A3 A4	16.5	32.98	30.60	12.00		0.49		0.62	12.64	
mmonia-W	21-Mar-09	8:42	180	A5	15.8	32.08	28.60	13.82		0.80		0.66	14.62	
2000m 0.0										****				
mmonia-TR	21-Mar-09	9:58	10	A6	15.8	37.74	35.60	7.71		0.06		1.72	7.77	
mmonia-TR	21-Mar-09	10:40	52	A7	15.7	37.88	34.30	8.56		0.21		1.43	8.77	
mmonia-TR	21-Mar-09	11:23	95	A8	15.7	33.40	33.40	9.92		0.37		1.60	10.29	
mmonia-TR	21-Mar-09	12:05	137	A9	15.6	35.36	30.40	11.52		0.52		1.58	12.04	
mmonia-TR	21-Mar-09	12:47	179	A10	15.6	33.25	29.80	13.22		0.66		1.21	13.88	







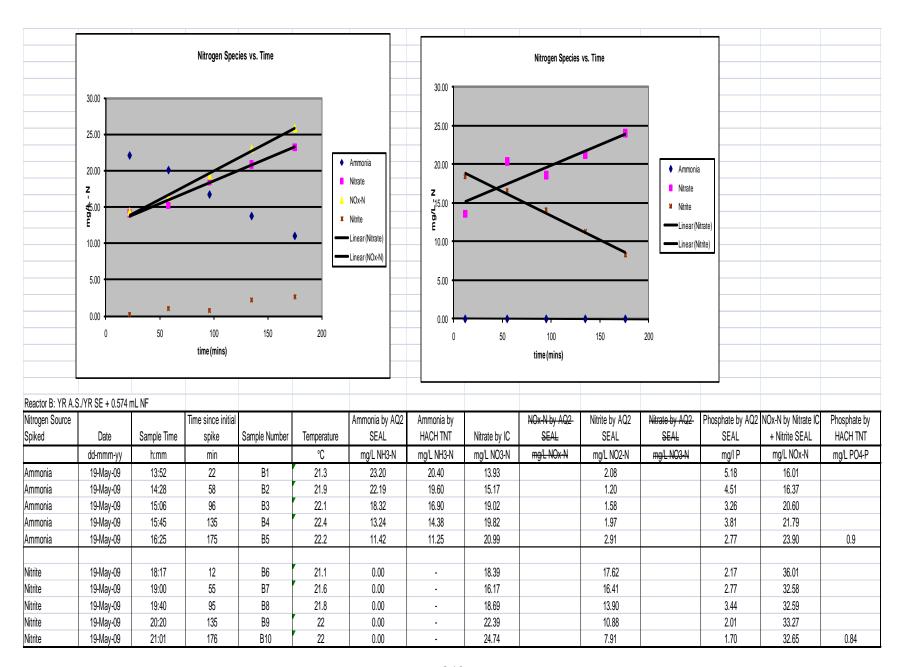


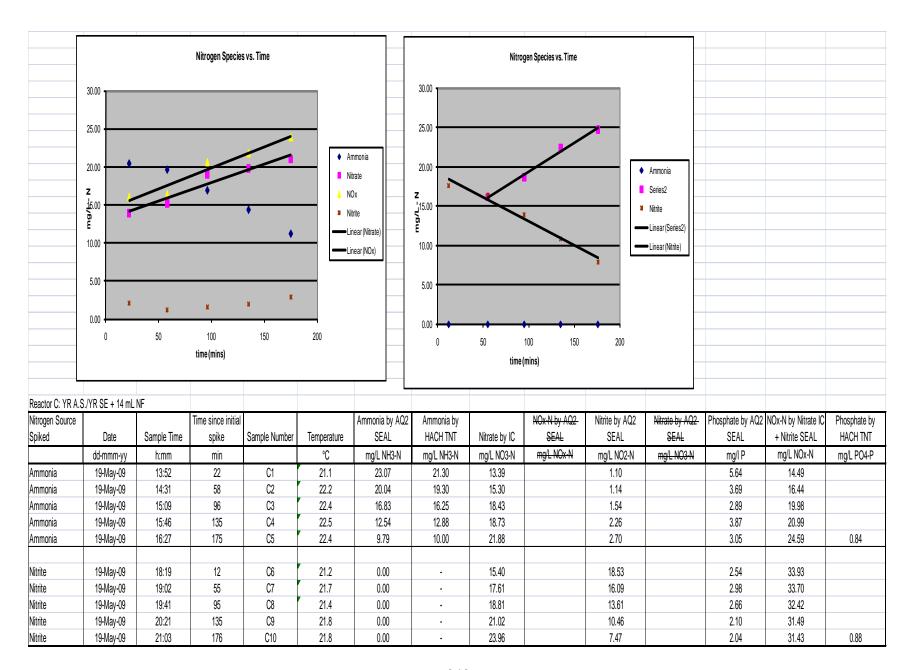


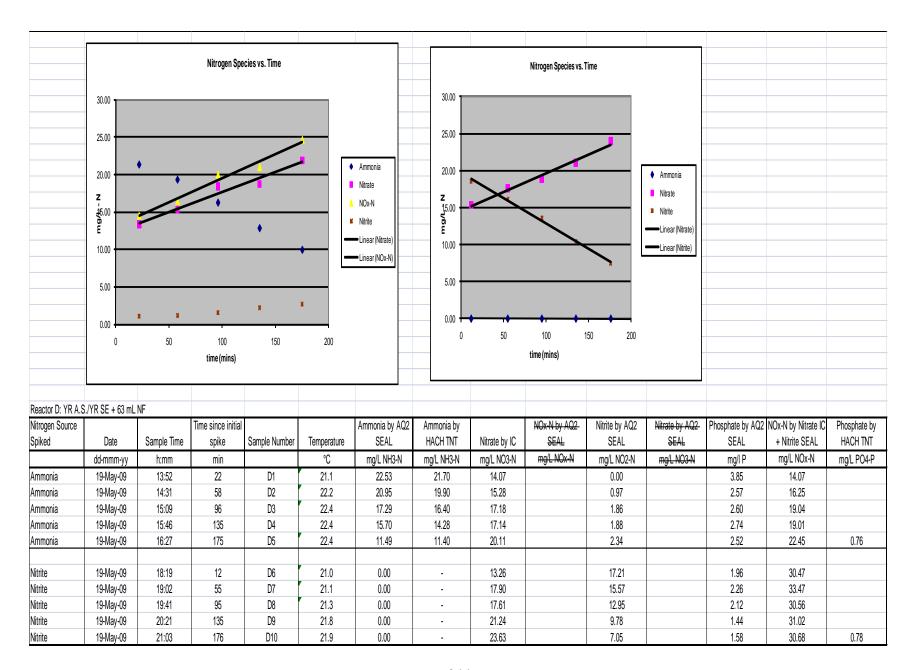
Veek 15 – QACs Day 1 AOB/NOB					
ample Data Report for Nansemond Nitrification Inhibition Study					
AOB & NOB Experimentation:					
I.AOB					
a. Spiked four 3L reactors with 25 mg/L NH ₄ .					
b. Each reactor is running continously by use of stir bars .					
c. 2 L of the diluent source is added to the reactors.					
d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the bioma	s was concentrated from 201 to 91				
e. Constant DO and pH were monitored and logged throughout the experiment.	3 Wa3 concentrated 11011120 E to 3 E	•			
f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valv					
70					
 g. LabView software was used to manage DO and pH recording as well as program and implem h. 5 samples were collected over a period of 3 hours through sampling ports on the reactors are 		millinara 0 4E um filtara			
i. Reactors B, C, and D in the diluent source incorporated varying concentrations of Nature Fre	n Chemical Tollet Additive based or	different assumptions			
(See Additive Dose Calculation worksheet)					
II. NOB					
a. Spiked four 3L reactors with 15 mg/L NO $_2^{\cdot}$ after ammonia levels were <1 mg/L NH $_3$ -N which	as checked through the use of HAC	H TNT 831 Ammonia method.			
In addition all reactors were allowed to run overnight to consume all present ammonia.					
b. Each reactor is running continously by use of stir bars .					
c. Constant DO and pH were monitored and logged throughout the experiment.					
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valv	5.				
e. LabView software was used to manage DO and pH recording as well as program and implem	nt the solenoid valves.				
f. 5 samples were collected over a period of 3 hours through sampling ports on the reactors are	I filtered into sample tubes through	millipore 0.45 μm filters.			
g. Reactors B, C, and D in the diluent source incorporated varying concentrations of Nature Fre	h Chemical Toilet Additive based or	different assumptions			
(See Additive Dose Calculation worksheet)					

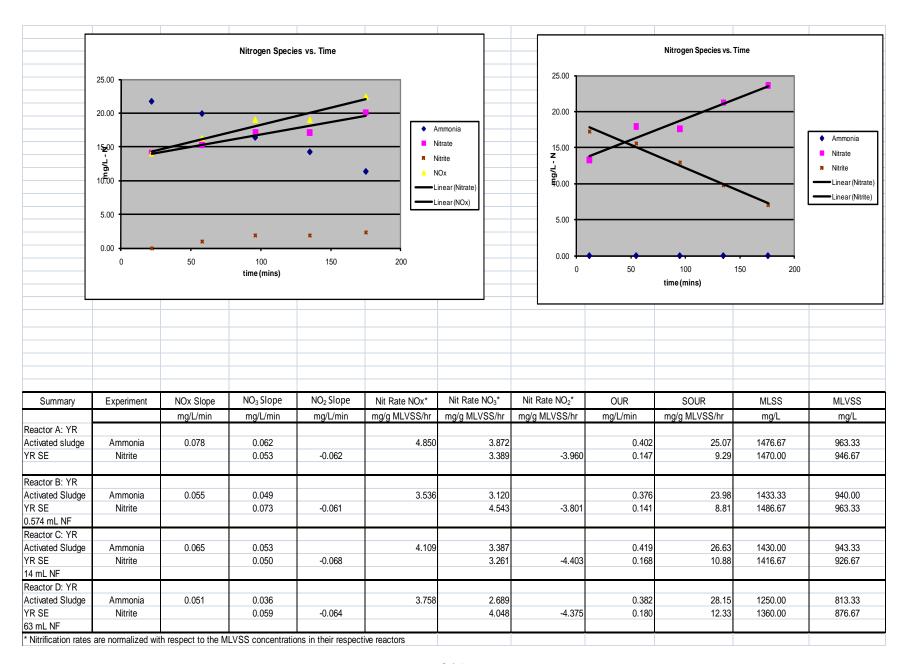
Danatas	۸.	VD	A C	/ VD	٥E
Reactor	Α.	YK	AS	/ YR	SE

Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOx-N by AQ2 SEAL	Nitrite by AQ2 SEAL	Nitrate by AQ2 SEAL	Phosphate by AQ2 SEAL	NOx-N by Nitrate IC + Nitrite SEAL	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		င့	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	19-May-09	13:52	22	A1	21.2	25.16	22.10	14.10		0.25		5.32	14.35	
Ammonia	19-May-09	14:28	58	A2	21.8	22.27	20.10	15.25		1.03		5.25	16.28	
Ammonia	19-May-09	15:06	96	A3	22	17.32	16.75	18.53		0.80		3.34	19.34	
Ammonia	19-May-09	15:45	135	A4	22.3	15.21	13.80	20.83		2.16		3.08	22.99	
Ammonia	19-May-09	16:25	175	A5	22.2	10.78	11.08	23.21		2.68		2.90	25.89	0.78
Nitrite	19-May-09	18:17	12	A6	21	0.00	-	13.61		18.38		2.61	31.99	
Nitrite	19-May-09	19:00	55	A7	21.5	0.00	-	20.40		16.63		3.47	37.03	
Nitrite	19-May-09	19:40	95	A8	21.7	0.00	-	18.62		14.17		3.13	32.80	
Nitrite	19-May-09	20:20	135	A9	21.9	0.00	-	21.19		11.32		2.36	32.51	
Nitrite	19-May-09	21:01	176	A10	21.9	0.00	-	24.06		8.27		2.19	32.34	0.92









Week 15 – Day 2 AOB/NOB

· · · · · · · · · · · · · · · · · · ·	
Sample Data Report for Nansemond Nitrification Inhibition Study	
AOB & NOB Experimentation:	
I. AOB	

a. Spiked four 3L reactors with 25 mg/L NH_a.

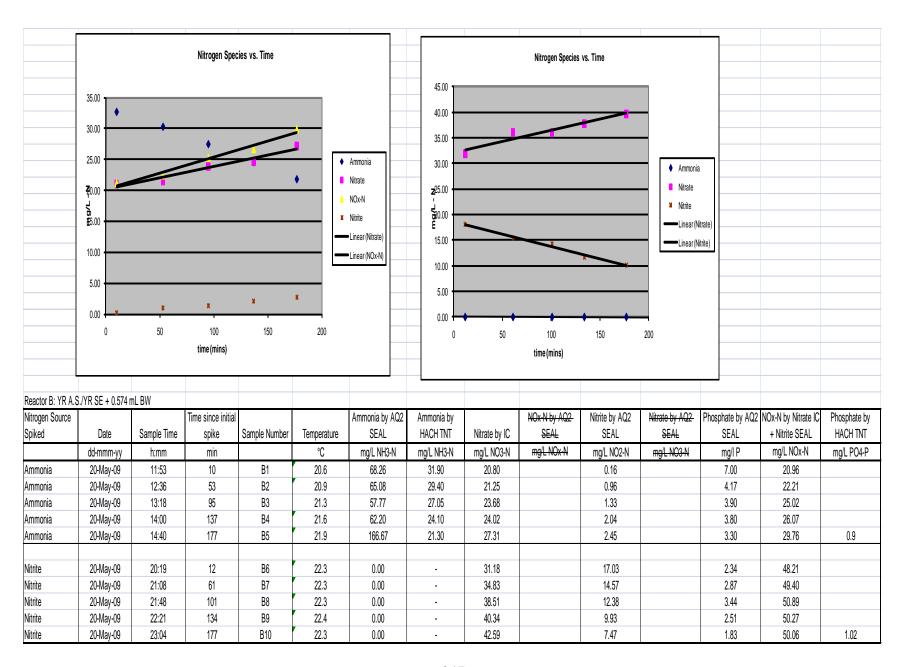
- b. Each reactor is running continously by use of stir bars.
- c. 2 L of the diluent source is added to the reactors.
- d. 1L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20L to 9 L.
- e. Constant DO and pH were monitored and logged throughout the experiment.
- f. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.
- g. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.
- h. 5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 μ m filters
- i. Reactors B, C, and D in the diluent source incorporated varying concentrations of Blue Works Chemical Toilet Additive based on different assumptions (See Additive Dose Calculation worksheet)

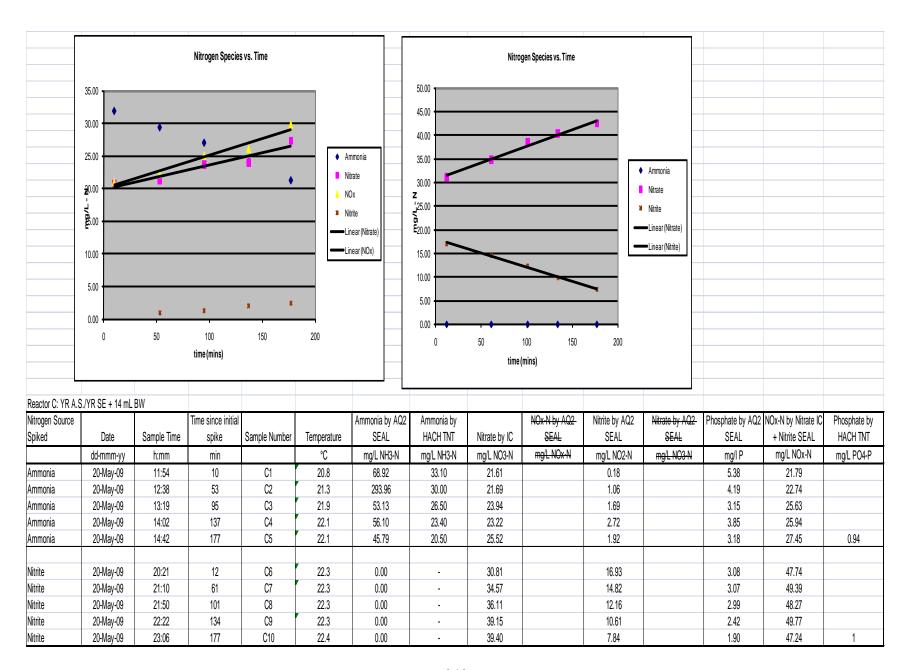
II. NO

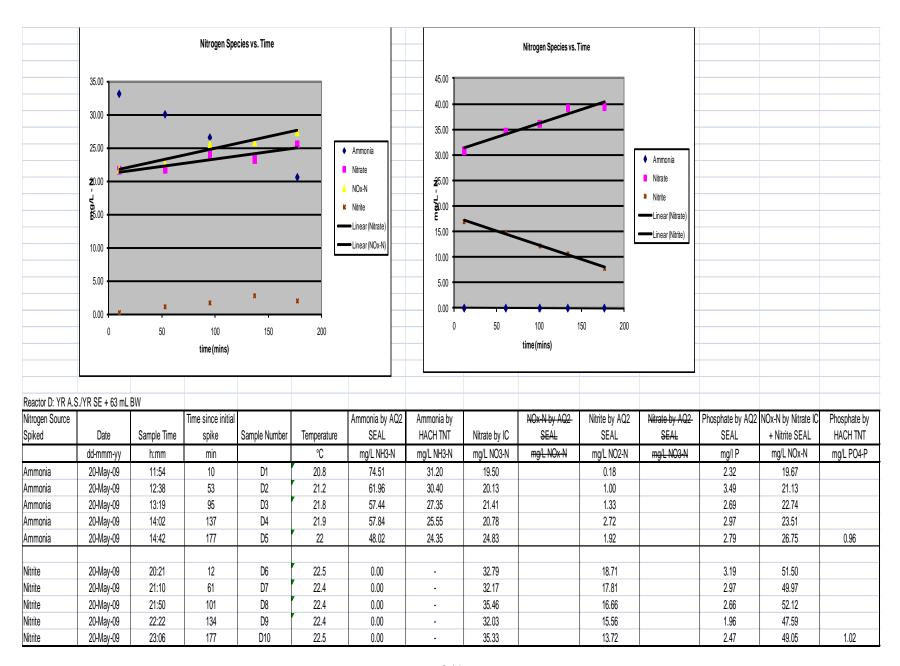
- a. Spiked four 3L reactors with 15 mg/L NO₂ after ammonia levels were <1 mg/L NH₃-N which was checked through the use of HACH TNT 831 Ammonia method. In addition all reactors were allowed to run overnight to consume all present ammonia.
- b. Each reactor is running continously by use of stir bars.
- c. Constant DO and pH were monitored and logged throughout the experiment.
- d. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.
- e. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.
- f. 5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 µm filters.
- g. Reactors B, C, and D in the diluent source incorporated varying concentrations of Blue Works Chemical Toilet Additive based on different assumptions (See Additive Dose Calculation worksheet)

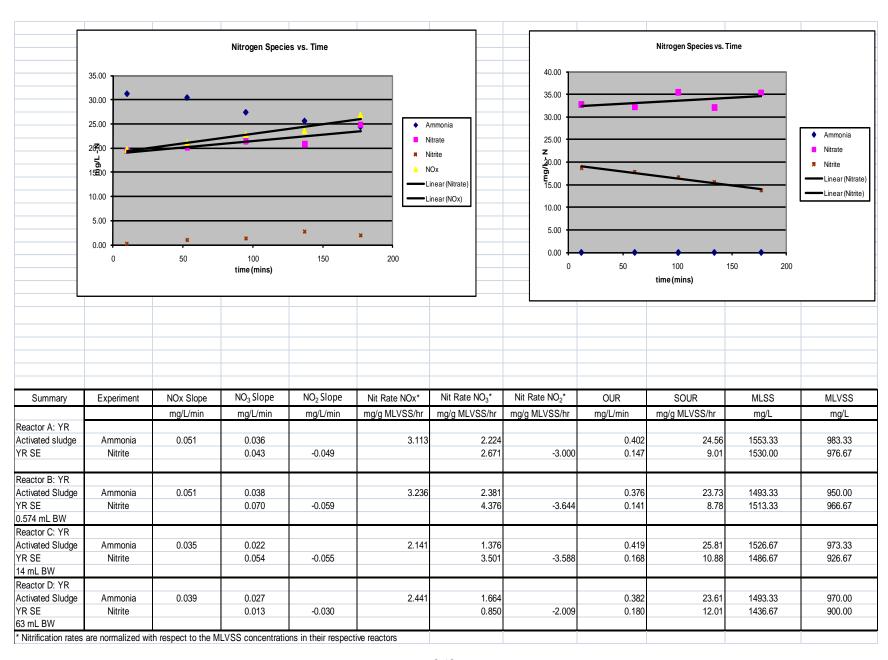
Reactor A:	YR	A.S	./YR	SE	

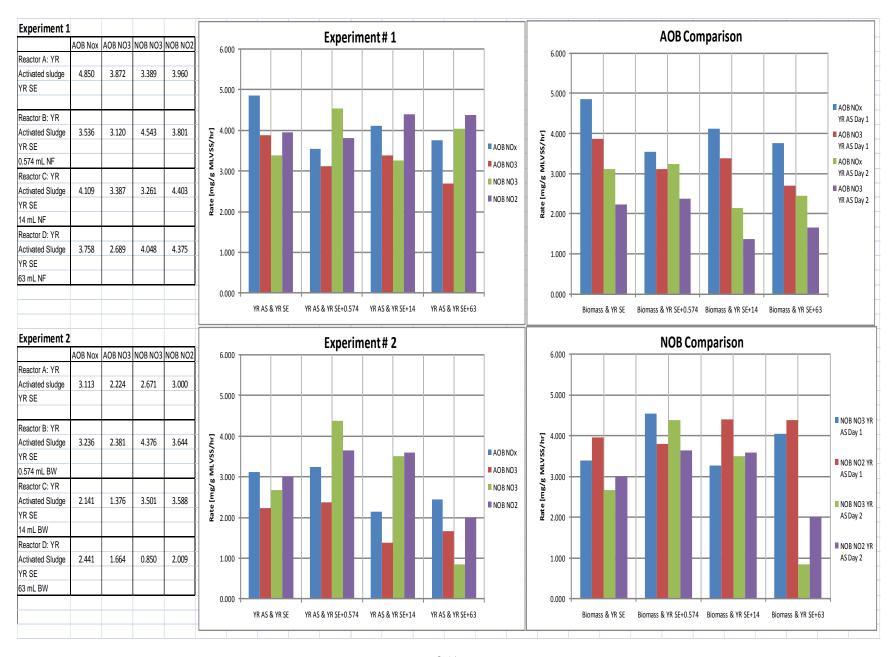
Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by IC	NOx-N by AQ2- SEAL	Nitrite by AQ2 SEAL	SEÁL	Phosphate by AQ2 SEAL	+ Nitrite SEAL	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/l P	mg/L NOx-N	mg/L PO4-P
Ammonia	20-May-09	11:53	10	A1	20.6	68.22	32.70	21.03		0.17		9.14	21.20	
Ammonia	20-May-09	12:36	53	A2	21.0	326.24	30.30	21.49		0.97		5.51	22.46	
Ammonia	20-May-09	13:18	95	A3	21.3	55.82	27.50	23.87		1.35		4.45	25.21	
Ammonia	20-May-09	14:00	137	A4	21.6	298.77	24.60	24.59		2.05		3.12	26.64	
Ammonia	20-May-09	14:40	177	A5	21.9	57.13	21.85	27.12		2.67		2.81	29.79	0.86
Nitrite	20-May-09	20:19	12	A6	22.1	0.00	-	31.93		18.11		3.36	50.04	
Nitrite	20-May-09	21:08	61	A7	22.1	0.00	-	36.04		15.52		3.73	51.56	
Nitrite	20-May-09	21:48	101	A8	22.2	0.00	-	36.13		14.27		3.28	50.40	
Nitrite	20-May-09	22:21	134	A9	22.2	0.00	-	37.85		11.68		2.85	49.53	
Nitrite	20-May-09	23:04	177	A10	22.3	0.00	-	39.70		10.16		2.12	49.85	1.2







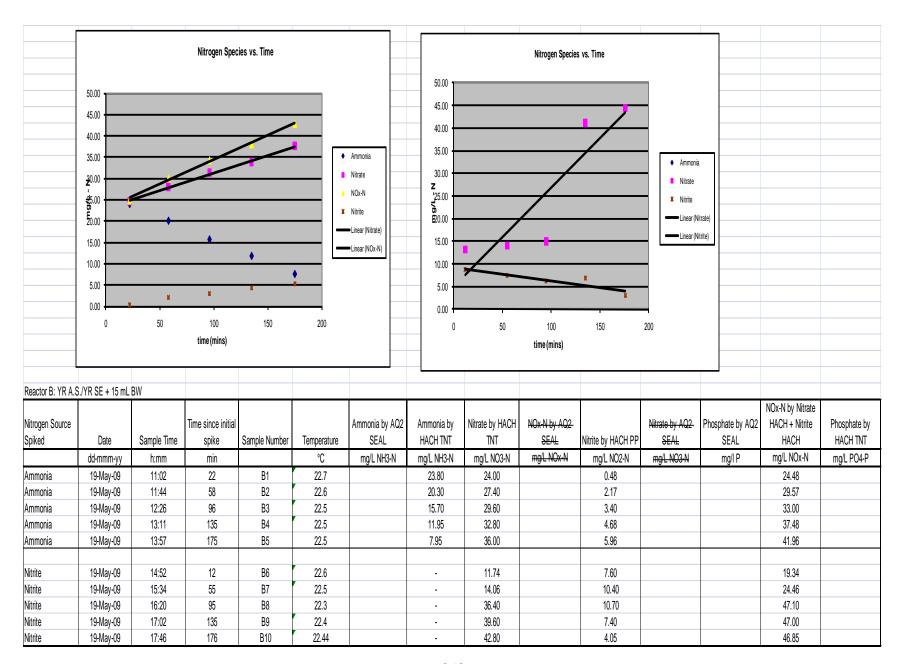


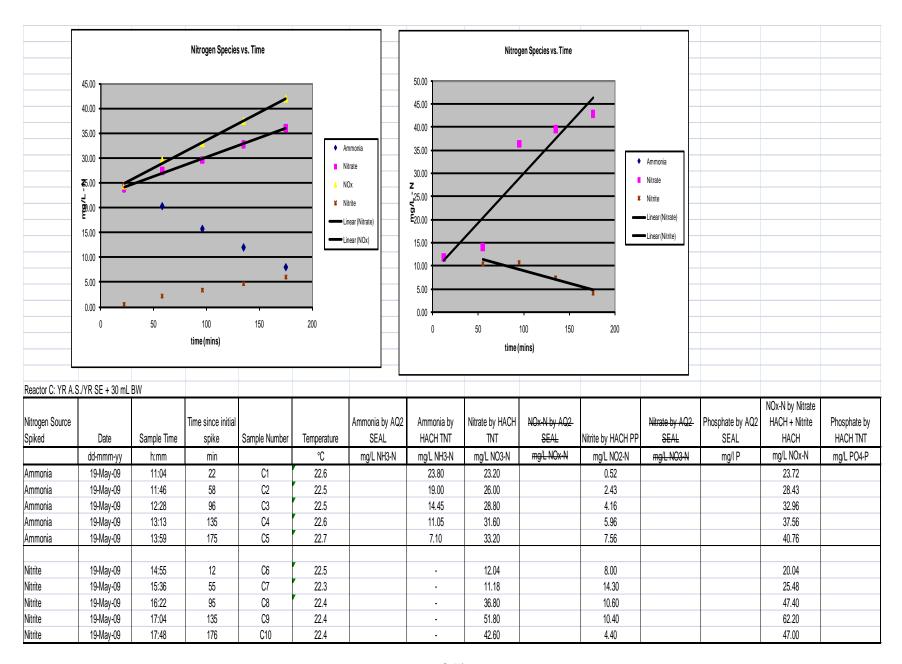


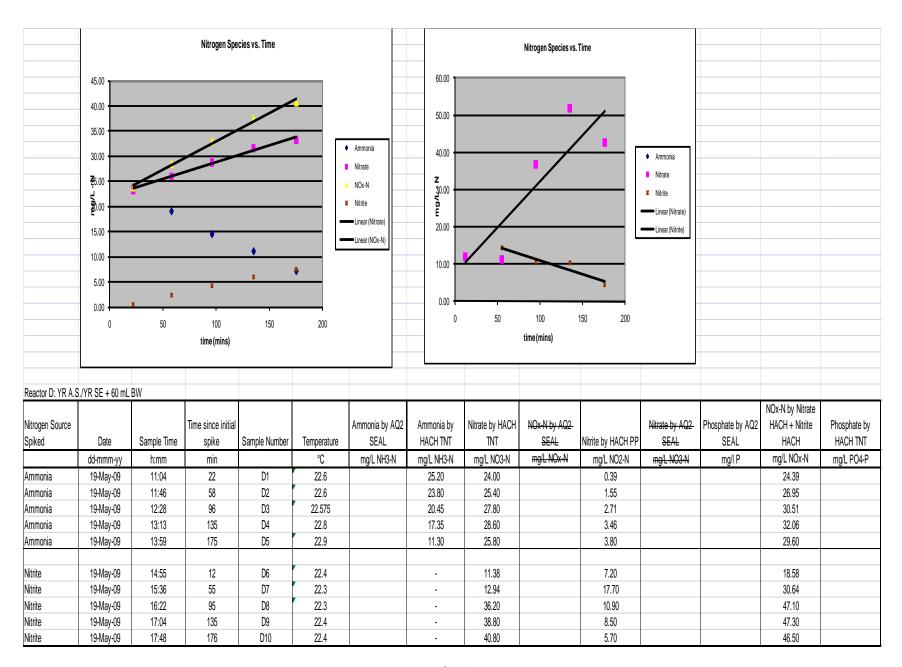
Week 16 – QAC Product B Day 1 AOB/NOB

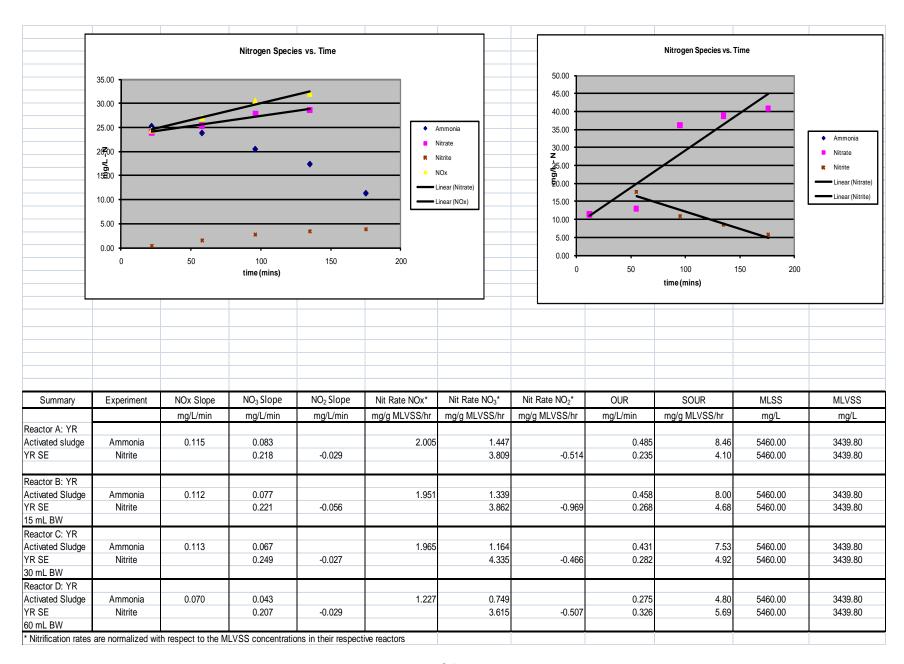
ample Data Report for Nansemond Nitrification Inhibition Study							
AOB & NOB Experimentation:							
.AOB							
a. Spiked four 3L reactors with 25 mg/L NH ₄ .							
b. Each reactor is running continously by use of stir bars .							
c. 2 L of the diluent source is added to the reactors.							
d. 1L of concentrated biomass is added to the reactors. For this weeks experime	nts the hinmass was o	oncentrated from 2	201 to 91				
e. Constant DO and pH were monitored and logged throughout the experiment.	into the biomass was e	.oncentratea mont	.021032.				
f. Oxygen/air blend was sparged into the reactors to maintain DO levels through	solenoid valves						
g. LabView software was used to manage DO and pH recording as well as progra		sayley hinnalos					
h. 5 samples were collected over a period of 3 hours through sampling ports on 1			c th rough millinare (1 15 um filtars			
i. Reactors B, C, and D in the diluent source incorporated varying concentration:			• .	•			
(See Additive Dose Calculation worksheet)	Of blue Works Cheffin	cai Tollet Additive t	idseu on uniterent ds	sumptions			
(See Additive Dose Calculation Worksheet)							
NOD							
I.NOB	AU		CLIA CLI TAIT 004				
a. Spiked four 3L reactors with 20 mg/L NO ₂ after ammonia levels were <1 mg/L	,	cked through the u	se of HACH INT 831	Ammonia metnod			
n addition all reactors were allowed to run overnight to consume all present amn	ionia.						
b. Each reactor is running continously by use of stir bars.							
c. Constant DO and pH were monitored and logged throughout the experiment.							
d. Oxygen/air blend was sparged into the reactors to maintain DO levels through							
e. LabView software was used to manage DO and pH recording as well as progra							
f. 5 samples were collected over a period of 3 hours through sampling ports on t							
g. Reactors B, C, and D in the diluent source incorporated varying concentrations							
(See Additive Dose Calculation worksheet)							
actor A: YR A.S. / YR SE							

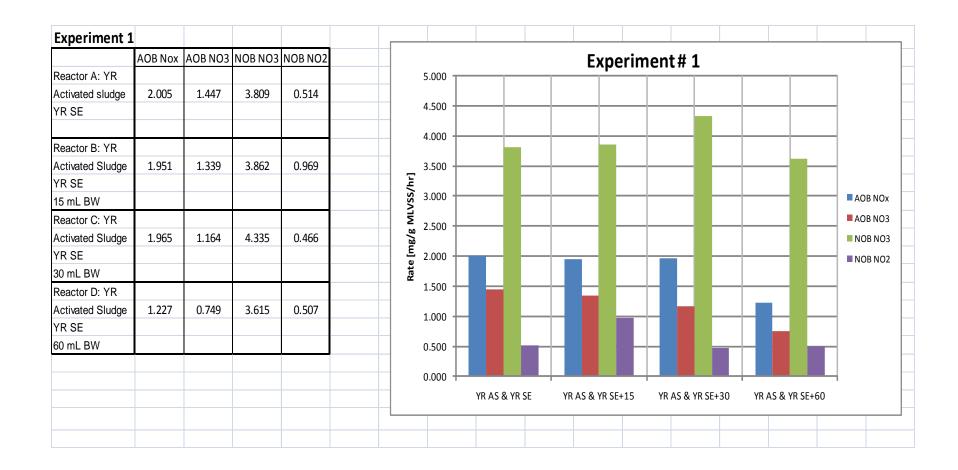
Reactor A: YR A.S	S. / YR SE												
Nitrogen Source Spiked	Date	Sample Time	Time since initial spike	Sample Number	Temperature	Ammonia by AQ2 SEAL	Ammonia by HACH TNT	Nitrate by HACH TNT		Nitrite by HACH PP	Nitrate by AQ2 SEAL	NOx-N by Nitrate HACH + Nitrite HACH	Phosphate by HACH TNT
	dd-mmm-yy	h:mm	min		°C	mg/L NH3-N	mg/L NH3-N	mg/L NO3-N	mg/L NOx-N	mg/L NO2-N	mg/L NO3-N	mg/L NOx-N	mg/L PO4-P
Ammonia	17-Jun-09	11:02	22	A1	22.6		24.00	24.60		0.34		24.94	
Ammonia	17-Jun-09	11:44	58	A2	22.45		20.10	28.00		2.12		30.12	
Ammonia	17-Jun-09	12:26	96	A3	22.4		15.75	31.40		2.92		34.32	
Ammonia	17-Jun-09	13:11	135	A4	22.4		11.90	33.80		4.34		38.14	
Ammonia	17-Jun-09	13:57	175	A5	22.425		7.68	37.60		5.36		42.96	
Nitrite	17-Jun-09	14:52	12	A6	22.6			13.22		8.80		22.02	
Nitrite	17-Jun-09	15:34	55	A7	22.4		-	14.06		7.40		21.46	
Nitrite	17-Jun-09	16:20	95	A8	22.4		-	15.04		6.30		21.34	
Nitrite	17-Jun-09	17:02	135	A9	22.4		-	41.20		6.90		48.10	
Nitrite	17-Jun-09	17:46	176	A10	22.4			44.40		3.05		47.45	











ple Data Report for Nansemond Nitrification Inhibition Study		
IB & NOB Experimentation:		
OB		
. Spiked four 3L reactors with 25 mg/L NH ₄ .		
. Each reactor is running continously by use of stir bars .		
. 2 L of the diluent source is added to the reactors.		
. 1 L of concentrated biomass is added to the reactors. For this weeks experiments the biomass was concentrated from 20 L to 9 L.		
. Constant DO and pH were monitored and logged throughout the experiment.		
Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.		
. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.		
. 5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 μm filters		
Reactors B, C, and D in the diluent source incorporated varying concentrations of Blue Works Chemical Toilet Additive based on different assumptions		
(See Additive Dose Calculation worksheet)		
NOB		
. Spiked four 3L reactors with 20 mg/L NO ; after ammonia levels were < 1 mg/L NH ;-N which was checked through the use of HACH TNT 831 Ammonia method.		
addition all reactors were allowed to run overnight to consume all present ammonia.		
. Each reactor is running continously by use of stir bars .		
. Constant DO and pH were monitored and logged throughout the experiment.		
. Oxygen/air blend was sparged into the reactors to maintain DO levels through solenoid valves.		
. LabView software was used to manage DO and pH recording as well as program and implement the solenoid valves.		
5 samples were collected over a period of 3 hours through sampling ports on the reactors and filtered into sample tubes through millipore 0.45 µm filters.		
. Reactors B, C, and D in the diluent source incorporated varying concentrations of Blue Works Chemical Toilet Additive based on different assumptions		
(See Additive Dose Calculation worksheet)		

Reactor A:	YR A.S. /	YK SE

Troductor / II TTT/IIC													
Nitrogen Source Spiked	Date dd-mmm-vy	Sample Time	Time since initial spike	Sample Number	Temperature °C	Ammonia by AQ2 SEAL mg/L NH3-N	Ammonia by HACH TNT mg/L NH3-N	Nitrate by HACH TNT mg/L NO3-N	NOx-N by AQ2- SEAL ma/L NOx-N	Nitrite by HACH PP	Nitrate by AQ2 SEAL mg/L NO3-N	NOx-N by Nitrate HACH + Nitrite HACH mg/L NOx-N	Phosphate by HACH TNT ma/L PO4-P
	- //	11.111111	1		,	IIIg/L INDJ-IN	IIIg/L INFIS-IN	IIIg/L NO3-IV	HIGHE NOX-14	IIIg/L INOZ-IN	HIG/L NOO-IN	IIIg/L INOX-IN	IIIg/L FO4-F
Ammonia	23-Jun-09	14:00	22	A1	22.6		23.30	22.00		0.73		22.73	
Ammonia	23-Jun-09	14:43	58	A2	23.2		15.50	26.60		4.38		30.98	
Ammonia	23-Jun-09	15:26	96	A3	23.5		7.00	32.40		7.16		39.56	
Ammonia	23-Jun-09	16:07	135	A4	24		0.00	36.40		9.85		46.25	
Ammonia	23-Jun-09	16:48	175	A5	24.2		0.00	41.20		5.70		46.90	
APA 24	041 00	0.00	40	40	00.0			04.00		47.40		40.00	
Nitrite	24-Jun-09	8:33	12	A6	22.9		•	31.20		17.40		48.60	
Nitrite	24-Jun-09	9:15	55	A7	23.6		-	33.80		14.40		48.20	
Nitrite	24-Jun-09	9:56	95	A8	24.3		-	36.80		10.70		47.50	
Nitrite	24-Jun-09	10:47	135	A9	24.9		-	41.20		5.80		47.00	
Nitrite	24-Jun-09	11:29	176	A10	25		-	44.20		2.05		46.25	

