

Using Pressure Transducers for Noninvasive Heart and Respiratory Monitoring

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Abstract

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Detecting heart and respiratory rates is an essential means of providing emergency medical care. Current methods of detecting such signals include the widely used electrocardiography (ECG) method. Other more manual methods of heart and respiratory rate estimation require a practitioner to constantly observe the patient. These methods are time consuming and detract valuable time from emergency medical care. This thesis presents a novel, hands off, heart and respiratory monitor (HARMONI). It uses pressure transducers and medical tubing placed on a person's chest. The tubing is plugged off at one end, and then attached to a pressure transducer at the other end. The transducer sees spikes in voltage whenever the pressure inside the tubing changes. Heart and respiratory rates both cause expansion in the chest, increasing the pressure in the tubing, and causing the transducer to see a change in voltage. The method was first validated, and then tested in a simulated environment. Finally, the device was transformed in to a full system prototype. Human tests were conducted to correlate the signal with that of an industry standard ECG device. This thesis explains how heart and respiratory rates can be derived using signal processing techniques and a simple non-invasive sensor. This device is a rapidly deployable tool that has the potential to save lives specifically in mass casualty situations. It would be a force multiplier, allowing a single responder to monitor multiple casualties, saving time and lives.

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Receiving a degree, at any level, is an accomplishment of not just one individual, but rather an accomplishment of many. It is a celebration of all those teachers who have had an impact on a person, throughout their life. A good teacher is one who helps you succeed in the subject in which they teach. A great teacher is one who helps you succeed in all aspects of life. I have had the rare fortune of having many great teachers in my life. It is these great people whom I dedicate this thesis.

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Chapter 1: Introduction

Tracking heart and respiratory rates has always been at the foundation of emergency medical care. For years, Emergency Medical Technicians, or EMT's, have been told to rapidly estimate a casualty's heart and respiratory rate in order to better triage their patients. However, there are often cases where patients are over-triaged or even under-triaged [1]. This study is intended to explore the idea of using pressure transducers to actively detect heart and respiratory rates in a rapidly deployable form factor for use in emergency response situations. The objective is to provide more information throughout the triage process, without hindering the caregiver's ability to respond to multiple casualties.

Emergency medical personnel, have been using skin electrodes to monitor heart rate since the early 1900's [2]. This method of gathering vital signs is widely considered non-invasive [3] to a patient, when compared to more manual methods. However it is still expensive and requires direct skin contact. An example image showing the placement of skin electrodes is shown below in Figure 1-1.

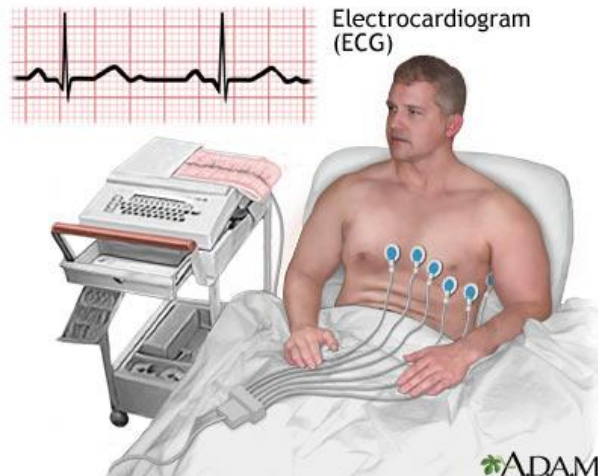


Figure 1-1. ECG image from the UMD Medical Center [4].
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The importance of monitoring heart and respiratory rate of casualties in a mass casualty incident (MCI) is to determine the current state of that casualty prior to triaging them. For instance, if the casualty has a rapid heart rate, they may be going in to shock. If their respiration is shallow, they may have some sort of injury or bleeding that is blocking their airway. If the initial reading of vital signs allows an EMT to respond to these issues immediately, would a continuous reading not allow the EMT to monitor such issues over time? The value of reading heart and respiratory rate over time is that the patient's state can be monitored for improvement or deterioration. The value of an EMT in an MCI is to rapidly assess the casualties for symptoms of more serious injuries. Then, the EMT either addresses those injuries on the scene, or sends the casualty to a hospital for further medical attention. The job of an EMT in an MCI is an important one, requiring extensive training, and constant improvement.

1.1 Emergency Medical Response Situations

When an emergency MCI occurs, the overall response is usually a three step process. The responders typically begin by setting up a command center in a safe location near the incident scene. The second step, and perhaps the most critical one, is to triage the patients. Triage is the process of determining the condition of all victims involved in the incident. During this stage, patients are prioritized based on severity of their injuries. The third step is to transport patients to a hospital in the prioritized order determined during the triage stage. There are several different acronyms and methods used to help first responders become better at the entire process. In general, they all follow the same three step approach as outlined. Similarly, the individual response to each casualty is a three step process. This individual response is known as the ABC's of EMT's. This prompts checking the airway for obstruction, then the breathing rate, and finally checking the casualty's circulation.

1.1.1 *Civilian Emergency Medical Applications*

In a study related to the use of triage protocols for assessing patients, a well accepted protocol was evaluated for a mass casualty triage of 132 patients. It was found that 64 patients (48%) were triaged correctly, 65 were over-triaged (49%), and 3 were under-triaged (2%) [1]. During a MCI, triage decisions must be made after taking the availability of resources in to consideration. As the National Guideline for Mass Casualty Triage reports, "Responding to 30 victims with 4 ambulances is different from responding with 16 ambulances and numerous responders" [1].

It is widely accepted in the emergency medical community that a minimum 15 second time period is required to estimate a patient's pulse. The same time period is required for respiratory rate [5]. However, that is based on the assumption that the EMT is capable of doing near instantaneous calculations in their head, and that they did not have difficulty finding the pulse in the first place. A rapidly deployable heart and respiratory monitor would allow for emergency responders to concentrate more on triaging the patients appropriately, than on estimating the patient's vital signs.

1.1.2 *Military Emergency Medical Applications*

EMTs are constantly being thrown in to situations on a daily basis where they are accustomed to restore order out of chaos. However, there are times when the need for their services far exceeds their capacity to provide them [6]. No one can attest to this more than, Thomas A. Middleton, who served as a combat medic in Iraq as part of Task Force Saber from June 2005 to June 2006, before returning home to serve as assistant fire marshal in Burlington, Vermont. Thomas wrote a book on his experiences, but relates several very important tips based on his time as a combat medic. The first thing to do in a MCI is to shout at all of the casualties and ask them to move toward the sound of your voice. Those that respond are likely of lowest priority, those that can at least walk are termed delayed and those that don't move are termed immediate. Middleton goes on to note that the responder's knees should never touch the ground, because that results in a tendency to focus on a single patient. A general rule of thumb is that each patient should be re-assessed every five minutes for an immediate patient, and every fifteen minutes for a delayed patient [6]. Note that the main concern of an emergency responder in an

MCI is to tag and assess patients so that further treatment can be made down the line. It is a difficult but necessary task, and emergency responders need all the help they can get. Middleton himself admits that “no one ever gets it exactly right, and there will always be room for improvement” [6].

1.1.3 The Need for Vital Sign Monitors in Mass Casualty Incidents

There are incidents every day where the number of casualties outweighs the resources of the emergency responders. Therefore, any monitor that would seamlessly fit in to such a situation, without requiring extensive training, or without interrupting patient care, would be a worthwhile technology. There are several methods that exist that allow vital sign monitoring, from respiratory belts, to wet electrodes, to manual estimation. However, there is no single device that can seamlessly, noninvasively, and rapidly be applied during an MCI. There is no single device that would allow an individual EMT to monitor multiple casualty's heart and respiratory rates simultaneously.

What if there was such a device? What if there were methods that allowed an EMT to monitor casualty's heart rates without having to place electrodes on to the skin? What if there were ways for one EMT to wirelessly monitor all casualties, without being distracted from the most critical ones?

1.2 Detecting Vital Signs without Using Bioelectric Signals

There are several areas of research in to noncontact patient monitoring. While some technologies such as an ECG monitor bio-electric signals, other technologies monitor pure mechanical signals within the body. One such method, of mechanical monitoring, is using highly sensitive pressure transducers to monitor patients. It works by plugging off one end of a length of tubing, then attaching the other end to the pressure transducer. The mechanical response of the chest due to heart and respiratory rate causes the pressure in the tubing to change. The tubing replaces the skin electrode as the nearest part of the device to the patient. It can be used in a chest strap form factor, and placed over clothing near the area of maximum cardiac impulse. This would greatly reduce the time that otherwise would be required to remove the patient's clothing and to place the electrodes properly.

The ability to create a sensor that does not require skin electrodes would be valuable in many ways beyond its reduced invasive nature. Consider a MCI such as the ones EMTs are called to. Many different people require aid and monitoring, yet the ratio of patients requiring attention to EMTs is often too high. Being able to swiftly install a wireless monitoring device on patients over their clothing would be a valuable time saver. Also, if the system can generate alerts when vital signs fall outside normal parameters, this would be of tremendous value as well.

This technology could also be utilized for monitoring military and civilian underwater technicians. Skin electrodes are a reliability risk when being used underwater. In contrast, medical tubing and the pressure transducers can easily be waterproofed for use in monitoring the heart and respiratory rates of a diver. Combine this with a wireless transceiver, and this device would be capable of providing underwater situational awareness. All of these benefits make the technology a worthwhile application in an MCI. What other technologies exist that can be applied in an MCI? What are their apparent benefits and drawbacks?

Chapter 2: Literature Review

There are several different methods of heart and respiratory monitoring that can be applied in an emergency response situation. Understanding what existing technologies have been able to accomplish, is the first objective. To do this, a review of heart and respiratory rate information is conducted, followed by a look at mobile health monitors that are currently on the market. This includes a look at both bioelectric and mechanical monitors.

Regardless of the approach, it is important to recognize that each technology has its pros and cons. For instance, ECG provides an in depth look at the heart's operation, giving more than just heart rate. It also provides respiratory rate by comparing the signals received by multiple leads. However, it is expensive, requires direct skin contact, and can easily be disrupted by movement or moist environments. Other industry standard technologies include the pulse oximeter, which is a small sensor usually placed on the fingertip that measures the oxygenation of a patient's hemoglobin. It is small, affordable, and provides a very clear heart rate signal. However, it is easily interrupted by small body movements, and its most practical application is in the intensive care unit (ICU) where patients are extremely stationary over time. Other current technologies include impedance and pressure belts for monitoring respiratory rate. They are great for monitoring respiratory rate, but do not necessarily make an attempt to monitor heart rate simultaneously.

While these traditional ECG monitors have provided the basis for heart and respiratory monitoring for decades, there has been a wealth of research on improving the technology. Such improvements include making them less invasive, more mobile, and by adding wireless capabilities. Integrating wireless heart and respiratory monitors is also one of the fields of improvement. It is an idea that has been around since the early 2000's. The first application of wireless technology in a medical environment was wireless local area networks [7]. Since then, integrating wireless technologies in to medical devices has been in the forefront of noninvasive medical research. One attempt at making ECG's less invasive is non-contact ECG's which utilize capacitively coupled leads to measure heart rate through thin layers of material such as a t-shirt. Another attempt to truly push the definition of the term non-invasive is a vital sign monitor of humans inside a vehicle. A mechanical explanation of what they are attempting to do might disprove their results. However, their methods are helpful in understanding how to use pressure sensors to monitor vital signs.

2.1 Background Information on Heart and Respiratory Rate

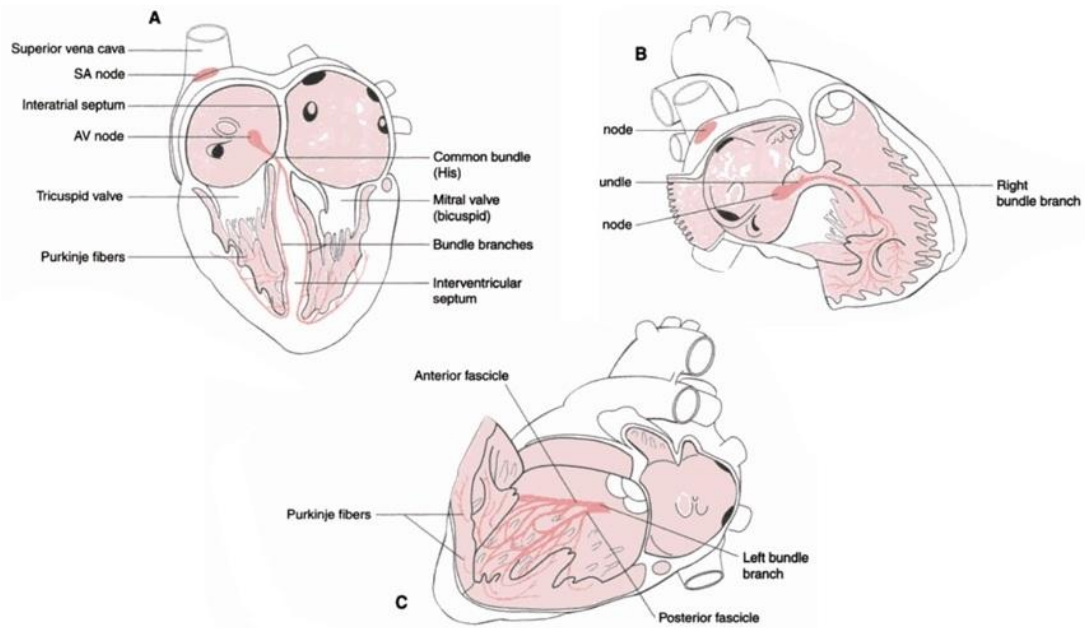
When considering heart and respiratory rates in emergency situations, one must be prepared to detect both the upper and lower limits of each. Tachycardia refers to the abnormally rapid heart rate, while bradycardia refers to abnormally slow heart rate [8]. Similarly, tachypnea and bradypnea refer to the abnormal high and low respiratory rates respectively. All four limits vary with age and health. However, in general, safe heart rate can vary as low as 40 beats per minute (HR) for experienced adult athletes, and as high as 160 HR for babies. In general the safe respiratory rate is as low as 12 breaths per minute (RR) for older adults, and as high as 40 RR for newborn babies [5]. With such a range of values, there is a need for methods and devices that are both accurate and versatile.

2.2 Manual Methods for Heart and Respiratory Rate Estimation

The most common method of estimating heart and respiratory rates is to do so manually. Nurses, doctors, and in this case EMT's have been trained to do this in a very simple manner. For observing the radial pulse, they place their middle and index fingers gently along the inside of the wrist. For a quick estimate, count the number of pulses felt during 15 seconds, then multiply that number by 4 to get the bpm. The same thing goes for breathing. Watch the patient's chest and count the number of times it cycles during 15 seconds, and then multiply that number by 4 to get the breaths per minute. Assuming the patient is cooperative, and the EMT is rapid in their estimations, this task takes a minimum of 30 seconds to complete. The result of manual heart and respiratory rate estimation is a single value at a specific instance in time and does not provide the attention that many patients require over long term care. This is the instigating factor behind creating monitors that actively track the patient's vital signs over time.

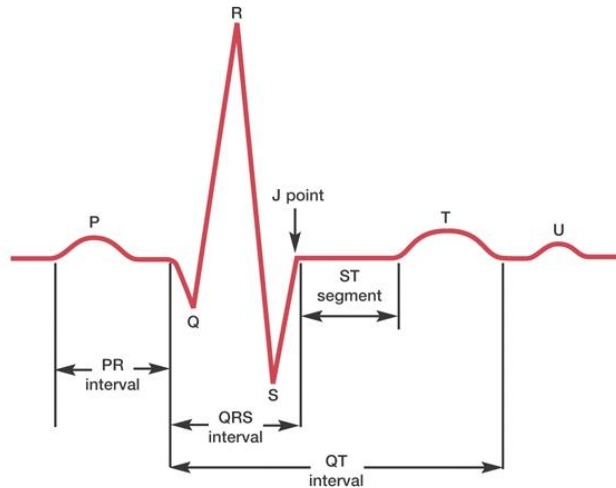
2.3 Electrocardiography (ECG)

The most widely used form of heart monitoring is the electrocardiography. Originally invented in 1872, electrocardiography is "the recording of the electrical activity generated by the cells of the heart that reaches the body surface" [9]. It is this electrical activity of the heart that initiates the muscle to contract and pump blood to the rest of the body. The most common application of an ECG is to observe abnormal rhythms of the heart in patients. An ECG wet electrode is what connects the measurement device to the patient. Electrodes are considered to be wet when they contain an electrolyte gel. The purpose of this gel is to decrease the influence of the skin on the impedance by moisturizing its dry outer layer and making it more ion-conductive [10]. They are required to be placed directly against the patient's skin, and often require the area to be cleaned with alcohol, or even shaved. The standard 12 lead ECG is capable of representing the heart's electrical activity from several different locations on the patient. The results of an ECG are in terms of voltage and are always comprised of different waves representing different contractions and behaviors of the heart. A diagram of the heart can be seen below in Figure 2-1.



**Figure 2-1. Diagram of the human heart [9].
Used under fair use, 2012.**

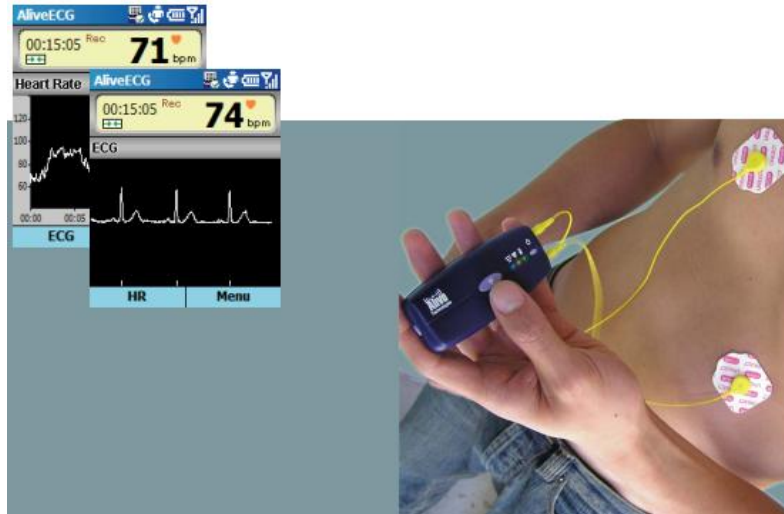
Each wave can be described as a P, QRS, or T wave. The P wave is a deflection wave that is caused by atrial depolarization. The PR interval is the time between the first variation in the P wave, and the first variation on the QRS spike. The three point QRS wave represents ventricular depolarization. The first downward deflection is the Q wave, and represents the depolarization of the interventricular septum. The large upward peak is the R wave, and represents the depolarization of the main mass of the ventricles. Finally the second downward deflection is the S wave, and represents the final depolarization of the ventricles at the base of the heart. The T wave corresponds with ventricular repolarization, hence the fact that the ST segment, is the time between ventricular depolarization and repolarization [11]. An example of the resulting waveform can be seen below in Figure 2-2.



**Figure 2-2. The basic pattern of electrical activity across the heart [11].
Used under fair use, 2012.**

Based on reliability and availability, the ECG is the clear industry standard when it comes to heart monitoring. This also makes it the most widely attempted base device for mobile heart monitoring. While it allows users to have a comprehensive view of their heart's performance, it still involves constant direct skin contact in order to function. As will be discussed, it is clear that all electrode based mobile heart monitors have their benefits and drawbacks due to this constraint. The most notable drawbacks being that they are invasive to the patient, take time to attach, and usually require the surface to be wet, but not so much as to disrupt the bioelectric signal.

One mobile technology that delivers ECG data to an active patient is the Alive Heart and Activity Monitor by Alive Technologies. This device has a wide variety of applications, from "cardiac rehab, to cardiovascular screening, home monitoring, disease management, falls monitoring, fitness monitoring, sports training" [12], and more. It utilizes a 3 axis accelerometer and a 2 lead electrode based ECG system. The small form factor allows it to be worn in nearly any activity. However, it is still limited to the fact that it requires the wet electrodes in order to operate. This means that it shares the same drawbacks of all wet electrode based approaches. Firstly the material used to create the electrode or paste could cause irritation and discomfort to the skin, especially if the subject is in the middle of a heavy physical activity and may be perspiring [13]. Another issue is that during bodily movements, the leads may loosen, breaking electrical contact and causing noise spikes in the signal [13]. Figure 2-3 below shows the Alive Mobile Cardiac monitor according to their product brochure.

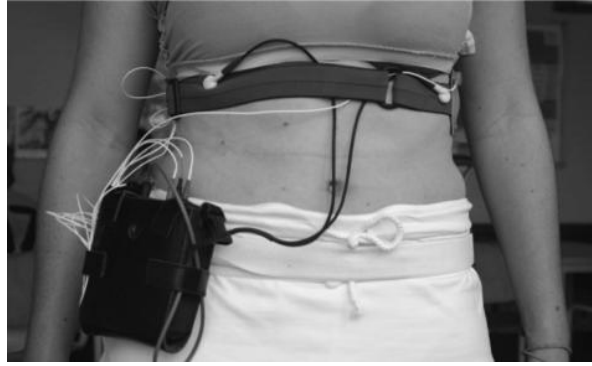


**Figure 2-3. Alive Mobile Cardiac Monitoring [12].
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While this device is capable of finding bioelectric signals, it does not seem to adhere to the physical lifestyle of the consumers it is geared towards. It does not account of the issue of excess sweat, nor does it deal with the issue of discomfort over long periods of time. This mobile cardiac monitoring unit may be portable but it does not have the ability to save time in an emergency response environment, because it still requires the precise placement of electrodes on the patient.

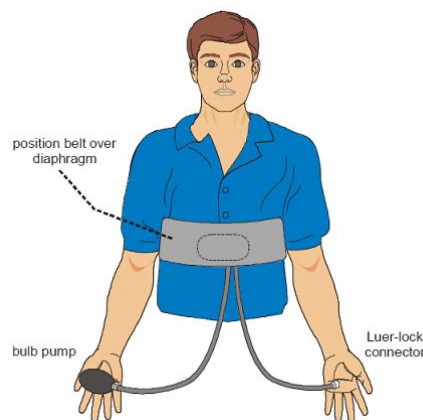
2.4 Respiratory Monitoring Using Impedance and Pressure Belts

Another technology that is akin with HARMONI in terms of recording respiratory rate is the Cardio Respiratory Belt by Twente Medical Systems International (TMSI). Specializing in physiological instrumentation, TMSI develops multi-channel amplifiers and data acquisition (DAQ) systems. Their cardio-respiratory belt utilizes a dual purpose belt that can measure both the resistance and inductance the belt when it undergoes changes due to breathing. The direct result is a respiratory signal, where apnea and other respiratory issues can be seen and evaluated. The belt does not specify that it needs to be worn directly on the skin, and is simply too large of an item to be used in the emergency MCI's where HARMONI is intended to be utilized. The cardio respiratory belt is seen below in Figure 2-4.



**Figure 2-4. Cardio Respiratory Belt by Twente Systems [14].
Used under fair use, 2012.**

Yet another technology that actually does utilize pressure transducers to detect respiratory rates is the Respiration Monitor Belt by Vernier Software and Technology. Their monitor is composed of a belt strap that is secured around the chest of a patient using Velcro. The belt is positioned directly over the diaphragm, with an air bladder resting near the base of the rib cage as seen in Figure 2-5 below.



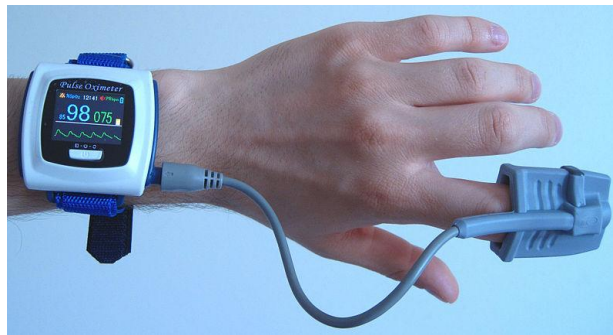
**Figure 2-5. Respiration Monitor Belt diagram [15].
Used under fair use, 2012.**

When the user has the belt in position, they pump up the air bladder to about 106 kPa, and then begin data collection. As the user breathes, their chest compresses against the air bladder, causing a change in pressure. The results are then analyzed using peak to peak estimation, to determine the respiration rate [15]. The device does not have any wireless connectivity options, and it requires a secondary sensor in order to estimate heart rate. That being said, it does utilize the same proposed method for monitoring heart and respiratory rate.

While all of these products described are capable of detecting heart and respiratory rate, all of the heart rate monitors require direct contact with the patient's skin. However, there are several non-contact methods for heart rate monitoring using newer technologies such as dry electrodes.

2.5 Pulse Oximeters

The traditional pulse oximeter uses a totally different signal than what ECG's use to monitor heart rate. While ECG takes advantage of variations in the heart due to electrical impulses due to contractions of the heart, a pulse oximeter utilizes an optical density monitor in the red and infrared wavelengths to compute arterial oxygen saturation. Invented in 1972, it originated during an attempt to measure tissue density in the ear lobe [16]. Later studies would show that the device could also read pulse and oxidation through a fingertip, or even a baby's foot. Since 1972 the technology has come a long way, as seen in the wearable pulse oximeter shown in Figure 2-6 below.



**Figure 2-6. A mobile pulse oximeter worn on the fingertip.
Used under public domain.**

Pulse oximeters operate under a basic concept. Oxygenated blood absorbs light at a red wavelength while deoxygenated blood absorbs light at an infrared wavelength. By sending both wavelengths of light through a thin layer of skin, and observing the amount of absorbed light on the other side, the device is able to estimate oxygen saturation levels.

Pulse oximeters are small, affordable, and offer a great method for monitoring heart rate and are the industry standard for monitoring oxidation of the blood. However, they are not easily waterproofed, require direct skin contact, and can be easily removed, or knocked off the hand.

2.6 Introduction to Non-contact Heart Monitoring

There are several issues with wet electrodes as have been discussed, which led to research in the area of dry electrodes. Electrodes are considered to be dry when they use an impedance transformation at the sensing site through active electronics [17]. First accomplished in 1968, capacitive coupling for detecting bioelectric signals has been limited by the ability to measure electric potential in free space. Therefore, it was not possible to measure bioelectric signals without direct skin contact. Finally, between 2001 and 2002, Quantum Applied Science and Research (QUASAR) developed “a new class of sensor that measures the electric potential in free space, i.e. without physical contact to any object” [17]. Since then, the field of non-contact ECG has grown tremendously.

2.6.1 *Capacitive Coupled Non-contact Electrocardiography (CCNE)*

One of the first evaluations of dry electrode technology was done in 2004 by Walter Reed Army Institute of Research (WRAIR). Their objective was to evaluate the use of CCNE's, for use on the future Objective WarFighter uniform. The results of the study showed that there was a high correlation between the normal ECG, and the CCNE. Furthermore, it was shown that the CCNE method provides unbiased estimates of ECG results [17].

Yet another evaluation of dry electrode technology was done in 2006 by a collaborative team of the University of California, and QUASAR, which was the group that originally developed capacitively-coupled electrodes. In the report, the engineers stated that wet electrodes are simply not capable of being used in high activity situations. The team was able to develop a 3 lead ECG monitoring system that weighed less than 17 grams. In addition the team says that their system integrates capacitive ECG sensors, which have been demonstrated to have 99% correlation with conventional electrodes [13].

The ECG chair, created by the Ford European Research and Innovation Center in Germany is another technology that utilizes non-contact ECG's to detect heart rate. The purpose of the technology is to actively monitor heart rate of a driver and in the case of a heart attack, to provide an automated vehicle response to ensure the safety of the driver of the vehicle. It is claimed that the seat produces correct results during 98 percent of driving time for 95 percent of drivers [18]. The seamless integration of this technology in to the seat is seen below in Figure 2-7.



**Figure 2-7. Ford ECG Seat [18].
Used under fair use, 2012.**

With all of these noncontact electrodes being researched and evaluated, a group at IEEE developed a methodological review on dry electrode technologies. The study adds that wet electrodes are used almost universally in clinical applications today, provide an excellent signal, but are “cumbersome and irritating for mobile use” [19]. The group divides the performance of all electrodes in to two categories. First is the signal to noise ratio (SNR) of the device, and motion sensitivity. Second is the comfort of the device based on its interface to the skin, either by contact or proximity. Wet electrodes simply do not have the ability to transmit bioelectric signals through cotton clothing. Even a thin layer of cotton acts like a large resistor. The clothing introduces large noise signals, rendering wet electrodes useless in noncontact applications. Likewise, movement of the electrodes can cause an

artifact in the signal, creating bias. It was found that while dry electrodes allow the devices to be worn over clothing, they still often succumb to the same motion induced artifacts that wet electrodes do. The main way around this is to require a tight vest and chest band [19], in order to secure the electrodes in place, and reduce movement. This also leads in to the second issue with dry electrodes, of friction based artifacts from the sensors rubbing up against the clothing. The tighter the strap around the body, the less likely motion or friction induced artifacts will distort the signal. It can be concluded that while wet electrodes provide direct and accurate results, they were simply prone to issues with skin irritation, as well as motion induced noise. Meanwhile dry electrodes, still only provide accurate results after being applied within a micrometer of the skin, and with tendencies to fluctuate under motion and friction induced noise.

2.6.2 Non-contact Heart Monitoring Without Bioelectric Signals

While capacitively-coupled ECGs have been proven effective in strict test environments, they still require very close proximity to the patient. Often times an electrode based approach is simply not a practical method for gathering vital signs. However, there are very few devices that are capable of non-contact heart monitoring without the use of bioelectric signals.

One device that has been researched as a method for detecting vital signs without using bioelectric signals actually uses a similar approach. However, instead of trying to detect vital signs through clothing, they attempt to detect vital signs through vehicle tires. The project is aimed at helping homeland security through border control. The final criteria are that a vehicle must pull on to a place of highly sensitive pressure sensors, turn off their engine, and all occupants must get out of the vehicle. If a person is hiding inside the vehicle, then the pressure sensor is supposed to detect their heart beat or breathing through the vehicles tires. Their proposed method relies on pneumatics. A silicon tube with one end plugged by a highly sensitive pressure sensor and the other end capped is sandwiched between two rigid boards and placed on the ground at the entrance gate of the border. “When one wheel of the car is on the board and the engine is stopped, the pressure sensor can detect human vital signs such as the heartbeat, which cannot be concealed. Due to the high sensitivity of the pressure sensor, consideration was given to the effect of external disturbances such as ground vibration and wind force acting on the car” [20]. Specifically, the device consists of a low-frequency microphone with 10,000 mV/kPa sensitivity. This microphone listens for small changes in pressure in the silicon tubing underneath the vehicle. Diagrams of this technology are shown below in Figure 2-8 and Figure 2-9.

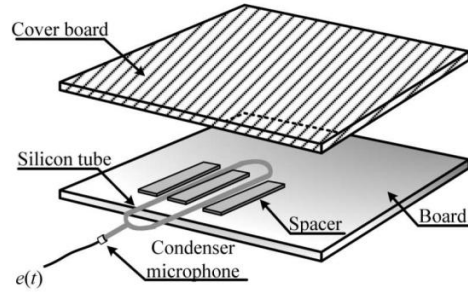


Figure 2-8. Development device for sensing a person hiding in a car [20].
Used under fair use, 2012.

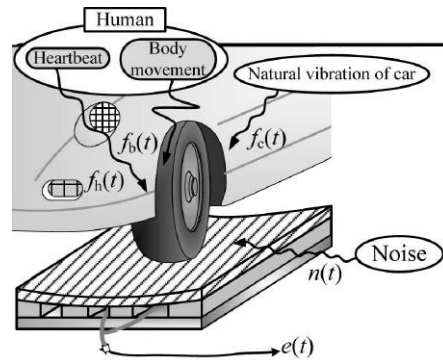


Figure 2-9. Illustration of how to sense a person hiding in a car [20].
Used under fair use, 2012.

The researcher makes the claim that they are able to filter out the 0.7 to 3.0 Hz vibration signals that is the common fundamental frequency range for an adult heart beat. The paper addresses the issues of wind and external forces by describing a technique for focusing on just the heart beat. They observed the higher harmonic components of 8 to 12 Hz [20] where there was less noise in the signal. After being placed through a bandpass filter, the component is “rectified and smoothed using a bandpass filter with a cutoff frequency of 0.5 and 2.0 Hz, which yields a shift back to the fundamental frequency range” [20].

What Hurikara fails to recognize is that by taking the original bandpass filter of 8 to 12 Hz, to look at the higher harmonic of the heart rate signal, then you are removing all lower frequency content from the signal. This means that the fundamental frequency of the person inside the vehicle would be removed because it is lower than the 8 Hz cutoff frequency. It is also impractical to utilize the higher harmonics of the vital signs signal, if you don’t know what the original fundamental frequency was to begin with. This is because simply choosing a harmonic does not imply that one knows how many times above the original fundamental frequency they are at. It may be any even or odd number times the original frequency. Also the reason Hurikara claims to have chosen the higher harmonics of the heart rate is to avoid the low level noise at the fundamental frequency range of the heart rate. However, if there is such significant noise, then there will be noise proportional to this at whatever higher harmonic range is chosen.

Yet another issue that is not even mentioned in Hurikara’s research is that the cushion, the suspension, and the tires underneath the vehicle are not designed to transmit energy directly from the

road to the passenger. Rather they are designed to dampen out vibrations that might interfere with the driving experience. It is highly unlikely that such low amplitude, low frequency vibration of a person's heart rate is going to be transmitted directly through the tires to the pressure pad. A method of monitoring heart rate through clothing rather than through a vehicle's tires will encounter less noise and therefore will require less signal processing than what Hurikara has proposed.

2.7 Summary of Current Technologies

With all of the technologies that have been explored, there is no single device like the one proposed. An affordable device that can be rapidly deployed in to a MCI, that is capable of being a force multiplier for a single EMT to monitor multiple casualties, would be a practical addition to the current market of health monitors. The method of using pressure transducers to accomplish this was the result of three main objectives. The first objective was that the device be sensitive enough to detect heart and respiratory rates over clothing. This ability would greatly reduce the time required for an emergency responder to administer the device. The second objective was that the device be compact and affordable. The more compact and affordable, the more of these devices an EMT could carry in to a MCI. Finally, the third objective was that the device be rugged. The more rugged the device, the more likely it can withstand the hazardous environments where it might be utilized.

Chapter 3: Methods

Now that existing techniques and research in heart and respiratory monitoring has been explored, a new method of using pressure transducers to detect both heart and respiratory rate will be explained, developed, and tested. Three stages of research were implemented in order to accomplish this. These stages were feasibility, simulated testing, and human testing. In feasibility, the concept undergoes its first reduction to practice. During the simulation stage, the technology is implemented on a printed circuit board (PCB), and then tested on a simulated breathing mannequin. Finally during the human testing stage, the device is refined in to a prototype and tested on humans in order to correlate its data with an industry standard ECG. Each stage involved materials, schematics, testing procedures, and goals that will be explained in detail. The final step is to understand the applicable signal processing techniques which will prove useful to analyze the testing and results. All of the details of the stages described as well as the signal processing involved will be explored in this section.

3.1 Feasibility

The initial key to the technology was to prove whether or not the combination of pressure transducers and flexible medical tubing can be used to adequately detect the desired signals. Preliminary tests had to be designed to validate this concept. By attaching a small bladder or piece of medical tubing to a pressure transducer, it is possible to create a movement sensor that is compact, rugged, and sensitive. The idea is simple. Power the pressure transducer, attach a piece of medical tubing, plug off its free end, and then place the apparatus against the chest. When a human breathes, the rise in volume in the lungs causes the chest cavity to expand, resulting in a displacement in the chest. You can think of the beating of the heart in the same way. When a human's heart beats, the result is a displacement in the chest, except usually at a much higher frequency and lower amplitude when compared to breathing. It was determined that it might be possible to detect these displacements through clothing, by pressing a piece of medical tubing to the chest, and monitoring the change in pressure in the tubing using a pressure transducer. The feasibility results will show that it is possible to extract heart and respiratory information using pressure transducers and tubing.

3.1.1 *Test and Materials*

With the conception of this idea came the need for a viability test. The experimental setup consisted of three main components. These were the pressure transducer, the flexible medical tubing, and an op-amp. The pressure transducer used was a Freescale temperature compensated and calibrated MPVZ4006G series integrated silicon gauge pressure sensor with on-chip signal conditioning. This sensor had a sensitivity of 766 mV/kPa. The nozzle was attached to one end of a section of common medical tubing. Next, a 30 gain op-amp circuit was created with adjustable offset voltage. Once the circuit was properly configured, the tubing was placed inside a strip of elastic medical wrap. This medical wrap was then placed around the chest of a test subject. Images of this assembly can be seen in Figure 3-1.



Figure 3-1. Feasibility test setup.
Photo by author, 2012.

Next, a National Instruments LabVIEW Virtual Instrument (VI) was programmed to take in the signal from the NI-USB-6009 device. The resulting signals were recorded and observed in the frequency domain using MATLAB. Several sets of data were taken, most placing the chest sensor right over the area of maximum cardiac impulse, and others with the chest strap at rest on the table. The results from this test were successful in validating that heart and respiratory signals can be observed using pressure transducers.

3.1.2 Feasibility Breadboard Circuit

The breadboard circuit that was used to read in the pressure transducer results and amplify the signal was made up of multiple components. These included the pressure transducer, resistors, and an operational amplifier arranged to be inverting. The resistor and capacitor values were chosen based on the desired gain values. The operational amplifier (op-amp) was a National Semiconductor LMC6482. This rail-to-rail op-amp operates at an ultra low input current and is capable of a high voltage gain. The pressure transducer's datasheet specified a voltage offset of 0.2 V, and therefore a differential amplifier circuit was designed to reduce that offset to zero. The resulting circuit can be seen below. The circuit used is seen below in Figure 3-2.

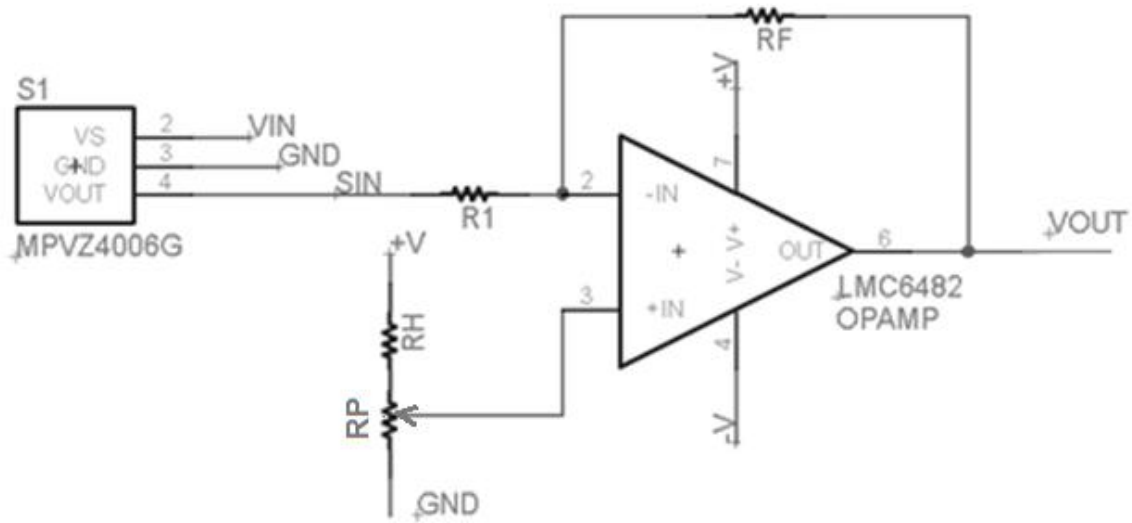


Figure 3-2. Feasibility amplifier circuit.

Note that the resistor values are not shown in this figure, but were chosen based on the 5 V rail of the NI-USB-6009 used to read in the VOUT analog signal in to the computer. The resulting equation for this circuit can be seen below in Equation 3-1.

$$V_{OUT} = \left(V_{IN} - (V_+) \left(\frac{R_P}{R_H} \right) \right) \left(\frac{R_F}{R_1} \right) \quad \text{Equation 3-1}$$

The breadboard circuit resulted in the feasibility of finding both heart and respiratory rate in one signal using pressure transducers. The feasibility results were verified by comparing the autospectrum output in MATLAB against a pulse oximeter and manual estimations as seen in the next section.

3.1.3 Feasibility Results

The feasibility of using pressure transducers to detect movement was immediately successful. The NI-USB-6009 recorded data at 100 Hz for around 35 seconds using LabVIEW. The data was then imported in to MATLAB for signal processing. The data was disrupted by movement at the beginning and end of data collection, and as previously discussed a 30 second block of data was separated for evaluation. The resulting normalized data results can be seen below in Figure 3-3.

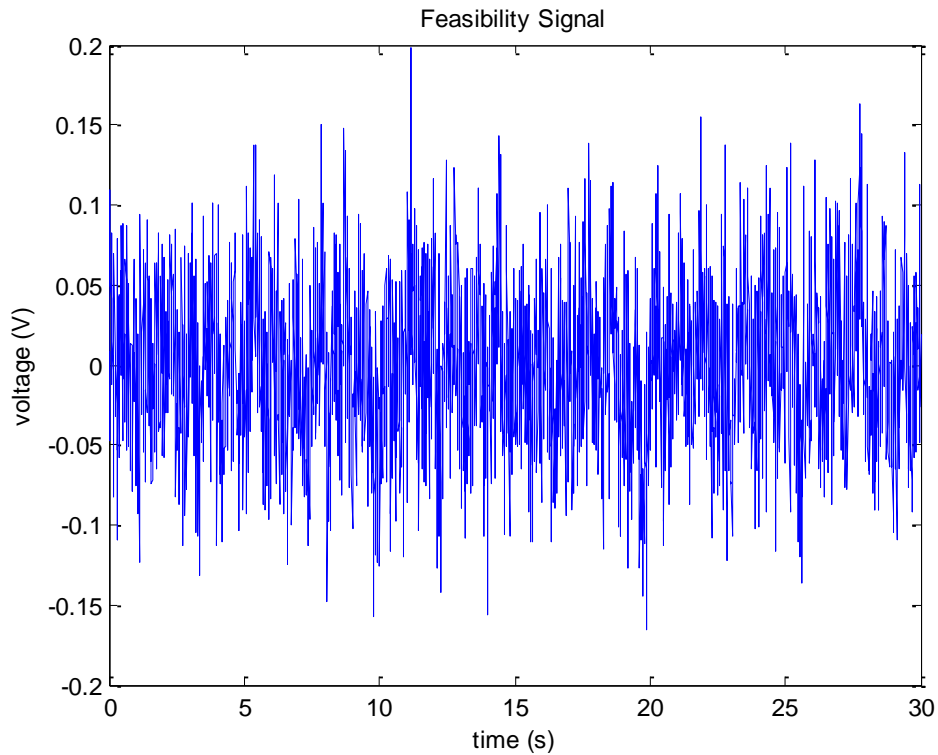


Figure 3-3. Feasibility normalized data results in the time domain.

The feasibility normalized data results show that the maximum amplitude of the chest sensor signal when attached to a person was around 0.35 V. This data, which was originally a 3000 point array, was broken up in to 5 averages, made up of 600 point arrays. Each sample record was passed through an FFT, to create a discrete Fourier transform array. Each DFT was then multiplied by its complex conjugate, in order to achieve the auto spectrum for that sample record. These results were summed up to create an ensemble. Once all sampler records had been evaluated, the results were averaged. The final result is the autospectrum, G_{xx} for that set of data.

All frequency domain results are only valid up to the Nyquist frequency, which was 50 Hz in this case. However, the relevant data for the desired signals in this case was well below 50 Hz. Therefore, the first plot will be shown up to the Nyquist, while subsequent plots will only be shown up to 4 Hz.

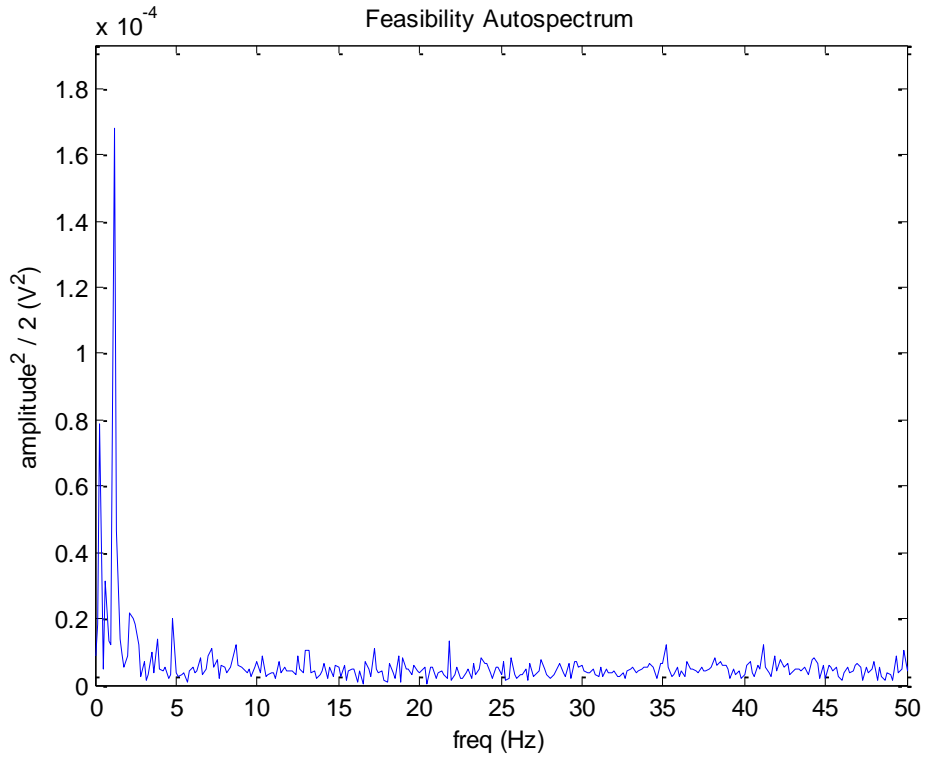


Figure 3-4. Feasibility autospectrum results up to the 50 Hz Nyquist frequency.

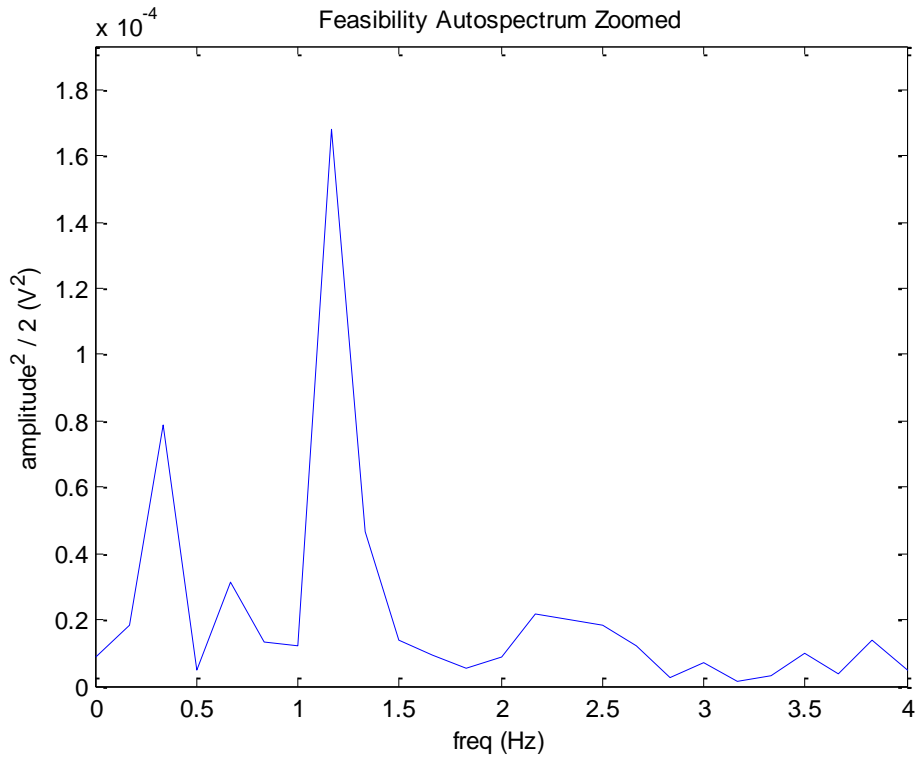


Figure 3-5. Feasibility autospectrum results up to 4 Hz.

Note that the autospectrum results show observed frequencies at 0.3333 and 1.167 Hz. These spikes correspond with 20 RR and 70 HR accordingly. This was verified using a pulse oximeter and manual estimation at the time that the person was being tested. The expected values were 20 RR and 70 HR, which matched with the autospectrum results. The noise in the autospectrum can be compared to the signal using an SNR estimate.

Here, the SNR will be defined as the ratio of the RMS value of the signal to the RMS value of the noise floor. The autospectrum results are already in terms of power, V^2 , so the SNR can be estimated by taking the square root of the magnitude for the signal divided by the square root of the mean magnitude for the noise. This is seen below in Equation 3-2.

$$SNR = \left(\frac{RMS_{signal}}{RMS_{noise}} \right) = \sqrt{\left(\frac{Power_{signal}}{Power_{noise}} \right)} \quad \text{Equation 3-2}$$

The resulting SNR value for the respiration signal during feasibility testing was 4.014, while the SNR for heart rate was 5.866. The SNR during feasibility testing was not expected to be very high, due to the fact that the signals were found and amplified using a breadboard circuit and through-hole components. However, it is important to note that any future circuits should be compared for effectiveness against these initial SNR results. The success of the feasibility tests resulted in the push towards a second generation concept, to be used in a simulated environment.

3.1.4 *Advancements Post Feasibility Tests*

The road towards a working prototype involved further research in several areas, even as early on as the feasibility stage. Firstly, there needed to be a model for the relationship between pressure in tubing and the expected compression of the chest. Once the pressure transducer and tubing could be properly specified, a PCB would have to be designed with built in op-amp. This intermediate device would act as a simulation, so that further data could be gathered. The Carilion Center for Experiential Learning at Carilion Roanoke Memorial Hospital was specified as an appropriate location by the Pediatric Medical Device Institute. This location possesses a simulated breathing mannequin that was ideal for data collection.

3.2 Simulation

In order to develop the best MCI force multiplier, several questions must be answered. Firstly, what kind of amplitude should be expected from the heart or from the respiratory signals on the human when observed through clothing? Next, what pressure sensors and data acquisition tools are capable of detecting such a response? Once these questions have been answered, and the components are arranged on a PCB, the device must be tested. It was for the purposes such testing, and to prove the model, that the device was taken to Carilion Center for Experiential Learning, to be tested on a simulated breathing mannequin.

3.2.1 Modeling the Change in Volume with the Change in Pressure in the Tube

Understanding the physics and the relationship between volume and pressure inside the tubing is a necessary step prior to choosing the tubing itself. The impulse of the heart or respiratory rate is transmitted through the clothing and in to the tubing, which then compresses, decreasing its air volume, and increasing its air pressure, which then causes an increase in voltage reading from the pressure transducer. Figure 3-6 below shows a simple illustration of how the impulse causes a decrease in volume in the tubing.

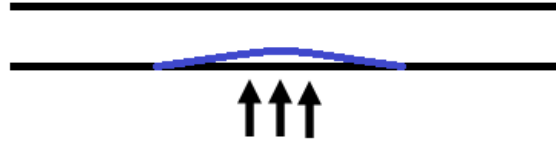


Figure 3-6. Visual interpretation of tubing being compressed by an impulse.

Note that the final relationship or equation can be determined through several basic equations. Boyle's law states that so long as temperature remains constant, pressure and volume remain inversely proportional to each other. This can be seen in the equations below.

$$pV = k \quad \text{Equation 3-3}$$

$$p_1V_1 = p_2V_2 \quad \text{Equation 3-4}$$

Following this relationship, one must know what the air volume in the tubing is. The volume of a cylinder is the product of inner circular area and the length of the tubing as seen in the equation below.

$$V_1 = \pi r^2 L \quad \text{Equation 3-5}$$

Assuming that the deflection of a specific length of tubing causes a change in radius, Δr , over a small tube length, L_x , then the change in volume can be calculated as seen in the following equations.

$$\Delta V = (\pi(\Delta r)^2)L_x \quad \text{Equation 3-6}$$

$$V_2 = V_1 - \Delta V = \pi r^2 L - (\pi(\Delta r)^2)L_x \quad \text{Equation 3-7}$$

Combining Equation 3-4, Equation 3-5, and Equation 3-6, it is possible to approximate the change in pressure in the tube as shown below. P_1 is assumed to be 100 kPa, or near atmospheric pressure.

$$100 (kPa) \cdot \pi r^2 L (cm^3) = p_2 (kPa) \cdot (\pi r^2 L - (\pi(\Delta r)^2)L_x) (cm^3) \quad \text{Equation 3-8}$$

$$p_2 (kPa) = \frac{(100 kPa)(\pi r^2 L cm^3)}{(\pi r^2 L - (\pi(\Delta r)^2)L_x cm^3)} \quad \text{Equation 3-9}$$

$$\Delta p (kPa) = \frac{(100 kPa)(\pi r^2 L cm^3)}{(\pi r^2 L - (\pi(\Delta r)^2)L_x cm^3)} - 100 kPa \quad \text{Equation 3-10}$$

Based on these equations it is possible to estimate what kind of changes in pressure to expect from compressing the tubing. However, before calculating the expected change in pressure, it is necessary to understand the anatomy of heart and respiratory rates, and their impact on the chest cavity.

3.2.2 *Anatomy of Heart and Respiratory Signals to the Chest*

The transfer of motion that is seen in the human chest is known as the apex beat, cardiac impulse, or the apical thrust. It is created by the left ventricular contraction of the heart. It is located at the lowest point on the chest at which the contractions of the heart can be felt. While this impulse is usually the point of maximum impulse, a diseased heart will often have pulsations correlating with other problems such as an enlarged right ventricle. It may also be noted that it might not be observable in the supine position in some subjects [21]. Regardless, the expected peak to peak displacement of the chest at the area of maximum cardiac impulse is around 0.3 mm [22].

Similar to heart rate, but with larger amplitude, and at a lower frequency, is respiratory rate. When humans breathe, the chest cavity expands, allowing the flow of oxygen in to the lungs. The peak to peak displacement of the chest due to respiration is between 4 and 12 mm [22].

3.2.3 *Application of the Model to the Transducer and Tubing Specifications*

Based on the information about the anatomy of the vital signs, it is possible to define the appropriate tubing as well as define the specifications for the pressure transducer. When it comes to tubing, a material that is malleable enough to displace without requiring a large amount of force, will allow the heart and respiratory signals to be more easily observed. A material, such as super-soft latex, is often used in medical applications, and fits this requirement. The first tubing ordered was super-soft latex rubber tubing with a 1/4" outer diameter, a 1/8" inner diameter, and a 1" bending radius.

The pressure transducer would have to be capable of observing changes in pressure as low as those defined previously in Equation 3-10. Plugging in approximate values for the tubing should give a good idea of the expected changes in pressure. The table below is composed of values based on a 10" strip of Tygon® super-soft latex rubber tubing. This table will be used to estimate the pressure change based on heart and respiratory impulses.

Table 1. Approximate values for estimating pressure change.

Parameter:	Description:	Value (Units):
r	Tubing Inner Radius	6.350 mm
L	Overall Tubing Length	254.0 mm
Δd (H)	Change in Inner Radius	0.3 mm
Δd (R)	Change in Inner Radius	4 mm
Δr (H)	Change in Inner Radius	0.15 mm
Δr (R)	Change in Inner Radius	2 mm
Lx (H)	Length of Tubing Impulse Effects	20.000 mm
Lx (R)	Length of Tubing Impulse Effects	100.000 mm

Note that the change in inner radius of the tube was determined to be equal to half of the displacement of the chest. Cutting the displacement in half allowed for the rest of the design process to actually take into account a lower level of pressure change than what actually might be expected. Also the total length of tubing that is affected by the heart signal is 20 mm for heart rate, and 100 mm for respiratory rate. This is based on the 10" tubing being configured in a spiral form factor and placed against the chest near the epicenter of the heart impulse, as approximated in Figure 3-7 below.

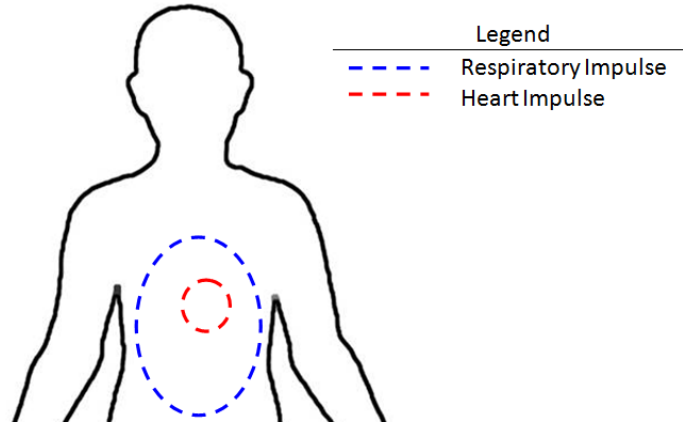


Figure 3-7. Approximate epicenters for heart and respiratory impulse not drawn to scale.

Plugging the values from Table 1 in to Equation 3-10, we get the following pressure changes, first for heart rate and second for respiratory rate.

$$\Delta p \text{ (kPa)} = \frac{(100 \text{ kPa})(\pi(6.35)^2 254 \text{ mm}^3)}{(\pi(6.35)^2 254 \text{ mm}^3 - (\pi(0.15)^2)(20) \text{ mm}^3)} - 100 \text{ kPa} \quad \text{Equation 3-11}$$

$$= 0.004394 \text{ kPa}$$

$$\Delta p \text{ (kPa)} = \frac{(100 \text{ kPa})(\pi(6.35)^2 254 \text{ mm}^3)}{(\pi(6.35)^2 254 \text{ mm}^3 - (\pi(2.0)^2)(40) \text{ mm}^3)} - 100 \text{ kPa} \quad \text{Equation 3-12}$$

$$= 1.587 \text{ kPa}$$

Such low pressure changes are going to require a high sensitivity pressure transducer. The equation below shows the impact of sensitivity, S , and pressure change, ΔP , on voltage response.

$$\Delta V = \Delta P(S) \quad \text{Equation 3-13}$$

This equation provides a simple calculation for the transducer response. For example, a pressure transducer with 100 mV/kPa will only show a 0.4394 mV raw peak to peak voltage change for a heart rate response, before any amplification. The table below shows several of the options for higher sensitivity pressure transducers. Included in the table are their corresponding supply voltages, sensitivities, and accuracies.

Table 2. Pressure transducer metrics.

Pressure Transducer:	Supply Voltage (V):	Sensitivity (mV/kPa):	Accuracy (V):
Freescale Semiconductor MPVZ4006G	5	766	0.230
Freescale Semiconductor MP3V5004G	3	600	0.027
Freescale Semiconductor MPX5010	5	450	0.225

While not the highest sensitivity, the MP3V5004G has the finest accuracy combined with a 3 V supply, which is ideal for embedded applications that run off of the supply voltage. Meanwhile, ignoring the supply voltage and accuracy, the MPVZ4006G has the highest sensitivity. However, of these sensors, the MPX5010 was the one chosen for simulation. This sensor had a sensitivity that was proven to be sufficient by comparing what the expected voltage output would be versus the resolution of the DAQ.

3.2.4 Data Acquisition and Gains

The data acquisition device that was chosen for initial data collection was the same NI-USB-6009 that was chosen for feasibility testing. This device has a 14 bit resolution over a 5 V range. The resulting DAQ voltage resolution can be calculated as seen below.

$$V_{res} = \frac{(V_{range})}{(2^{\# bits})} = \frac{(5 V)}{(2^{14})} = 0.3052 mV \quad \text{Equation 3-14}$$

The 450 mV/kPa pressure transducer that was chosen must be verified by comparing the voltage response against the DAQ's voltage resolution of 0.3052 mV. Plugging in the 450 mV/kPa sensitivity and the change in pressure from Equation 3-11 and Equation 3-12, in to Equation 3-13, provides voltage response for heart and respiratory rate as shown below.

$$\Delta V = (0.004394 kPa)(450 mV/kPa) = 1.977 mV \quad \text{Equation 3-15}$$

$$\Delta V = (1.587 kPa)(450 mV/kPa) = 714.2 mV \quad \text{Equation 3-16}$$

This voltage response due to the expected change in voltage due to cardiac impulse is 6 times the voltage resolution of the DAQ. This means that the resolution of the DAQ is capable of reading in the low voltage change of the pressure transducer due to heart rate, without having significant interference from noise. Recall that when advancing towards a final embedded system prototype, the MP3V5004G is more practical due to the 3 V supply and 600 mV/kPa sensitivity. Therefore, the same calculations that were just discussed on the MPVZ4006G will be re-evaluated when discussing the design specifications for the embedded prototype board.

In addition to having a low enough resolution, the DAQ also has a maximum possible voltage input. The NI-USB-6009 that was chosen has a 5 V supply voltage, and therefore the signal that is being read in by the DAQ must not be larger than 5 V. Anything above this rail would cause the DAQ to become saturated and the output would no longer be meaningful. This means that the amplifier circuit that was shown previously must not have a gain that would cause the output voltage in to the DAQ to be higher than 5 V.

3.2.5 Signal Processing Considerations

One of the other areas that must be explored prior to building the first PCB is to thoroughly understand which signal processing technique is most applicable based on what kind of data is expected. It has been shown that the expected data should be visible by the DAQ. A technique that can show what frequency signals are embedded in the data.

An autospectrum was the ideal method to accomplish this task. The next step is to discover what resolution signal is sufficient for the simulation device. After consulting several doctors, it was determined that an accuracy of 2 bpm, would be more than sufficient. The amount of time (T) that would be required based on this frequency resolution is shown below in Equation 3-17. Likewise the actual time that would be required to acquire a 2 bpm resolution would be 30 seconds, as shown in Equation 3-18.

$$\frac{1}{T} = \text{frequency resolution} \quad \text{Equation 3-17}$$

$$T = \frac{1}{2 \left(\frac{\text{beats}}{\text{min}} \right) \frac{1}{60} \left(\frac{\text{min}}{\text{sec}} \right)} = 30 \text{ seconds} \quad \text{Equation 3-18}$$

The 30 seconds worth of data proved to be less than sufficient, as the act of averaging the results would lower this frequency resolution from 2 bpm to 6 bpm. However, this will be discussed in further detail in the results section. Before designing the experiment, the materials for the PCB had to be chosen.

3.2.6 Circuit Board and Materials

As with all electronics based platforms, the first revision of a PCB was the next logical step after feasibility testing. As expected, the transition from breadboard to surface mount parts improved the SNR as well as improved the overall reliability of the electronics.

The PCB that was designed included a revised differential amplifier circuit. There were several factors that led to this revised version. The number one reason was to improve the signal to noise ratio. However, another important change was to remove the DC component from the signal. This DC component or slow changes in voltage were due to a drift in atmospheric pressure over time, combined with pressure escaping from the tubing over time. Recall that the tubing is plugged off at its free end, yet it is not perfectly sealed. Therefore over time, the overall pressure in the tubing would gradually drift, and begin to reach the rail voltage of the op-amp and saturate the signal. Since this drift was so slow, adding a capacitor on the output side of the op-amp, would remove this unwanted DC component. The op-amp used was a Texas Instruments OPA140 high-precision, low-noise, rail-to-rail output op-amp. The original pressure sensor was replaced with a new board mount version. This new pressure transducer was an on-chip signal conditioned, temperature compensated and calibrated, Freescale Semiconductor MPVZ4006G Integrated Silicon Gauge Pressure Sensor. It had a sensitivity of 450 mV/kPa as previously discussed. The PCB was created using EAGLE, as a schematic and then routed as a board. The schematic utilized the same amplifier circuit shown previously in Figure 3-2, except with the

capacitor in between the previous voltage output, and the new voltage output. The schematic for this simulation board can be seen below in Figure 3-8.

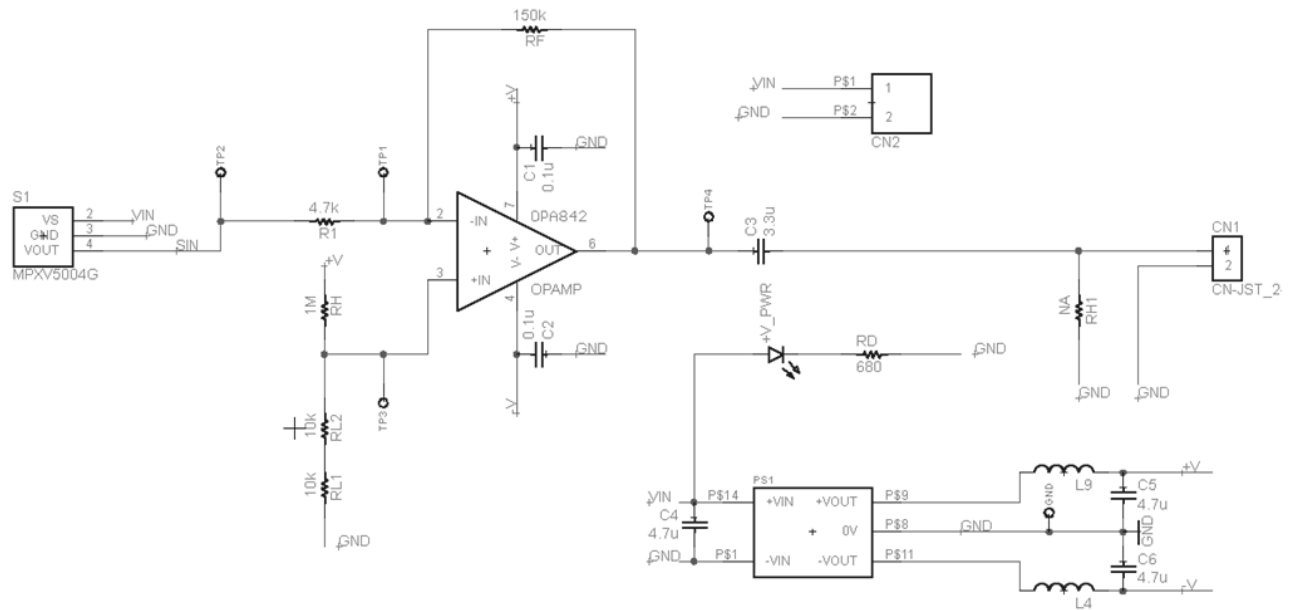


Figure 3-8. Simulation PCB schematic.

The board that resulted from this schematic can be seen below in Figure 3-9.



**Figure 3-9. Simulation PCB after populating the components.
Photo by author, 2012.**

Along with all the new components for the board, came the need to find tubing that would provide a highly sensitive output when placed on the chest. As described, it was determined that Tygon® latex tubing would be sufficient. It is flexible enough to be placed in various shapes without blocking the flow of air and pressure in the tubing. In addition it is elastic enough to deform under small forces such as those created by heart and respiratory response in the chest. In order to create a versatile and durable

system that could endure long periods of testing and analysis, the board was attached to a section of fiber glass just long enough to hold the PCB and a certain length of Tygon® tubing. The resulting simulation device is seen below in Figure 3-10.

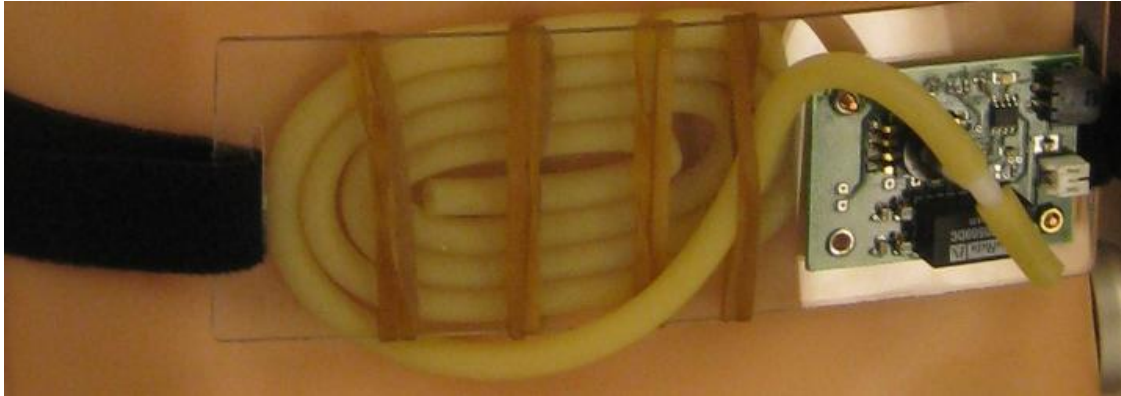


Figure 3-10. Simulation device with tubing and chest strap.
Photo by author, 2012.

3.2.7 Simulation Experiment

The device was taken to the Carilion Center for Experiential Learning at Carilion Roanoke Memorial Hospital, in order to compare its response to that of an actual simulated breathing mannequin. The simulated breathing mannequin is capable of outputting a programmable respiratory rate through a bladder in the mannequin's chest. However, it is incapable of providing heart rate in the same manner. Note that the experimental setup can be seen below in Figure 3-11, with the device placed right underneath the left breast of the mannequin.

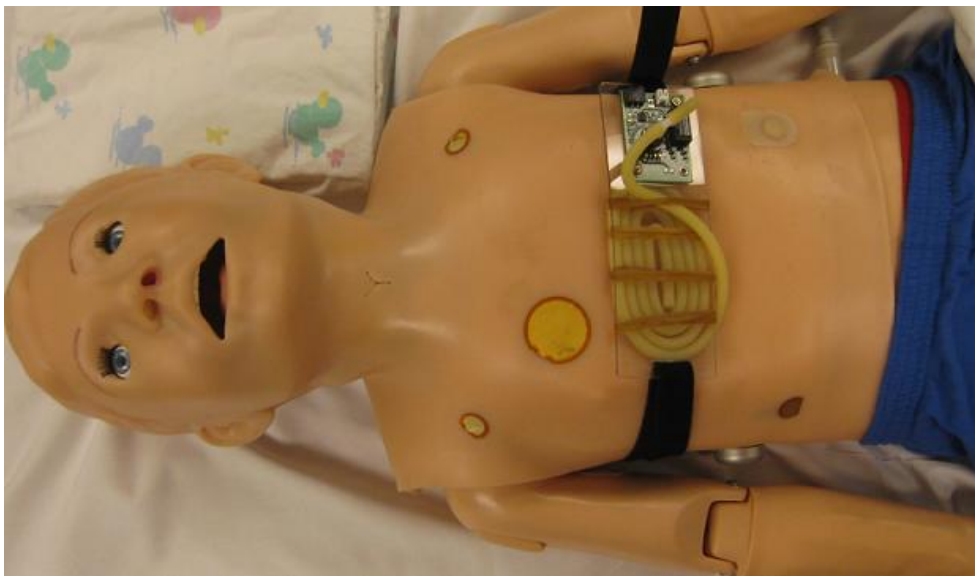


Figure 3-11. Simulation device on the simulated breathing mannequin's chest.
Photo by author, 2012.

The device was attached using a Velcro strap, and measurements were taken for six different simulated respiratory rates. A computer program at the test site was used to dial up the respiratory rates for at least thirty seconds for each of the six simulated respiratory rates. The data from the op-amp circuit was read through the NI-USB-6009 as previously discussed.

Data was recorded, and then its autospectrum was calculated in MATLAB. These autospectrum results were created using 30 seconds worth of data as previously discussed. The results suggested that there was correlation between the estimated breathing rate and the actual simulated breathing rate. It should be noted that the simulated breathing mannequin did not have the ability to emit a heart rate in addition to respiratory rate. Therefore, an experiment approved for human testing, would have to be developed to test the device for correlation with heart rate. In addition, qualitative assessment of the autospectrum does not provide the quantitative results needed, to conclude that this device is actually doing what it is intended to do. Having an industry standard tool, such as an ECG machine, would allow for such a quantitative assessment of the results through a cross spectrum and through coherence function analysis.

3.2.8 Need for Improvements

Despite the success of the simulation experiment, there was still a need for improvements to the PCB. Specifically, the SNR was low as will be shown. The changes that needed to be made in order to improve the SNR are to move to a higher sensitivity pressure transducer, or a higher resolution ADC. Another issue was that the tubing being used was not completely sealed at the end attached to the pressure transducer. This allowed air to often escape the tubing during compression, causing an increase in pressure over time. The result was a built up of pressure over time in the tubing. To solve this, the AC component would have to be decoupled from the signal using a high pass filter. With the need for all these improvements, came the transition to a full system prototype.

3.3 Full System Prototype

With the need for improvements to the hardware and software, came the next revolution of the design process. Firstly, new sensors needed to be specified, and a new more versatile and durable design would have to be incorporated. The first prototype should also have advanced software with real time capabilities. This meant that a new fully embedded prototype would have to be created that combines both data acquisition and analysis.

3.3.1 Overall Design Considerations

Given the fact that the next board would be the first embedded prototype, there were several changes that needed to be considered. In mass casualty situations, it has been shown that a durable tool that can act as a force multiplier would be most useful. It was also mentioned that the device must be capable of being utilized in multiple ways. Three ways that would be easiest for an EMT to communicate with the system would be through flashing LED's that blink with the heart and respiratory rates, an LCD screen that actively displays the rates via text, and finally a wireless monitoring capability

such as Bluetooth®. Outside of communication, the device must be small, easily deployable, and adjustable to different body types.

All of these considerations must be taken in to account prior to choosing the actual hardware that the prototype will be comprised of. Therefore, all software, sensors, and materials chosen must develop through these needs. Note that all of the individual components of the PCB are listed along with their part number and a URL to their corresponding datasheets are shown in Appendix F: Miscellaneous.

3.3.2 *Software and Sensors Chosen*

LabVIEW provides a great environment for coding techniques that are both rapid and versatile. A microcontroller that is capable of being programmed through LabVIEW, while at the same time having all of the requirements to control LED's, an LCD screen, a Bluetooth® device, and a pressure sensor is a requirement. In addition, if the microcontroller does not already have an analog-to-digital converter (ADC) with the required voltage resolution, then an external ADC must be specified.

The microcontroller chosen was the Texas Instruments LM3S8962. It utilizes a compact design, running on a 3.3 Volt supply. It has two serial inputs, JTAG debugging capabilities, can operate multiple inter-integrated circuit (I2C) devices, and has several 10 bit ADC inputs over a 3 V supply reference, as well as numerous general purpose input outputs (GPIO). Based on the resolution and reference, the resulting microcontroller voltage resolution can be calculated as seen below.

$$V_{res} = \frac{(V_{range})}{(2^{\# \text{ bits}})} = \frac{(3 \text{ V})}{(2^{10})} = 2.930 \text{ mV} \quad \text{Equation 3-19}$$

The new pressure transducer that was chosen was the Freescale Semiconductor MP3V5004G that was previously specified, has a 600 mV/kPa sensitivity, and a 3 V supply voltage, which matches that of the microcontroller. Plugging in the 600 mV/kPa sensitivity and the changes in pressure from Equation 3-11 and Equation 3-12, in to Equation 3-13, provides the following voltage responses for heart and respiratory rate.

$$\Delta V = (0.004394 \text{ kPa})(600 \text{ mV/kPa}) = 2.636 \text{ mV} \quad \text{Equation 3-20}$$

$$\Delta V = (1.587 \text{ kPa})(600 \text{ mV/kPa}) = 952.2 \text{ mV} \quad \text{Equation 3-21}$$

The voltage response due to cardiac impulse is actually less than the resolution of the microcontroller's ADC. This means that the resolution of the microcontroller is definitely not capable of reading in the low voltage change of the pressure transducer due to cardiac impulse.

An external ADC must be found that is capable of detecting the 2.636 mV cardiac impulse signal, in full, with a high SNR. Most ADC's afford about 2 bits just to noise, so at least a 16 bit ADC would be practical. In addition an ADC that can be coded in I2C, and provides an easy way for the microcontroller to communicate with the ADC without interfering with other devices, would be ideal. One such ADC that matches these requirements, is the LTC2471 made by Linear Technologies. With a small profile, a

1.25 V built in voltage reference, a 3 V supply voltage, a 1 kHz maximum sampling rate, and a 16 bit resolution. Its specifications match all of the needs for the prototype. The 2.63 mV response due to heart rate was 138 times the resolution of this ADC as calculated in Equation 3-22 below.

$$V_{res} = \frac{(V_{range})}{(2^{\#bits})} = \frac{(1.25 V)}{(2^{16})} = 0.01907 mV \quad \text{Equation 3-22}$$

This resolution is small enough to detect the heart rate signals. The improved resolution will be combined with a more advanced amplifier circuit. This new circuit will utilize the Texas Instruments OPA2301 low noise, high-speed, low single supply power rating, 16-bit resolution, op-amp. The 16-bit resolution matched the voltage resolution of the LTC2471 ADC.

Once the data is gathered properly, software will be used to calculate the estimated heart and respiratory rates, which will then be displayed to the user on an LCD screen. The LCD screen chosen was the Newhaven Display NHD-C0216. It was a 2 lines, 16 character, transfective, white LED backlit display with a 3 V supply voltage. This screen was chosen mostly due to its small form factor. While most embedded hobby LCD screens are around 1 cm thick, this one is 5.5 mm thick, and the screen takes up most of the front of the body making it much less bulky than the average LCD screen. Also note that this LCD screen can be programmed through the I2C interface in the microcontroller.

3.3.3 Ergonomics

The first full system prototype needs to be developed, keeping in mind that it needs to be something an EMT can easily pick up and use in any emergency response situation. There are several major factors that go in to making this device ergonomic.

Firstly, it must fit in the palm of the user’s hand. Second it must be lightweight, such that several can be carried on to the field. However, it must also have a firm feel, such that users do not drop it or have trouble strapping the device around a patient. Another requirement is that the chest strap does not get in the way of the emergency technician during triaging of the patient. The first prototype was created, taking in to account these requirements, which were delegated in to the following design constraints.

Table 3. Prototype design constraints.

Constraint:	Parameter:	Target:
Fits in the palm	Size (W x L x D) in.	2 x 3 x 1 (inches)
Lightweight but Firm	Weight (Min < Max) oz.	4 < 8 (ounces)

These design constraints were chosen based on common handheld devices such as the Apple® iPhone series and the Motorola® Droid series, which are common devices that people are used to holding for prolonged periods of time. This device will likely be bulkier than the common Smartphone upon first revision, due to the need to include a bladder, battery, and circuit board, all on top of one another.

3.3.4 *Power Requirements and Scheme*

Similarly to ergonomics, the power requirements are of great importance when designing the first full system prototype. From the on/off mechanism to battery life, everything must be considered and evaluated against how this device is intended to be used.

Most modern devices have a power scheme based around a switch mode charge manager. This manager is what allows a device to be powered by an AC adapter, which simultaneously charges a battery in the device. In this way, when the device is unplugged, it can be used for a given amount of time on battery life, before needing to be recharged. The first step in developing the power scheme for this device was to choose a switch mode charge manager, a voltage converter, a power switch, a low power voltage detector, and then to specify all of the current draws of each component on the board.

The switch mode charge manager that was chosen for this device was the BQ24123 by Texas Instruments. It was chosen because it is suitable for 1-, 2-, or 3-Cell Li-Ion and Li-Polymer Battery Packs. It includes battery temperature monitoring, and an automatic sleep mode for low power consumption. In addition it is small, cost effective, requires a low 3.5 uA supply, and includes a “charging, adapter present, and done charging” pins for LED indicators. The BQ24123 on its own is not capable of providing all of the power scheme functionality of modern devices. Rather it works in conjunction with an on/off controller to allow the device to be easily enabled and disabled at the push of a button. The charge manager also requires a step down converter from the 5 V lines of the adapter and battery, to the 3.3 V lines that the microcontroller and other components operate on.

The on/off controller chosen was the MAX16054 by Maxim. It is a pushbutton on/off controller with a single switch debouncer and built-in latch. It requires a 3 volt single supply, and operates at 7 uA. It is intended to be used in conjunction with a step down converter to enable and disable the power at any given time. The step down converter chosen was the TPS62203 step-down or buck converter, by Texas Instruments. It operates from a standard 5 V rail, and steps down to 3.3 V output. Pulling this device’s enable pin to ground causes it to go into shutdown mode. Conversely, pulling the enable pin high causes it to start back up again. This will be a useful part of the power schematic.

All together, the system follows a block diagram operation for power operation based around the enable pin on the TPS62203. There are two methods by which the TPS62203 can be told to disable the power supply from reaching the rest of the circuit. Firstly an STM1061 low voltage detectors by ST Microelectronics, checks to see if the battery voltage is at dangerously low levels. The detector chosen considers anything below, 3.4 V, to be dangerous. When the battery voltage goes below a 3.4 V threshold, its output signal goes low. Secondly, the MAX16054 checks for whether or not the power button has been turned on or not. Whenever the power button is pressed, then “in” pin receives a momentary low pulse, causing the output logic to invert. Based on this, the desired logic output to the TPS62203 is such that the enable pin should be low, if either the MAX16054 outputs low, or the STM1061 outputs low. The desired logic conditioner is therefore an AND gate, as seen in the table below.

Table 4. Logic conditioning for an AND gate.

INPUT A:	INPUT B:	OUTPUT Y:
0	0	0
0	1	0
1	0	0
1	1	1

An SN74AHC1G08, Single 2-Input Positive-AND gate by Texas Instruments was chosen to calculate this logic prior to the enable pin on the voltage converter. The resulting block diagram for power operation in the full system prototype is seen below.

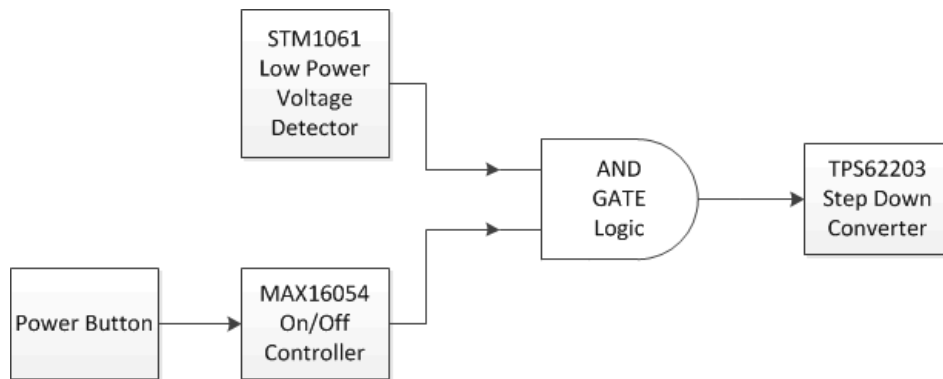


Figure 3-12. Prototype block diagram for power operation.

With all of these pieces of the puzzle coming together to create a full system prototype, a battery must be chosen based on the current draw of the system as a whole. The resulting table shows each component listed thus far, and their corresponding current draw.

Table 5. Prototype PCB current draw by component.

Component:	Description:	Current Draw:
BQ24123	Switch Mode Charge Manager	3.5 uA
LM3S8962	Microcontroller	100 mA
LTC2471	Analog-to-Digital Converter	3.5 mA
MAX16054	On/Off Controller	7 uA
MP3V5004G	Pressure Transducer	10 mA
NHD-C0216	LCD screen	20.5 mA
OPA2301	Op-Amp	9.5 mA
SPBT2532C2	Bluetooth® Module	40 mA
STM1061	Low Power Voltage Detector	0.9 uA
TPS62203	Step Down Converter	15 uA

The equation for required battery capacity can be found by summing these current draws and multiplying by the voltage supply, divided by the battery voltage multiplied by the efficiency of the

voltage converter. This equation and its subsequent calculations can be seen below in Equation 3-23 and Equation 3-24.

$$mAhrs\ Required = \frac{(24\ hours)(\sum I's)(V_{supply})}{(efficiency)(V_{battery})} \text{ Amp Hours} \quad \text{Equation 3-23}$$

$$mAhrs\ Required = \frac{(24\ hours)(0.1835\ Amps)(3.3\ Volts)}{(0.9)(3.7\ Volts)} = 4364\ mAhrs \quad \text{Equation 3-24}$$

Repetitive calculations for this value were made for different operating ranges, as seen in Appendix B: MATLAB Code under Battery Requirements for Full System Prototype. Also note that the efficiency from Equation 3-23, refers to the efficiency of the step down converter, which is 90% at a 3.3 V operating voltage.

The BQ24123 charge manager schematic chosen, and the TPS62203 step down converter, dictate that the ideal battery is a 1-cell, 3.7 volt battery. Since 24 hours of operation requires over 4364 mAhrs, then 8 hours of operation should require a 1456 mAhr battery. Common 1-cell, 3.7 volt devices are for handheld devices, and have around 1000 mAhr capacities. Taking this in to account, a battery was selected that was 3.7 volt, single cell, and had between 1000 and 1456 mAhr capacity. The NuPower Replacement Rechargeable Battery for Apple iPod 4th Generation Models was a capable 3.7 volt, single cell battery, with a 1350 mAhr capacity.

3.3.5 Product Design

When creating a full system prototype, the device must be something that is both familiar and ergonomic. Based on the number of components, a form factor the size of the average smart phone is more than practical. Knowing some of the components, a rough sketch was created as a design target for the full system prototype to be modeled after. The resulting sketch provided an idea of the end product, which is invaluable to circuit board design for a full prototype. This design can be seen below in Figure 3-13. Other renditions of the product design can be seen in Appendix F: Miscellaneous.

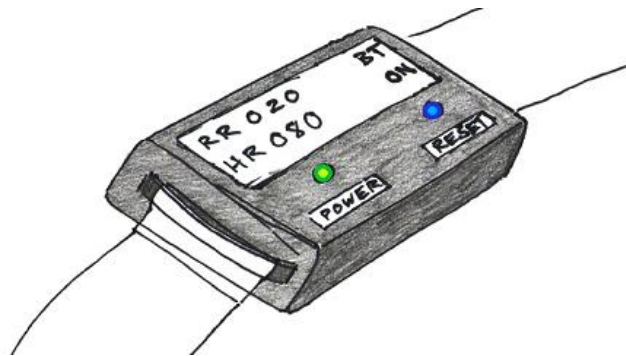


Figure 3-13. A rendition of the prototype targeted appearance.

Most smart phones fall well within the 5 x 3 inch range previously specified. Therefore, an enclosure was chosen that would house the PCB, the LCD screen, the LED lights, the auxiliary buttons, and the batteries. The Pactec® enclosures HM series was chosen, as it has a clean form factor, and can easily be milled to house the LCD screen, the LED lights, and any buttons. The enclosure chosen was approximately 3 ¼ inches long by 2 ¼ inches wide and 7/8 inches thick. While this project box was within sight of the original concept drawing, the final product design differed slightly. For instance, the power and reset lights were placed to the left of the screen, and two auxiliary lights were added to the right side of the screen. Finally the power, reset, and auxiliary buttons were placed on either side of the enclosure. The buttons chosen for the final prototype were TL1105B tactile momentary switches by ESwitch®. These switches were chosen because their buttons extend long enough away from the board to be accessible from outside the project box.

3.3.6 Circuit Design

All circuit design was done using EAGLE version 4.16r2. Dominant sections of this schematic are divided into components devoted to power, pressure acquisition, the LCD screen, and Bluetooth®. The board layout was situated based on these individual divisions of the schematic.

Power Scheme

The power schematic chosen followed the BQ24123 Charge Manager's typical applications schematic for 1-cell batteries. Note that this board was designed to attach to a PJ-053D, 3 mm, DC power jack to provide the 5 V supply voltage for the battery to charge off of. Also, instead of attaching 3 separate LED's directly to the power, charge, and done lights, a simple common anode Bivar 3 mm T1 bi-color LED combined the charge and done lights in to one LED. It was wired such that it will light up yellow when charging, and then green when done charging. A simple 3 pin JST connector was attached so either a battery or external power supplies could easily be used to power the board. A kill switch was also added such that the battery could stay attached at all times without requiring the project box to be constantly disassembled while testing. The resulting schematic surrounding the BQ24123 is as shown.

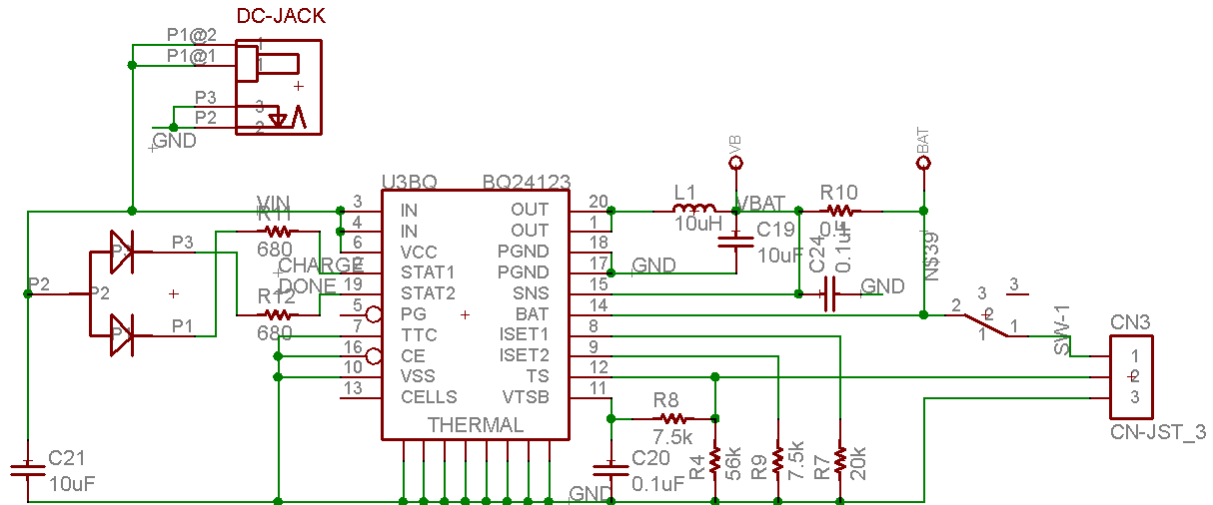


Figure 3-14. EAGLE Schematic surrounding the BQ24123 switch mode charge manager.

While the BQ24123 manages the charging and the battery interface, an on/off mechanism had to be developed to handle when the board actually receives power from the battery. The on/off mechanism as previously explained is based on two constraining factors. Firstly, that the battery voltage is not below the pre-specified 3.4 V safe battery level. Secondly, that the user is not attempting disable the device using the power button. In order to accomplish this, the AND gate was attached to both the STM1061 low-voltage detector, and the output on the MAX16054 on/off controller. A reset button was added to the microcontroller, which acts as a software reset, rather than as an actual power reset. The resulting schematic for the on/off mechanism can be seen below in Figure 3-15.

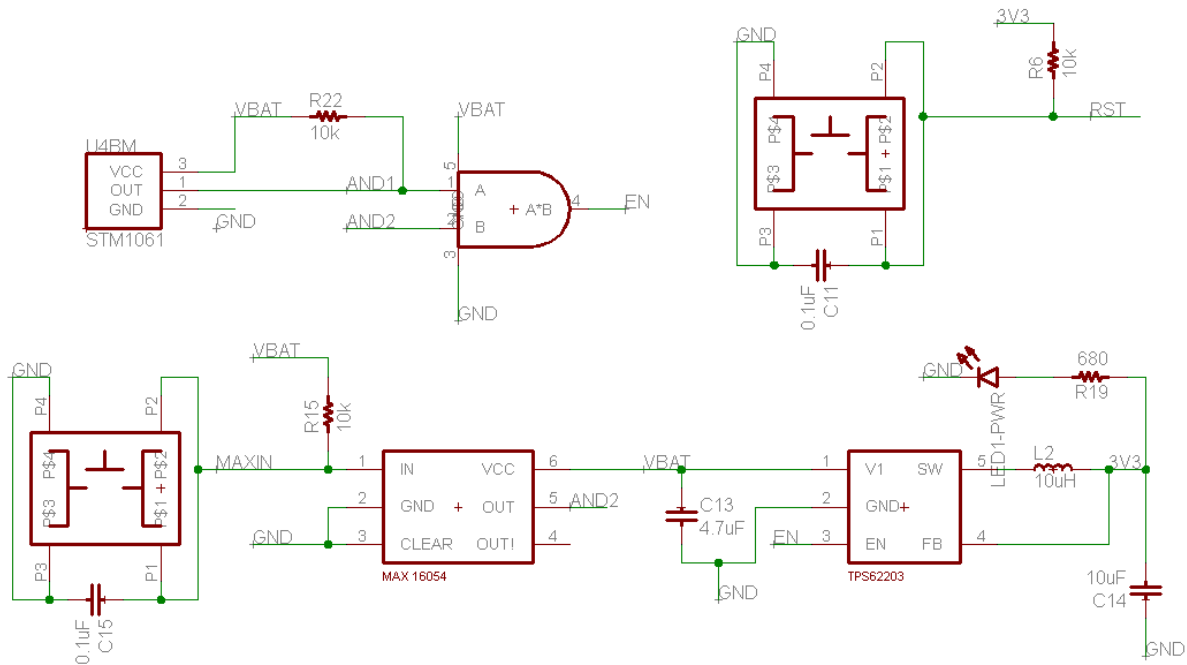


Figure 3-15. EAGLE Schematic containing the MAX16054 on/off controller.

Pressure Acquisition Scheme

Moving beyond the power scheme for the full system prototype, there are several peripherals that actually make the device do what it is meant to do. The first and most important peripheral is the pressure transducer. Surrounding the transducer is the amplifier circuit and the analog-to-digital converter. The schematic for this configuration can be seen here in Figure 3-16.

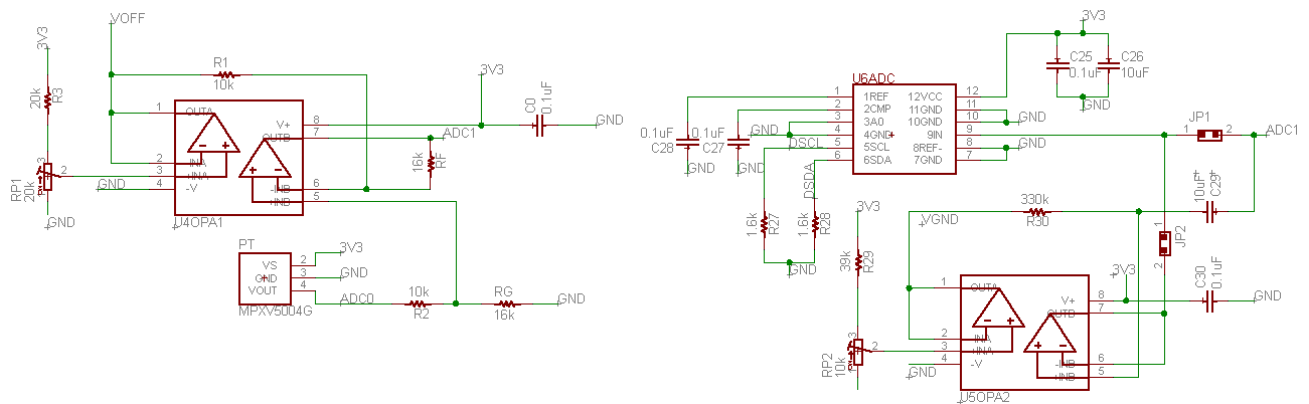


Figure 3-16. EAGLE Schematic containing the amplifier and data acquisition circuit.

There are five main phases within this amplifier circuit. The first is the voltage divider which accounts for the low DC offset voltage of the pressure transducer by taking a voltage rail in series with a resistor, then a potentiometer, and then ground. The voltage divider for the prototype includes a 3.3 V rail, a 20 kΩ regular resistor, and a 20 kΩ potentiometer. The potentiometer allows the voltage offset to

be manually adjusted, anywhere from 0 to 50% of 3.3 V. The normal voltage offset of the pressure transducer is around 0.2 V. However, the divider should be calibrated to whatever that individual pressure transducer's voltage offset is.

The second component in this amplifier circuit is the voltage follower, which is attached to the voltage divider. A voltage follower is implemented using a unity gain amplifier, and does not change the output voltage. The benefit of the voltage follower is that the input impedance of the op-amp is very high, meaning that it draws a low current, and outputs as if it were a pure voltage source. The value of this voltage output is specified by the voltage divider. A single supply op-amp uses a 0.1 uF bypass capacitor on its power supply to stop noise from being coupled back in to the power supply [23].

The third component is the differential amplifier circuit. This differential amplifier circuit uses a negative feedback op-amp configuration, then takes the difference between two input voltages, and finally magnifies this signal by some constant gain, which will be detailed later.

The fourth component decouples the AC component from the signal using a high pass filter. This means that any low signals, such as creep, or the pressure buildup in the tube, are filtered out from the signal. The voltage is then positioned at a virtual ground that is halfway between the ADC's maximum output voltage of 1.25 V. This will provide both positive and negative voltage responses based on an increase or decrease in pressure in the tubing.

The fifth and final component is the Buffer. This simple voltage follower circuit allows the ADC to receive a pure voltage source reading, with very low impedance. Figure 3-17 below shows a simplified schematic, highlighting all five components of these amplifier circuit components.

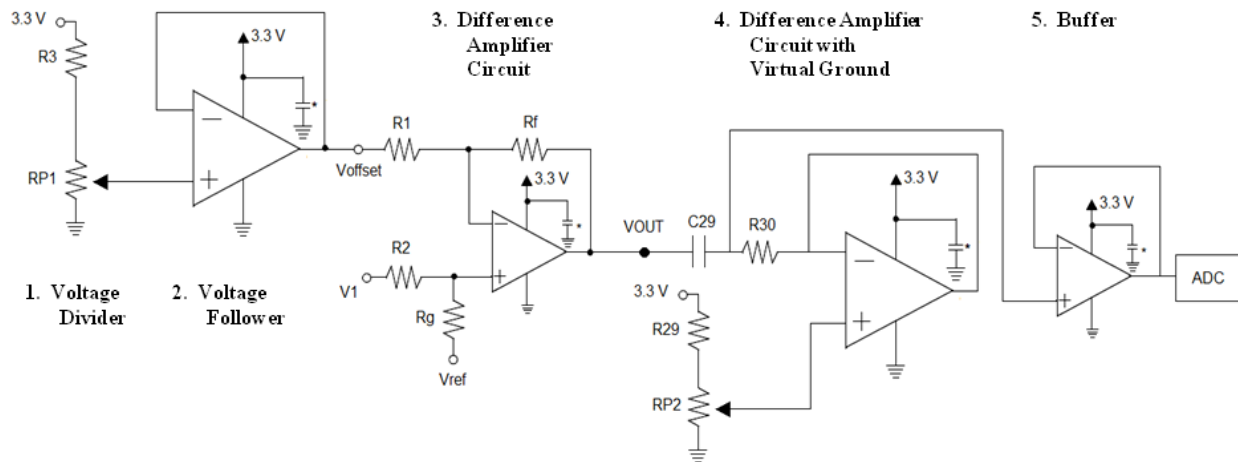


Figure 3-17. Prototype amplifier and DC decoupling circuit.

The governing equations for such a circuit are provided below. Note that Voffset is the voltage after the voltage follower which is governed by the voltage divider. Meanwhile, Vout is what voltage flows out at the differential amplifier circuit. Also note that Vref in this circuit is connected to ground.

$$V_{offset} = \left(\frac{R_{p1}(\% \text{ of potentiometer})}{R_3 + R_{p1}} \right) \quad \text{Equation 3-25}$$

$$V_{out} = (V_1 - V_{offset}) \left(\frac{R_f}{R_1} \right) + V_{ref} \left(\frac{R_g}{R_2} \right) \quad \text{Equation 3-26}$$

In addition to these equations are the equations for the high pass filter and virtual ground. The high pass filter filters out all frequencies below the cutoff frequency f_c . Meanwhile the virtual ground is determined by the same equation for the voltage divider. The resulting equations are shown below.

$$f_c = \frac{1}{2\pi R_{30} C_{29}} \quad \text{Equation 3-27}$$

$$V_{virtual_ground} = \left(\frac{R_{p2}(\% \text{ of potentiometer})}{R_{29} + R_{p2}} \right) \quad \text{Equation 3-28}$$

In terms of actual components, a Texas Instruments OPA2301 was chosen to read the output from the pressure transducer, because it had 16-bit resolution which matched the voltage resolution required for the lowest amplitude signal. In addition it had low noise, a low single supply power rating, and a compact form factor containing two amplifiers in one component. The linear technologies LTC2471 ADC was chosen also due to its 16-bit resolution, as previously described. The incoming pressure signals are converted and read in through the I2C interface on the microcontroller. Both the op-amp and the ADC were powered by the 3.3 V singly supply rail. The gain on the differential amplifier circuit was chosen to be 1.6, to keep the DC component of the signal less than the ADC's 1.25 V rail. This value may be lowered, if the system begins to have issues with oversaturation of the ADC or op-amp.

LCD Screen Scheme

With the data being gathered at the correct resolution, and with the voltage offset of the pressure transducer accounted for, the software can then receive a high resolution signal from the pressure transducer. Depending on available resources, the heart and respiratory rates can then display them to the NHD-C0216 LCD screen. Similar to the ADC, the LCD screen was written to using the I2C interface on the microcontroller. The schematic shown below in Figure 3-18, follows the LCD screen's datasheet for normal operation.

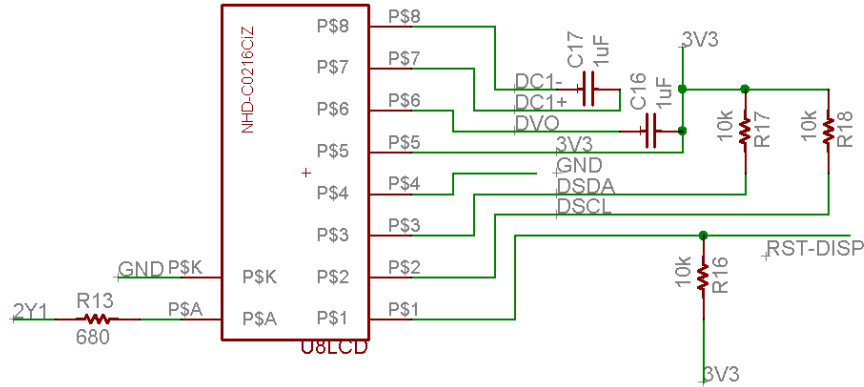


Figure 3-18. EAGLE Schematic showing the two JST connectors that are used for the LCD screen.

Both the screen itself and its LED backlight can take in a 3.3 V single supply voltage. An active low reset pin is connected to one of the general purpose pins on the microcontroller to provide a programmatic LCD screen reset option to the user. In addition, the LED backlight was connected to the LED driver such that it could also be controlled programmatically.

Bluetooth® Scheme

While the board is capable of taking data measurements and displaying estimated heart and respiratory rates to an LCD screen, having a wireless interface would allow the board to communicate with external devices, and transfer data seamlessly. For testing purposes, an external receiving device such as a laptop could receive data from the device. For prototyping purposes, a smartphone or tablet could be used to show the device's ability to communicate multiple sets of information on to a single screen. The end goal of course is for a simple touch-screen interface that EMT's could use to observe multiple casualties in the field. This will be explored in section 5.2 Future Work.

The module chosen to provide this wireless capability was a Bluetooth® V2.1, class-2 power module, 3.3 V single supply voltage, with serial communication using an AT command set. It was especially practical due to its small 10.5 x 13.5 mm module size. The only components that need to be connected to this module are an external antenna connected through a capacitor. It also requires connection to the I2C interface on the microcontroller, for programming purposes. The external antenna chosen was a Rufa 2.4 GHz Bluetooth® antenna. It is an antenna that is commonly used for mobile phones, headsets, and laptops.

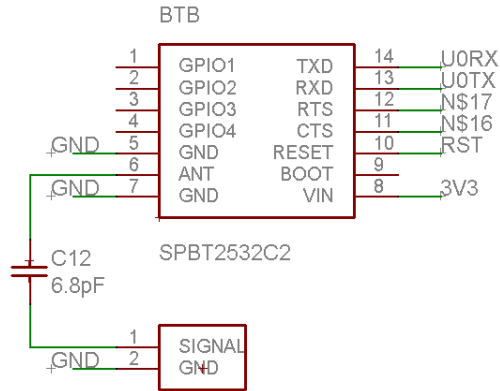


Figure 3-19. EAGLE Schematic showing Bluetooth® module, and the Antenna.

Board Layout

Prior to board layout, every component of the schematic was verified according to each components respective datasheets. Next, each component was placed on the PCB layout, which was sized according to the project box’s specifications to be 89.5 mm long by 54.4 mm wide. First the screw terminals were placed on the board prior to layout, so as not to route traces over top of them. Next, the microcontroller was placed in the center to allow for easier routing of all other peripherals. The JTAG connector was placed in the lower left corner, for easy debugging and programming using LabVIEW. The power scheme was placed on the left hand side, along with the PJ-053D DC power jack, the kill switch, and the 3-pin JST battery port. The LCD, Bluetooth®, and pressure acquisition schemes were all placed on the right hand side of the board. Note the pressure transducer was given room to connect to tubing underneath the board. Finally the LED’s for power and charging were placed on the top left of the board, next to their corresponding buttons. The opposite side contains two other LED’s and two auxiliary push buttons which can all be controlled programmatically. The resulting PCB layout, in Figure 3-20 below, includes dimensions and areas of interest.

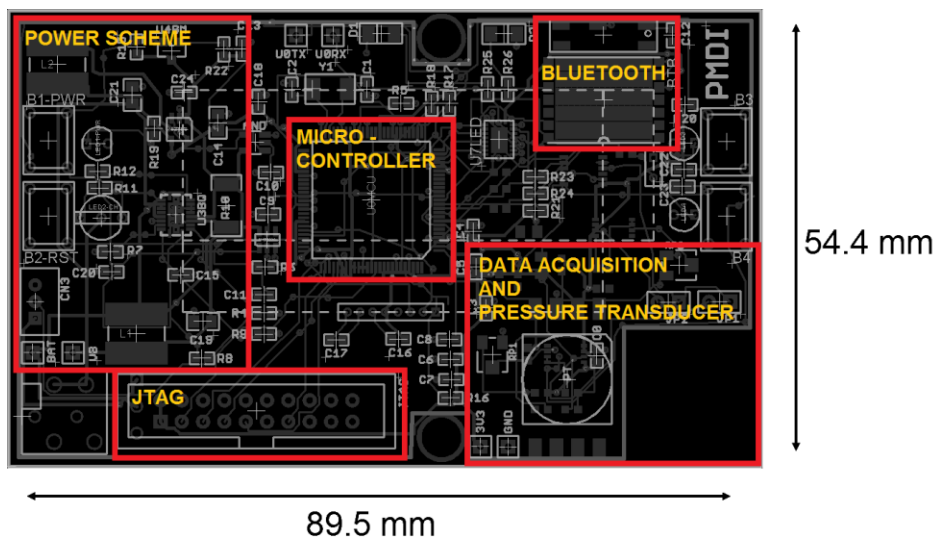


Figure 3-20. Prototype PCB board layout with dimensions.

The benefit of this layout is that the project box becomes a familiar and ergonomic interface for a user to be able to rapidly assimilate to. The power scheme and auxiliary buttons are something that anyone who is familiar with smart phones and small electronic devices, should be able to navigate.

The board was verified using “freeDFM,” a web service provided by Advanced Circuits. Once all design checks were verified, the board was sent to Advanced Circuits, and all components of the board were either ordered or sampled. The PCB was then populated in the Mechatronics Lab at Virginia Tech. After which, the hardware was tested using embedded software designed in LabVIEW. The final board in its project box can be seen in Figure 3-21 below.



**Figure 3-21. Prototype PCB after being populated.
Photo by author, 2012.**

3.3.7 Bladder Configuration

The bladder used for the full system prototype involved taking a 6 inch length of Tygon® super-soft latex rubber tubing and attaching it to the pressure transducer. A hole was drilled in to the bottom of the project box, such that the tubing could be routed directly underneath the board. Next the tubing was placed in a single spiral form factor, to create a maximum change in pressure for such a small length of tubing. A small piece of malleable metal ensures that it stays in a relatively uniform configuration during testing. An image of the resulting form factor for the bladder can be seen below in Figure 3-22.

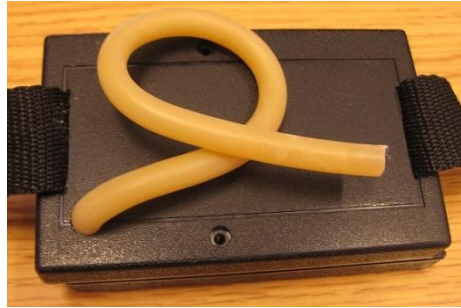


Figure 3-22. Prototype bladder configuration used for human testing.
Photo by author, 2012.

This change in tubing configuration from a pure spiral to a single crossed spiral should result in a different voltage response. It is anticipated that two layers of tubing will create a higher voltage response than a single layer of tubing, given the same input. The tubing or bladder configuration is considered a highly variable factor in the design, and future work concerning it will be discussed in section 5.2.2 Manufacturing a Custom Bladder.

3.3.8 Software Development

Once the full system prototype was fully assembled and ready to go, each component had to be programmed in LabVIEW. The three main components of the full system prototype that required programming were the NHD-C0216 LCD screen, the LTC2471 ADC, and the SPBT2532C2 Bluetooth® module. The LCD screen and the ADC both were programmable through the I2C interface. LabVIEW's ARM I2C palettes provide subVI's with the ability to configure, open, read, write, and close I2C ports. Therefore the only thing necessary to get the two to operate, was to provide LabVIEW the slave address and any configuration messages that might be required to operate each of the two respective devices. The LCD screen's slave address was x7C, while the ADC's slave address was x14. Due to a known configuration issue in the LCD screen by Newhaven, a bit shifting occurs when reading in the slave address. Therefore, the LCD screen chosen can only be recognized when bit shifted to the right or by dividing its address in decimal form by two. This means the LCD screen's new slave address becomes x3E. The rest of the configuration VI with some sample code can be seen throughout Appendix C: LabVIEW Code.

In order to prepare the device for gathering data, the Bluetooth module was configured to transmit samples from the ADC every second. The sampling frequency of the ADC needs to be twice that of the highest expected frequency, in order for it to be seen using signal processing techniques such as an autospectrum. Heart rate is therefore the confounding factor, because it occurs at a higher frequency than respiration, in most cases. The upper limit of heart rate, as previously stated, varies with age. However, one can safely assume that 200 bpm is well above the upper limit for any person. 200 bpm corresponds with a 3.333 Hz heart rate signal. The sampling frequency therefore must be at least 6.666 Hz. Due to the available hardware resources, a 20 Hz sampling frequency was chosen, such that there would be three times the required sampling frequency to extract the desired signals. The reason why a higher sampling frequency was not chosen has more to do with the available memory on the

board. Due to the fact that an array of data must be stored and sent over Bluetooth every second, the device is limited during data collection to a certain amount of memory. In fact the Bluetooth and ADC are both capable of operating at much higher sample rates. This will be addressed further in Chapter 5: Conclusions and Future Work.

With a 20 Hz sampling frequency, the Bluetooth would be sending 20 samples every second to the computer. The Bluetooth receiver on the computer would then concatenate an array of recent data, creating a larger array including all the received data. This data would be recorded using 6 significant figures and 5 decimal precision. This precision was chosen based on the 0.01907 mV resolution of the ADC, and the fact that the voltage response can never exceed the 1.25 V saturation limit of the ADC. The data recorded on the computer could then be manipulated using different signal processing such as a correlation function analysis. This data could also be used to correlate with other data, such as ECG analog voltage output, in order to benchmark its results based on industry standard devices.

3.3.9 *Institutional Review Board Human Testing*

The human test was devised on paper and passed through Virginia Tech's institutional review board. The test documents approved by the IRB can be seen in Appendix D: IRB Forms. The procedures as explained to test subjects in the consent form are as follows. Subjects are told that they are being asked to allow the researcher to place the heart and respiratory monitor on their chest and measure their heart and respiratory rates. In addition they may be asked to wear a multi-lead electrocardiography (ECG) tool, or pulse oximeter, and allow the researcher to take manual pulse estimation by holding your wrist. The ECG tool used for testing was an MDE Escort Prism Vital Signs Monitor. It consisted of a three lead setup, with two leads placed on either side of the chest and one placed on the stomach. The positions of the ECG leads can be seen in Figure 3-23 below.



**Figure 3-23. ECG lead locations during human testing.
Photo by author, 2012.**

The raw data from the ECG is transferred to a computer using the same NI-USB-6009 device that was used for estimation, meanwhile the pressure signal is transferred to a computer over Bluetooth. The two data sets were then investigated using MATLAB.

Subjects are told the test will take approximately 15 minutes, and that it will all take place at the Mechatronics Lab in Randolph Hall at Virginia Tech or wherever convenient for the subject. Subjects

may be asked to wear specific upper body clothing made of different materials such as body armor. The researcher would of course provide such clothing. All test results shown in this report are taken with the subject lying in a supine position, to simulate a mass casualty scenario. In all cases, the subject was also wearing no more than two layers of clothing, such as a t-shirt or undergarment. An image of the test setup can be seen below in Figure 3-24.

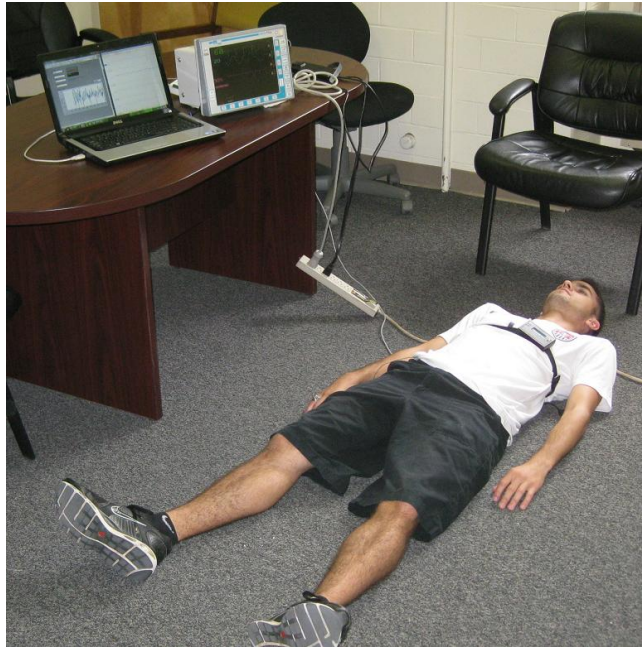


Figure 3-24. Human testing setup.
Photo by author, 2012.

It was determined that a 90 second block of data from both the device and the ECG would be the starting point. From there, a 60 second block of data would be extracted. The number of averages and resulting frequency resolution during signal processing will be explained further in the results.

3.3.10 Analysis and Signal Processing

One potential signal processing technique for data analysis would be a cross correlation. This would provide a comparison of the similarities between the two signals by a time lag applied to one of the signals. However, this would be most useful if the testing was concerned with synchronizing the device with an industry standard ECG machine. However, this is not the focus of the testing. Rather the testing is intended to compare the likeness of the two signals, in order to discover whether or not the embedded signal in one set of data is also what caused the embedded signal in the other set of data. To truly test this causality between the embedded signals in the device with those in an ECG, a coherence function analysis would be ideal. Coherence is a measure of the power transfer between two signals. In other words, it measures the causality between one event and another event. A high coherence would mean that one event is understood as a consequence of the other. In order to accomplish this, the coherence estimate utilizes a cross spectrum and the two autospectrum results. Such results and

analysis were vital in measuring the capability of this device versus the industry standard. However, survey results were vital to gaining input on the need and the future applications of the device.

3.3.11 *Institutional Review Board Explorative Survey*

In order to receive direct feedback from emergency medical responders, a survey was created, and passed through Virginia Tech's institutional review board. The survey, which can be seen in Appendix D: IRB Forms, was provided to emergency medical responders throughout Roanoke and Blacksburg, using contacts from researchers at Carilion Clinic, Carilion Roanoke Memorial Hospital, and the Virginia Tech Rescue squad.

As seen in the survey, there were very important questions that needed to be answered. These answers would help in development of the full system prototype, and in future applications of the project. Firstly, it was important to know by what means the average EMT takes heart and respiratory rate information from a patient. Secondly, how much time does it take them to do this, and do they ever have difficulty finding the information based on their current method? Finally, would they find it useful to have a deployable chest sensor such as the one that we are creating? The survey was distributed throughout the spring and summer of 2012. The data was then collected, analyzed, and is described in section 4.2.2 IRB Explorative Survey. Regardless of the results, it was expected that future applications of the technology will take in to account the comments found in the survey results.

3.4 Signal Processing Techniques

Through digital signal processing it is possible to extract information from a discrete time domain signal that is not always obvious to an observer. In order to observe both cardio and respiratory signals simultaneously, an understanding of several signal processing techniques is required. Several techniques have already been explored throughout the methods section. However, this section is meant to explain those techniques in further detail, as they will be utilized throughout the results section. These techniques include but are not limited to Fourier series, cross correlation, cross spectrum, autocorrelation, autospectrum, coherence analysis, and averaging.

3.4.1 *Fourier Series Analysis*

Fourier series analysis is a method by which time domain waveforms are decomposed in to magnitude and phase estimates in the frequency domain [24]. These decompositions are made up of sine and cosine functions, at a variety of frequencies. It is a widely utilized technique throughout signal processing. The discrete Fourier transform or DFT provides this decomposition. It is used to calculate the cross spectrum and the autospectrum. It is assumed throughout this discussion that there are two arrays of data, x and y .

3.4.2 *Cross Correlation and Cross Spectrum*

Cross correlation (C_{xy}) is a measure of the likeness of x and y as a function of time lag applied to one of them. Meanwhile the cross spectrum (G_{xy}) is the Fourier transform of this cross correlation.

Note that the conjugate of G_{xy} is equal to G_{yx} . It is always a complex valued function, and does contain phase information between two signals. The easiest way to find the cross spectrum, programmatically, is to take the DFT of sample record of y , and then multiply this by the complex conjugate of the DFT the same size sample record of x [24]. The result is the double-sided spectrum, because the amplitude of the cross spectrum is half of the desired result, or the single-sided spectrum.

3.4.3 *Autocorrelation and Autospectrum*

Autocorrelation (C_{xx} or C_{yy}) is the cross correlation between a signal and itself. This can mean the cross correlation between x and itself, or between y and itself. Autospectrum (G_{xx} or G_{yy}) is the Fourier transform of the autocorrelation. It is always a positive function, and it contains no phase information. One way to find the autospectrum, of x for example, is to take the DFT of a sample record of x , and then multiply this by the complex conjugate of the same sample record. The outcome is again the double-sided spectrum, because the amplitude of the autospectrum is half of the desired result, or the single-sided spectrum. Once G_{xy} , G_{xx} , and G_{yy} have been found one can use these to measure the causality between x and y . Such causality can be found by relating these spectral estimations in terms of correlated and uncorrelated content.

3.4.4 *Coherence Function*

Coherence is a measure of the causality between two events as a function of frequency. This will only be useful in showing what level of confidence exists that an event in x causes an event in y . The coherence function has a value between 0 and 1, where 0 suggests that the output measurement is composed entirely of uncorrelated content. Meanwhile a value of 1 suggests that there is no uncorrelated content in the output measurement.

3.4.5 *Averaging*

In order to gain true confidence in the results, several sample records of data must be evaluated from the original data. This is where the term averaging comes in to play. Averaging provides confidence behind the results of G_{xy} , G_{xx} , and G_{yy} , by creating an ensemble of sample records. The individual auto and cross spectrum results of each sample record are then averaged. The more averages, the more confidence one can have in the results. The number of averages that can be taken is limited by the original size of the data. If there is only one sample record, then there can be no confidence in the autospectrum or cross spectrum results. Meanwhile as the number of averages approaches infinity, the more confidence one can have in the result.

3.4.6 *Frequency Response Function*

When defining inputs and outputs, the term Frequency Response Function (FRF) is inseparable. The FRF can be measured from output to input or from input to output. In this case, the calculations are based on an H1 estimate, where the FRF is equal to G_{xy} divided by G_{xx} . If the calculations were based on an H2 estimate, the FRF would be equal to G_{yy} divided by G_{yx} [24]. Recall that the conjugate G_{xy} is equal to G_{yx} . Note that the coherence estimate would be the same, regardless of an H1 or an H2 estimate.

These techniques will prove useful, as explained throughout the following sections on simulated testing, and human testing. In the section on simulation, the autospectrum will be useful in order to understand the characteristics of the embedded signal. It will also be useful for determining the signal to noise ratio, which will be discussed. For the human testing results, the cross spectrum and coherence function will be useful in order to understand the causality between the signals embedded in the pressure transducer device output, with the signals embedded in the ECG machine's output. The number of averages used to estimate G_{xy} , G_{xx} , and G_{yy} will be explained according to each section.

Chapter 4: Results and Discussion

Now that all of the methods for simulated testing and human testing have been explained in detail, the results of each stage will be provided. The simulation results will compare the data results from the first PCB to the simulated breathing mannequin. Finally, the human testing results will showcase the coherence between the data from the final pressure transducer device with an industry standard ECG. In addition, some of the general trends that resulted from the IRB survey will be presented and explained. Note that the y-axis in all of the plots, as well as the rest of the time domain plots shown, is in terms of voltage (V) and not gauge pressure (kPa). The conversion between the two is based on the sensitivity of the pressure transducer being used, along with the offset, and the gains of the amplification circuit. However, such a conversion is not relevant to purpose of these tests.

4.1 Simulation Results

The simulation device was taken to Carilion Center for Experiential Learning and attached to a simulated breathing mannequin. Data was collected at a 1000 Hz sampling frequency for a minimum of 30 seconds. This sampling rate far exceeded what was needed in order to show the frequencies that were expected due to heart and respiratory rate. The highest frequency that was expected during simulation would be 1.0 Hz. Therefore the minimum sampling frequency required, would only have to be twice this, or 2.0 Hz. By decimating the 30,000 point array down in to 10 sets of 3,000 point arrays, the frequency range of the autospectrum results would now have a sampling frequency of 100 Hz, and therefore a Nyquist of only 50 Hz. However, these sample records were created differently for simulation results, than for any other results. In this case, the original array was decimated. The benefit of doing it this way is the user still has 30 second blocks of data to use for averaging. This method is therefore much more effective in this case, as long as no aliasing appears due to higher frequency content in the results. A plot below shows the autospectrum results for the 24 RR test, based on the original data before decimation.

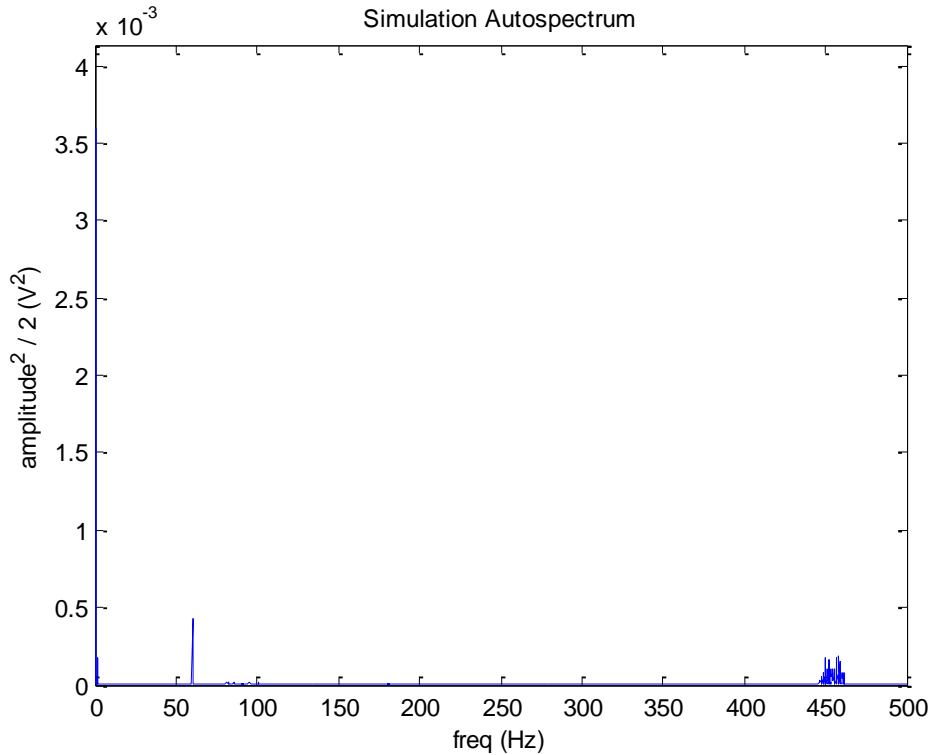


Figure 4-1. Simulation device autospectrum results at 24 RR shown up to the 500 Hz Nyquist.

As seen in this figure, there are two areas of higher frequency content at both 60 and 450 Hz. The 60 Hz noise is expected due to electrical interference, while the 450 Hz noise is may be related to something else in the testing environment. Regardless, their results are very low in magnitude and would likely not cause aliasing in the results due after decimation. Therefore, the rest of testing and the results after decimation will now be explained.

As described, tests were done at six different respiratory rates (RR's). These rates were 12, 18, 24, 30, 40, and 60 RR. The expected frequencies from these RR values were 0.20, 0.30, 0.40, 0.50, 0.67, and 1.0 Hz respectively. Each data set was imported in to MATLAB, where an autospectrum was calculated using averages based on the decimation described above. This data, which was originally a 30,000 point array, was broken up in to 10 averages, made up of 3,000 point arrays. By observing the spikes in the autospectrum, the respiratory rates were compared to the expected respiratory rates. The device was capable of detecting the actual respiratory rate in all six cases. The figures below show one of these simulated breathing rates, at 24 RR, and its corresponding pressure response in the time domain and its autospectrum in the frequency domain. Note that the other five data results not shown below can be found in Results, under Appendix E: Results, Simulation Testing Results.

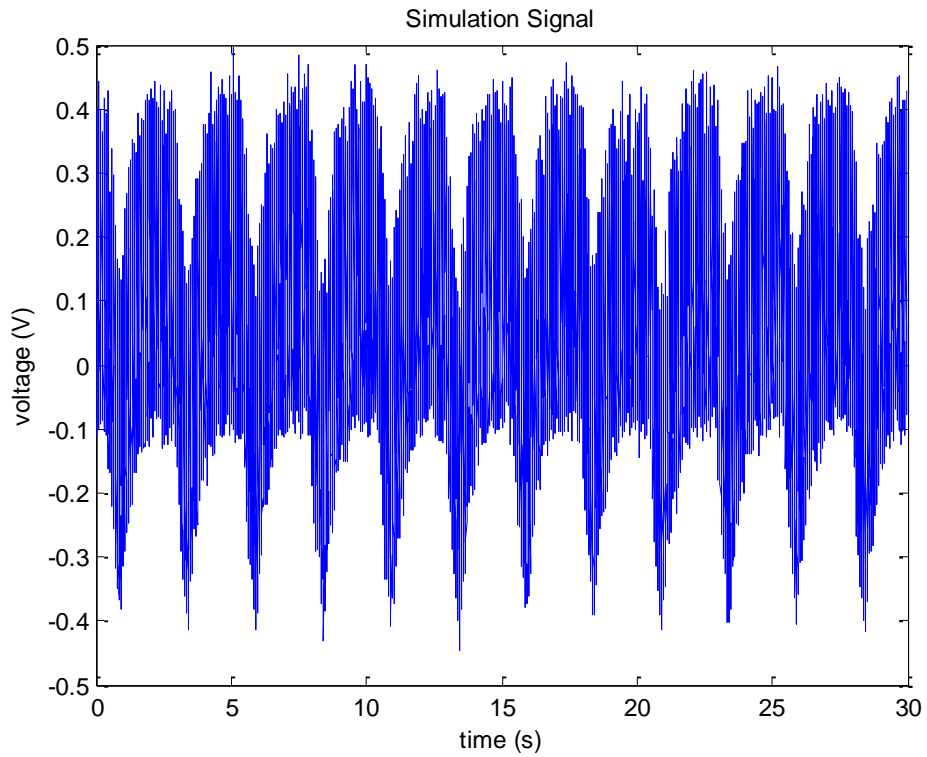


Figure 4-2. Simulation device output as a function of time for the 24 RR test.

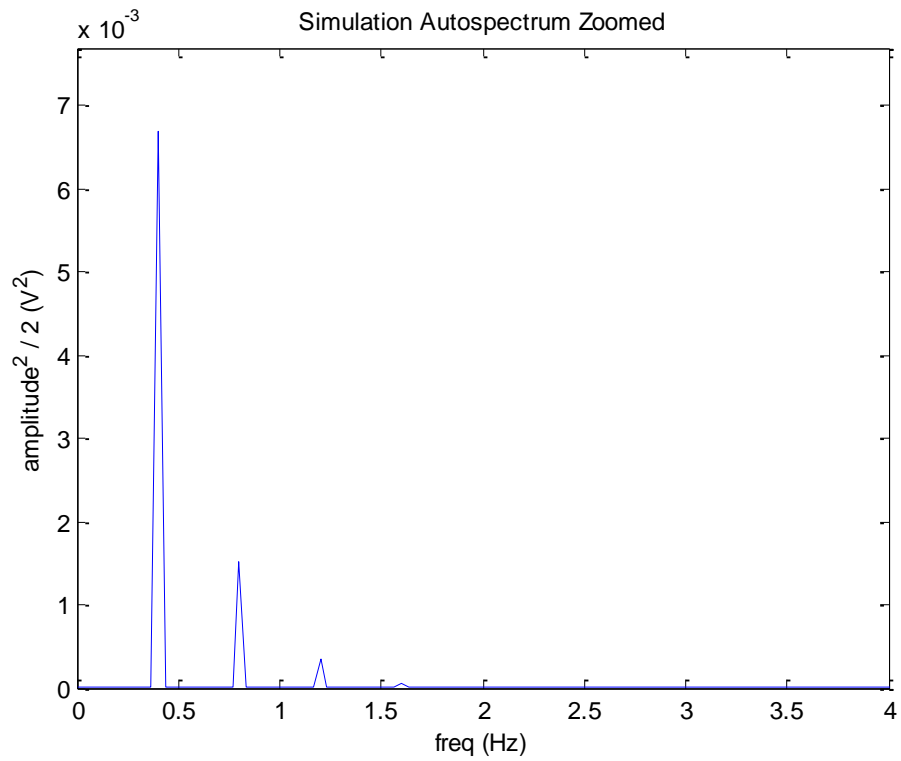


Figure 4-3. Simulation device autospectrum results at 24 RR.

It should be noted that the multiple peaks after the first observed frequency in each of the autospectrum results, are harmonics of the original signal. These peaks are expected in any Fourier analysis, as they represent whole number multiples, or harmonics, of the original frequency. The SNR for respiration improved from 4.014 during feasibility testing, to 48.915 for the simulation tests.

The increased SNR is a positive sign. It can be explained in two ways. The first explanation is that the feasibility test was done on a human. Humans have more variation in their breathing and other extraneous movements than the simulated breathing mannequin. The second explanation, as expected, is that the move from through-hole breadboard components, to solder mounted PCB components, reduced the transfer of noise. Besides improved SNR, the device must be capable of accurate estimation of the respiratory rates of the mannequin. The observed frequencies in each of the six tests are tallied in the table below. The plots for each of these tests can be found in Results, under Simulation Testing Results.

Table 6. Simulated versus observed frequencies during simulation testing.

Test #	Respiratory Rate (per minute)	Expected Freq. (Hz)	Observed Freq. Spike (Hz)
1	12	0.20	0.20
2	18	0.30	0.30
3	24	0.40	0.40
4	30	0.50	0.50
5	40	0.6667	0.6667
6	60	1.0	1.0

As shown the observed frequencies matched exactly to each of the six expected frequencies. Any coherence analysis would require another signal to compare to. The hypothesis being that the embedded signals in each are caused by each other. As mentioned previously, human testing was carried out that would verify whether or not these signals were caused by each other.

4.2 Human Testing Results

For the purposes of benchmarking this device versus the industry standard for heart and respiratory monitoring, a set of IRB approved human tests were carried out. Specifically, several subjects were attached to ECG leads, which would monitor their heart and respiratory rates. Simultaneously, they were attached to the full system prototype. Both the ECG and the full system prototype transmitted data to the computer for 90 seconds at a 20 Hz sampling rate. In addition to the benchmark testing, an explorative survey was distributed to EMT's. The results of both the benchmark testing and the explorative survey are shown below.

4.2.1 IRB Benchmark Testing

The raw data was taken in to MATLAB, where the following results were found. Firstly, the results varied from person to person. Some subjects had heart and respiratory rates that could easily be seen by both the pressure transducer device and the ECG. Meanwhile, other subjects' heart rates were

not as obvious in the data results. The test results with the most and least coherence between the prototype device and the ECG machine, for both heart and respiratory frequencies, will be detailed below.

Subject 4 showed perhaps the highest heart and respiratory rate coherence between the prototype and the ECG. As seen below, they exhibited a clear pattern of both heart and respiratory rate in the time domain signal for the pressure transducer device. The autospectrum was calculated using averaging. This data, which was originally a 1200 point array, was broken up in to 6 averages, made up of 200 point arrays. The time domain and autospectrum results from the prototype device can be seen below in Figure 4-4.

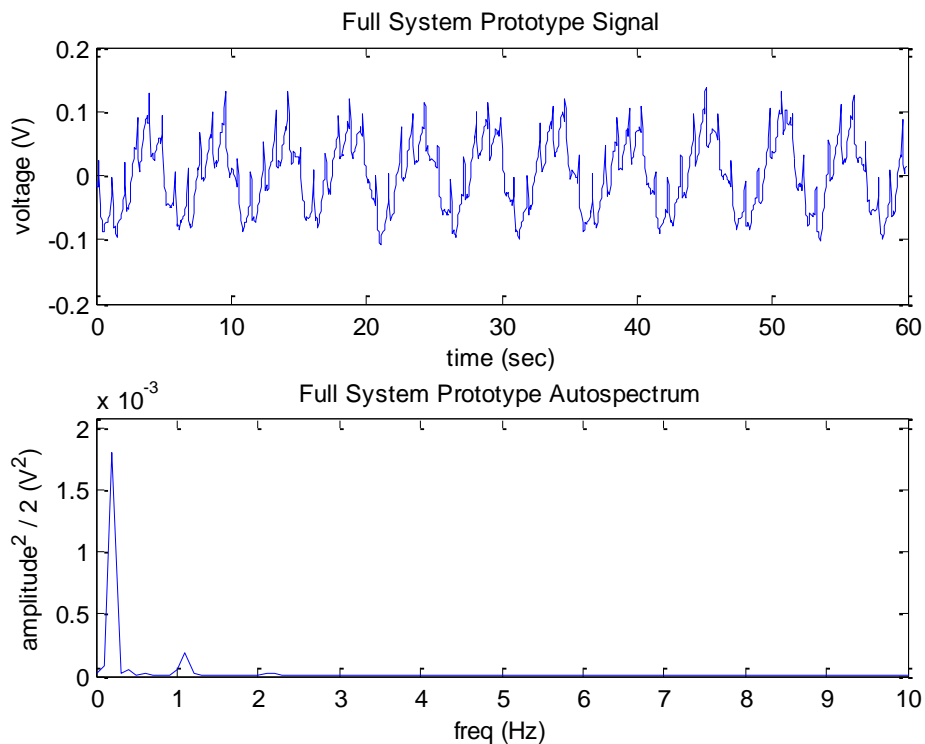


Figure 4-4. Subject 4 time domain and autospectrum results for the Full System Prototype.

The two notable frequencies in the autospectrum are at 0.2 and 1.1 Hz. The SNR for respiration for subject 4 was 622.3, while the SNR for heart rate was 201.9. Note that these SNR values were capable of being calculated subject 4, as their heart and respiratory rates could both be represented in the autospectrum of the prototype. Due to the fact that some subjects did not have an observed frequency matching that of heart rate, the SNR for heart rate could not be calculated. However, the SNR for respiration for all subjects, varied between 600 and 2000.

Meanwhile, the ECG results for subject 4 show similar results. This is evident in the autospectrum, which shows notable observed frequencies at 0.2 and 1.1 Hz. The two frequencies don't

mean anything, until they have been compared to the ECG results. The time domain and autospectrum results from the ECG can be seen below in Figure 4-5.

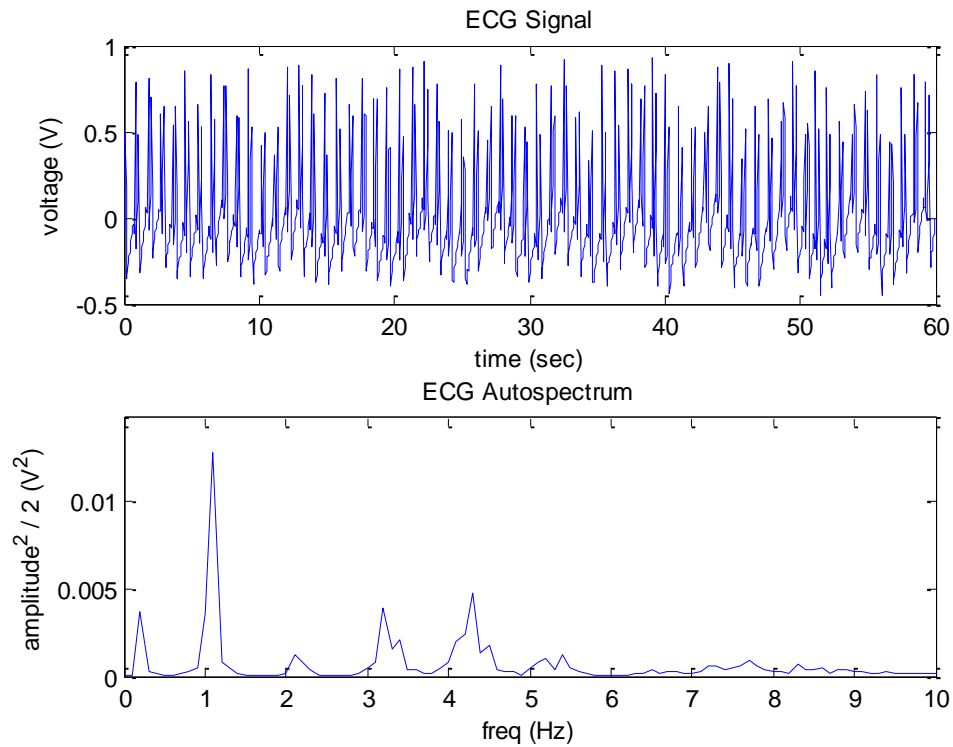


Figure 4-5. Subject 4 time domain and autospectrum results for the ECG.

The similarity between the two autospectrum results suggests that the two devices were picking up the same signals. However, the only way to prove the causality between the prototype and the ECG signals is through a coherence function analysis.

Coherence as previously stated, is represented by values 0 through 1. A coherence of 0 suggests that the output measurement is composed entirely of uncorrelated content. Meanwhile a coherence of 1 suggests that there is no uncorrelated content in the output measurement. The function is calculated by taking the product of G_{xy} and G_{yx} and then dividing by the product of G_{xx} and G_{yy} . The coherence plot for subject 4's pressure transducer and ECG results is seen below in Figure 4-6.

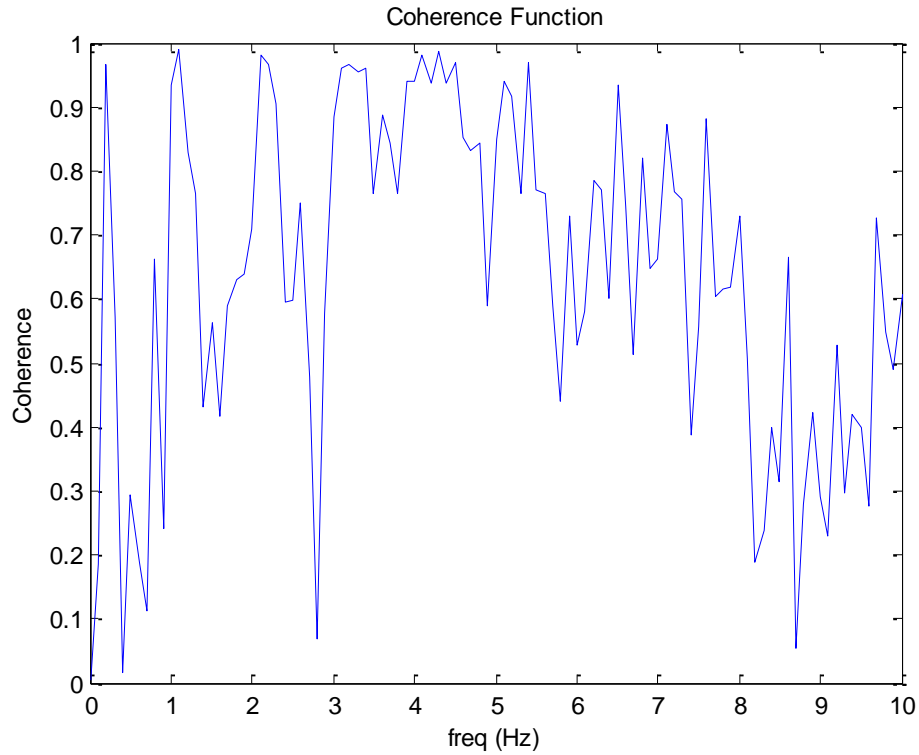


Figure 4-6. Subject 4 coherence function results between the Full System Prototype and the ECG.

The coherence plot shows a coherence of 0.9671 at 0.2 Hz. In addition, there is a coherence of 0.9897 at 1.1 Hz. This is interpreted to say that the signals had high coherence at both of the frequency ranges for heart and respiratory rate. In this case, high coherence is defined as any coherence over 0.9.

Subject 2's results varied dramatically from subject 4. Exactly as before, a 60 second portion of data using ten averages was used to create the autospectrum and cross spectrum. However, this time the pressure transducer resulted in only one observed frequency at 0.2 Hz. The time domain and autospectrum results for the prototype can be seen below in Figure 4-7.

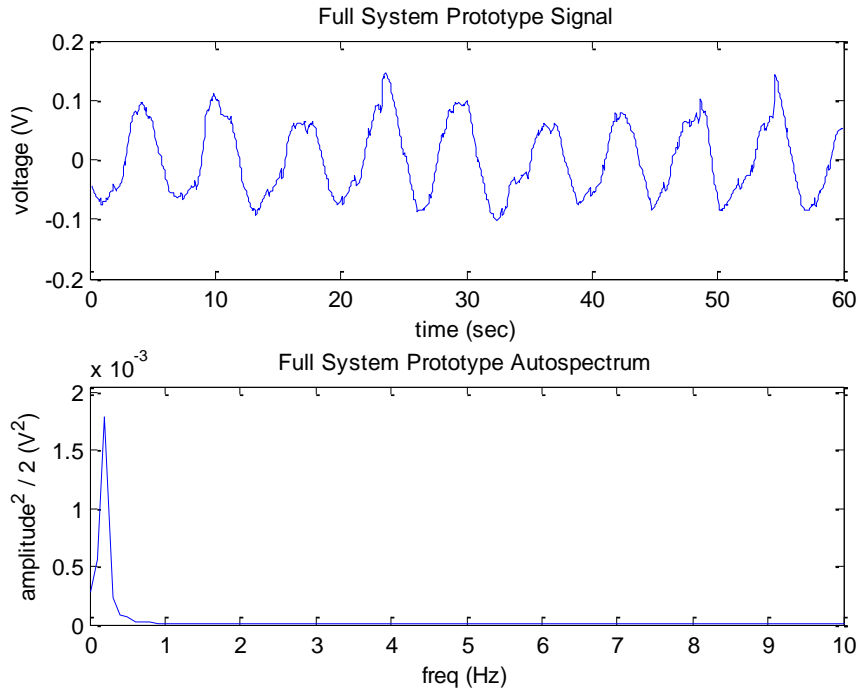


Figure 4-7. Subject 2 time domain and autospectrum results for the Full System Prototype.

The ECG results varied from the prototype. The autospectrum results for the ECG show two frequencies at 0.2 and 1 Hz. The time and frequency domain results from the ECG can be seen below in Figure 4-8.

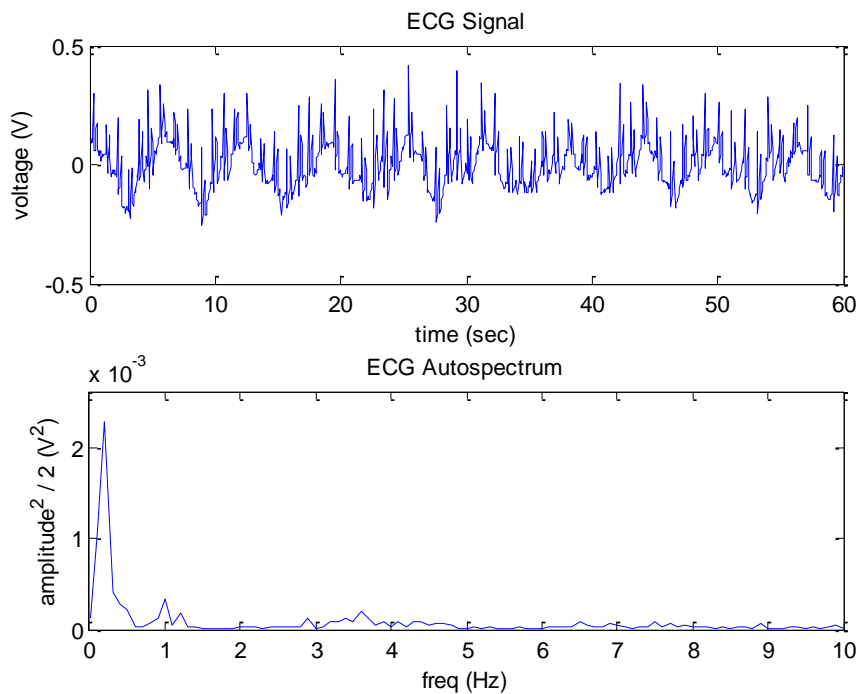


Figure 4-8. Subject 2 time domain and autospectrum results for the ECG.

The autospectrum results suggest that the two devices were picking up the same respiratory rate, however the heart rate was not clear. As with subject 4, the coherence plot will show the true test of the relationship between the prototype output and the ECG output. The coherence plot for subject 2's pressure transducer and ECG results is seen below in Figure 4-9.

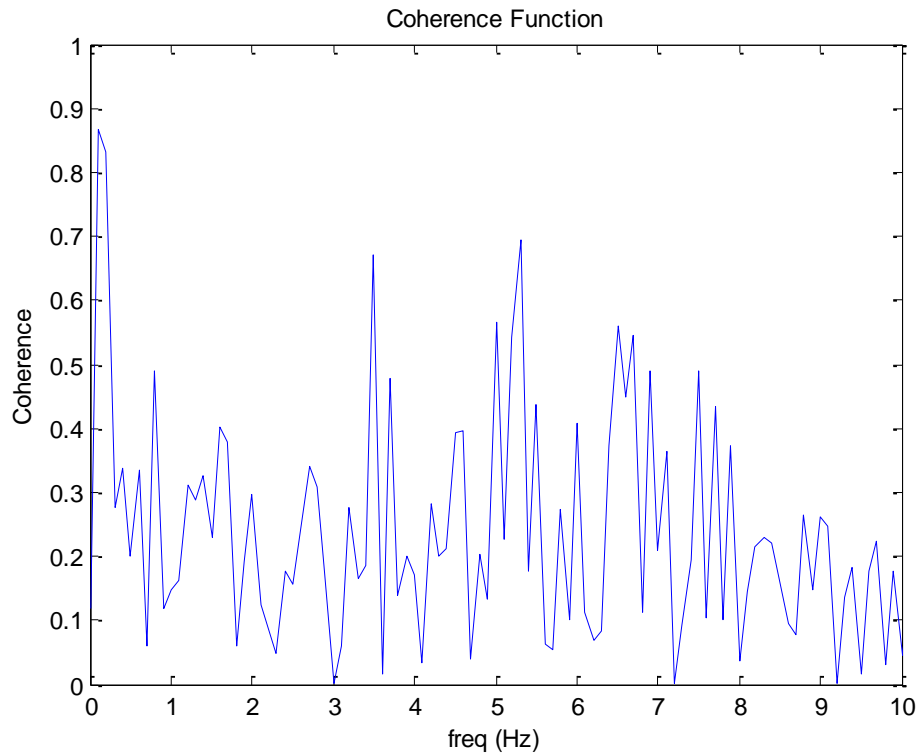


Figure 4-9. Subject 2 coherence function results between the Full System Prototype and the ECG.

The coherence plot shows a coherence of 0.8662 at 0.1 Hz. However, beyond that there were no areas of coherence that even approached the 0.9 limit. At the 1 Hz, where one would expect there to be high coherence between the prototype and the ECG, the coherence was 0.1462. This means that this subject's results had a high coherence at the frequency range matching that of the respiratory rate. However, there was very low coherence at the frequency matching that of the heart rate.

The rest of the data results varied between those of subjects 4 and 2. Every set of data results showed that respiratory signal could be detected using the pressure transducer device. Table 7 shows the coherence result at the expected respiratory frequencies for each subject.

Table 7. HARMONI vs. ECG observed respiratory frequencies and coherence.

Subject:	Respiratory Frequency (Hz):	RR:	Coherence:
1	0.3	18	0.4003
2	0.2	12	0.8662
3	0.2	12	0.8727
4	0.2	12	0.9671
5	0.1	6	0.2230
6	0.1	6	0.9163
7	0.1	6	0.5033

As seen in this table, 4 out of the 7 subjects had had a relatively high correlation value, above 0.8, at the respiratory frequency. These results support a relatively high correlation for respiratory rate between the pressure transducer device and the ECG for 4 out of the 7 subjects. The reason why more subjects did not show a high coherence at the respiratory frequency is because the raw ECG results do not necessarily include the respiratory signal. Rather the ECG machine itself has built in processing techniques which estimate the respiratory rate on its own. It is expected that by finding a way to extract this respiratory signal from the ECG machine, that the coherence estimate would be much higher. However, the machine used for testing only provided the raw ECG signal and not the conditioned respiratory signal.

The correlation for the heart rate signal between the ECG machine and the prototype provided coherence above 0.9 for only 2 out of the 7 subjects. However, the results showed a relatively high coherence, above 0.8, for 4 out of the 7 subjects. This can be seen in Table 8 below.

Table 8. HARMONI vs. ECG observed heart rate frequencies and coherence.

Subject:	Heart Rate Frequency (Hz):	HR:	Coherence:
1	1.2	72	0.8807
2	1.0	60	0.4883
3	0.9	54	0.0977
4	1.1	66	0.9897
5	0.9	54	0.8137
6	1.0	60	0.6816
7	1.0	60	0.9812

These results suggest that there was a relatively high coherence for heart rate between the pressure transducer device and the ECG for 4 out of the 7 subjects. While human testing supported this method against the industry standard, survey results were gathered to support an understanding of the need and the environment in which the technology might be applied.

4.2.2 *IRB Explorative Survey*

The surveys were distributed using several different resources, including the VT Rescue squad, and the EMT's at Carilion Roanoke Memorial Hospital. The results from VT Rescue are presented in this report. There were some notable trends in these results that should be pointed out.

VT Rescue's survey results provided broad perspectives. Firstly, there were fourteen survey's completed through VT Rescue. Those who participated had career lengths varying anywhere from a few months to eight years. The average career length of the participants was 3 years. 42.86% had responded to a mass casualty situation. By definition a mass casualty situation is usually one in which the number of patients outweighs the resources of the responders. Many responded to MCI's such as at football games where up to a hundred patients require monitoring. Meanwhile, others responded to car accidents and some to more threatening incidents. Of those that participated, every single one answered that detecting heart rate was often times difficult. The difficulty measuring heart rate was sometimes a result of the type of injury, while other times it was due to the body type of the patient.

71.43% of the participants said that having a deployable chest strap that wirelessly monitored multiple patients simultaneously would be considered a worthwhile technology, if certain conditions were met. The most common condition was that the device not be expensive. Several of the secondary comments concerned sanitation, privacy of the patient, and in what way it would be implemented in to their response environment. The complete collection of survey results can be seen in Appendix E: Results under IRB Survey Results.

4.3 Discussion

With all of the results shown simulated testing and human testing, it is important to discuss the implications that come with those results. Firstly, how does this technology compare to the industry standard technologies that exist? Secondly, how would the technology be implemented in the real world? A discussion of what needs to change, and the assumptions made prior to real world implementation, is required.

4.3.1 *Comparison of Current Technologies with HARMONI*

As seen in the simulation results, the device was capable of accurately estimating respiratory rate on a simulated breathing mannequin. In addition, the human testing results showed that the prototype was capable of detecting the same heart rate signal as an industry standard ECG. While an ECG can provide a complex analysis of the human heart, the pressure transducer device only provides heart rate. However, an ECG is time consuming and expensive, while the pressure transducer device is rapidly deployable and inexpensive. These are the tradeoffs that make the pressure transducer device ideal for mass casualty situations.

There are several areas that require further research. Firstly, signal processing techniques used to condition the data might be implemented such that the estimation of heart and respiratory rates could be done on the board itself. Secondly, how do body weight and gender affect the usefulness of

this technology? It would be expected that a larger subject pool, with a random sample, would provide insight into how these factors affect the results. Despite the need for further research, it has been shown that the prototype was capable of showing high coherence with the ECG results at the expected frequency range of both heart and respiratory rate. The fact that only a few of the subjects showed high coherence for both heart and respiratory rate, suggests that further testing, research, and development is needed to ensure that the device is more reliable regardless of which subject is being tested.

4.3.2 Application of HARMONI in the Real World

The original goal for where this device would be implemented is in a mass casualty situation. As discussed in the introduction, it is a well accepted rule of thumb that each casualty in a mass casualty situation should be reevaluated every 5 minutes. Also as seen in the survey results, an EMT usually allots at least 30 seconds to estimate the heart and respiratory rates of each person. This means that if the device could be attached to a patient and give accurate readings within at least 30 seconds, then it would not only provide an initial reading, but it would provide continuous readings over the next 4 minutes and 30 seconds, before that patient should be reevaluated. The tracking of said patient's vital signs over the course of minutes, is an unparalleled ability in an MCI, and has the potential to be a life saving tool. However, there are several assumptions for such an application.

4.3.3 Assumptions for Real World Application

The first assumption of applying this device in the real world is that it is easy to use for the EMT, such that it would take them less than 30 seconds to apply and get results. A second assumption is that the device can measure heart and respiratory rates through certain layers of clothing, making it a more rapidly applied tool than the industry standard ECG. A third assumption would be that the device is capable of doing all of the signal processing and estimation on the board, without having to send data to a central processing unit. That being said, a central unit could still gather the estimated heart and respiratory rates. This will be discussed further in the next chapter on conclusions and future work.

4.4 Summary of Results

- The method of using pressure transducers to detect heart and respiratory rate was validated.
- Simulated testing revealed that the device was capable of detecting respiratory rate on a simulated breathing mannequin at 6 different respiratory rates.
- A PCB that can be used as a base platform for future work was designed and tested.
- Human testing revealed that 4 out of the 7 subjects had respiratory rates observed using the prototype, which had coherence over 0.9, with the data provided by an ECG machine.
- Human testing revealed that 4 out of the 7 subjects had heart rates observed using the prototype, which had coherence over 0.8, with the data provided by an ECG machine.
- Survey results revealed that over 70% of EMT's recommended that a deployable chest strap for monitoring patients in a mass casualty situation would be useful if certain conditions were met.

Chapter 5: Conclusions and Future Work

This final chapter reviews the results of the study on using pressure transducers for noninvasive detection of heart and respiratory rate, and how these results can be applied in to mass casualty situations. All of the results from the feasibility to human testing suggest that the device could be incorporated in to mass casualty situations to improve patient monitoring. Prior to being implemented in to such a high risk situation, there must first be further research and testing, as well as several improvements to the device's sensors, on board processing, and form factor. This will all be discussed under the section entitled Future Work. In addition, there are several applications of using pressure sensors to monitor movement that will also be explored.

5.1 Summary

Creating a new method for heart and respiratory monitoring for emergency response environments may seem unnecessary at first, when considering the accuracy of current methods. However, it has been shown that under the time constraints of mass casualty situations, a rapidly deployable heart and respiratory monitor that can go over clothing and wirelessly monitor multiple patients would act as a force multiplier for EMT's. Survey results showed that EMT's strongly agree with this statement. It was also shown that using pressure transducers for noninvasive detection of heart and respiratory rates is an affordable alternative to electrode based methods. When combined with a wireless transceiver, it would allow a single EMT to monitor multiple casualties simultaneously.

A literature review revealed that pressure sensors have been used to monitor vital signs. Yet, none of these attempts were targeted towards the mass casualty situation. The contributions of this research were to show that a single affordable pressure sensor and a small strip of medical tubing could be combined to create a cheap, durable, and compact vital signs monitor. In addition, a PCB was developed that would act as a base platform for future pressure transducer based movement sensors. The results of the testing showed relatively high coherence, above 0.8, between the prototype and the ECG device for respiratory signals in 4 out of the 7 subjects. Meanwhile the results also showed a relatively high coherence, above 0.8, between the prototype and the ECG device for heart rate signals in 4 out of the 7 subjects. Such results suggest that there needs to be further research, development, and testing concerning how the results vary based on the test subject and other factors.

Other contributions include the software basis for operating a Bluetooth based data acquisition device. Note this was done while simultaneous operation of both an ADC and an LCD screen over the same I2C bus. Additional contributions, in terms of software, include a sustainable state machine architecture that allows for sub routines to be added and removed from the program with ease. While these contributions make a strong foundation for the technology to be incorporated in to mass casualty situations, there is still much work to be done, and improvements to be made.

5.2 Future Work

The plan for the future application of this device is to have multiple deployable chest straps operating in the same network, with a central processing unit (CPU) located on the first responder in a tablet form. The tablet would display results to the user in the form of a companion application. This application would have options for the first responder to input as they apply the device to the casualty, such as their estimated age and weight. This data will be used to influence alerts on the first responder's CPU, while they are busy triaging other casualties. The CPU will take the information from each patient's heart and respiratory rates, and display it all on one screen for rapid evaluation. An example of this can be seen below in Figure 5-1.



The image shows an ASUS tablet displaying a table with patient data. The table has four columns: ID, ALERT?, HEART RATE, and RESPIRATORY RATE. The rows are labeled A through H. Row A has a yellow background for the alert and green for the heart and respiratory rates. Row B has a yellow background for the alert and green for the heart and respiratory rates. Row C has a green background for the alert and heart rate, and a light green background for the respiratory rate. Row D has a green background for the alert and heart rate, and a light green background for the respiratory rate. Row E has a red background for the alert and green for the heart and respiratory rates. Row F has a green background for the alert and heart rate, and a light green background for the respiratory rate. Row G has a green background for the alert and heart rate, and a light green background for the respiratory rate. Row H has a green background for the alert and heart rate, and a light green background for the respiratory rate. The tablet also shows a status bar at the bottom with the time 03:24 and battery level.

ID	ALERT?	HEART RATE	RESPIRATORY RATE
A	CAUTION BREATHING	74	60
B	CAUTION BREATHING	72	6
C	OK	72	22
D	OK	70	21
E	WARNING HEART RATE	30	20
F	OK	68	24
G	OK	69	19
H	OK	74	20

Figure 5-1. Suggested implementation of emergency response companion application.

Note that in this figure, the green areas are meant to be for quick reference, while the yellow and red highlighted areas are meant to give caution and warning to the user. In this way, an operator's attention could be drawn to the casualties who need most help at any given time, allowing their resources to be better distributed in the emergency medical environment. Once the information is made readily available to the user, it is important to incorporate that information throughout the triage process. This means integrating the information obtained with other medical equipment.

5.2.1 Integration with Other Medical Technology

While the current prototype does not currently communicate with other pre-existing medical equipment and software, a future revision may also have vital sign alerts that coincide with devices such as the Broselow Emergency Tape. In this way, the application of medicine to a patient can first be determined by their size, weight, and vital signs, while the duration and details of each individual triage process can be recorded and used for later evaluation. An SD memory card slot could easily be incorporated in to the device, so that data could be stored and saved for download by the hospital or by higher level care. In this way, all of the patient's vital signs and medical attention from the moment the device was turned on could be on record. The data could be seamlessly transferred directly using the SD

card or by using the Bluetooth technology and a simple .exe file placed on the user's computer station. Besides integrating the device with other medical equipment, the device itself can see improvements as the project moves towards implementation. This begins with one of the most variable factors in the device, the bladder and tubing configuration.

5.2.2 Manufacturing a Custom Bladder

While human testing showed that the current configuration of the pressure transducer proved to be sufficient. There was no evidence to suggest it is the best bladder configuration. In fact, the device likely had difficulty detecting the heart rate in some subjects, due to the bladder configuration. One would suspect that a flatter bladder with a wider area, would allow for a higher sensitivity on a greater number of individuals. Physically, this can be proven considering the change in surface area. A larger contact area between the bladder and the subject's chest will create a larger observed change in volume, and hence pressure. This new form factor could be realistically achieved by approaching a plastics company, such as Plastics One® in Roanoke. This company specializes in custom medical equipment, and would be capable of manufacturing a wider, flatter, bladder, with a nozzle attachment for the pressure transducer. Assuming that the bladder can be improved, the next set of improvements would be the on board sensors.

5.2.3 Improvement of Sensors and Chips

There are several on board sensors that can be reevaluated for improvement. These include the Bluetooth antenna, the ADC, the LCD screen, and most importantly the pressure transducer. Firstly, the Bluetooth antenna's range could be improved by replacing it all together with a broader ranging antenna. The ADC could be improved by finding an alternative ADC with the same precision but a higher reference voltage. A higher resolution as previously discussed would allow the smaller changes in pressure due to heart rate to be more easily detected. This could be accomplished by finding an ADC with a reference voltage above the current 1.25 V, while still having 16 bit precision and 3 V supply voltage. The current LCD screen accomplished all of the objectives laid out for the full system prototype. However, even it could be improved by incorporating a programmable dimming circuit. In this way the LCD screen could be dimmed just like any other smart device.

Finally, the pressure transducer is perhaps the most critical sensor, when considering the whole system, which could be improved. While there was no comparable device that had a 3 V operating range, was affordable, and had more than 600 mV/kPa sensitivity, this does not mean that a better sensor will not come along. Higher sensitivities would mean changes to the data acquisition circuit. Any future changes, should follow the steps outlined in section 3.3.6 Circuit Design. Assuming that the sensors and chips could be improved, the algorithm used to estimate the heart and respiratory rates should be integrated in to the board itself.

5.2.4 On Board Heart and Respiratory Rate Detection

While almost all of the signal processing done in this research has been done post data collection. The integration of the technology in to a mass casualty situation requires near real time

heart and respiratory rate estimation. In order to achieve this, the embedded device must be capable of performing simple signal processing techniques such as filtering, denoising, or Fourier analysis. These techniques would make it possible to separate two individual signals from the original combined signal. A simple peak-to-peak estimation technique could then estimate the heart and respiratory rates in real time, by measuring the time in between peaks, and extrapolating this to see how many beats or breaths per minute are occurring.

One change that would certainly allow for better peak to peak estimation is to increase the sampling frequency. This can be accomplished by finding a new processor with more RAM. More RAM will allow for larger arrays of data to be stored in the system memory, as well as allow for the signal processing to take place on the board. With all of the processing being transitioned from post processing in to real time on board processing, having accurate data results is perhaps the last piece of the puzzle when it comes to future work. To accomplish this, the pressure data could be fused with other small sensors' data to verify its results.

5.2.5 Fusing Pressure Data with an Accelerometer

The benefits of simultaneously correlating pressure transducer data with other sensors' data, is increased accuracy and robustness. A simple, cheap, sensitive, and small sensor that could easily be incorporated in to the design is a single axis accelerometer. The pressure transducer and bladder configuration, in a sense, provides displacement of the chest due to movement. Similarly a well placed single axis accelerometer on the same board would provide the acceleration of the chest due to movement. Double integrating these results over time, would provide displacement. A simple cross spectrum analysis between the pressure transducer results and the accelerometer's results should allow for the heart and respiratory signals to be extracted more easily. In addition, a coherence estimate would provide confidence to these results. The fusion of such data is the future of this technology, and is almost necessary as a backup, in case one of the sensors were to fail during operation.

5.3 Other Applications

With the assumption that the device is capable of detecting heart and respiratory rate in emergency mass casualty situations, there come several other applications in to which it could be incorporated. From patient tracking through a mattress, to smart phone applications, to work out monitors, there are several areas that this technology could be easily applied.

5.3.1 Tracking Patients Movement and Vital Signs through a Mattress

One of the first applications is in the scenario of tracking movement, respiration, and heart rate through a mattress. If the device can be improved to the point of tracking movement and vital signs through a mattress, then it could be implemented in nursing home environments. Elderly patients are very susceptible to falls, especially in nursing home environments. This device could therefore be a cheap, easily integrated technology, capable of monitoring patients' movements and then wirelessly alerting medical professionals that a patient has attempted to get out of bed unassisted.

Similarly it could provide a less invasive method for sleep monitoring, allowing a person to experience a more natural sleep environment without the respiratory impedance sensors and electrode based methods having to be used to track their vital signs. These sensors are often cumbersome and take away from the whole idea of sleep evaluation, because the people who are being studied are not used to being attached to such sensors in their normal sleep environment. A wireless monitor placed under the mattress would therefore be a worthwhile alternative that would in turn provide a more wholesome and normal sleep experience.

5.3.2 Creating a Smartphone Application for Every Day Use

Another application of the technology is to provide the everyday consumer with a method for monitoring their heart and respiratory rates. To do this, a smart phone application that could observe the results from the pressure sensor device would be ideal. Such an application would allow the smart phone to read information over Bluetooth, from the device, over an extended period of time. This information could then be displayed such that the user could see their vital signs tracked over the course of a day. In addition, the application could provide alerts based on abnormal results.

5.3.3 Creating Noninvasive and Waterproof Exercise Monitor

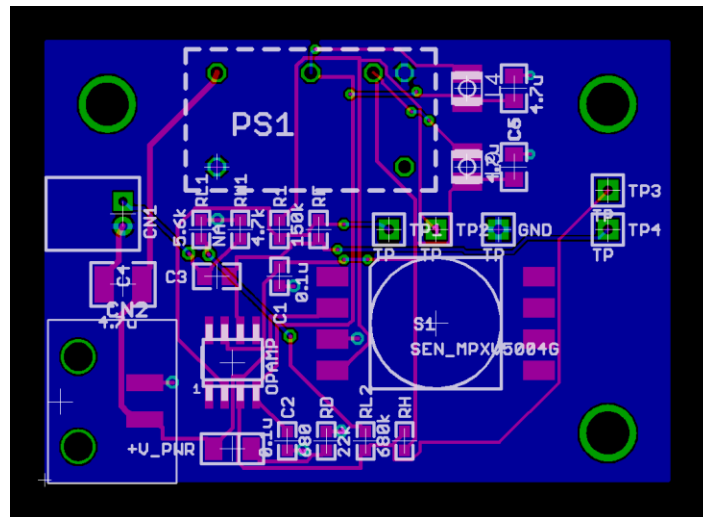
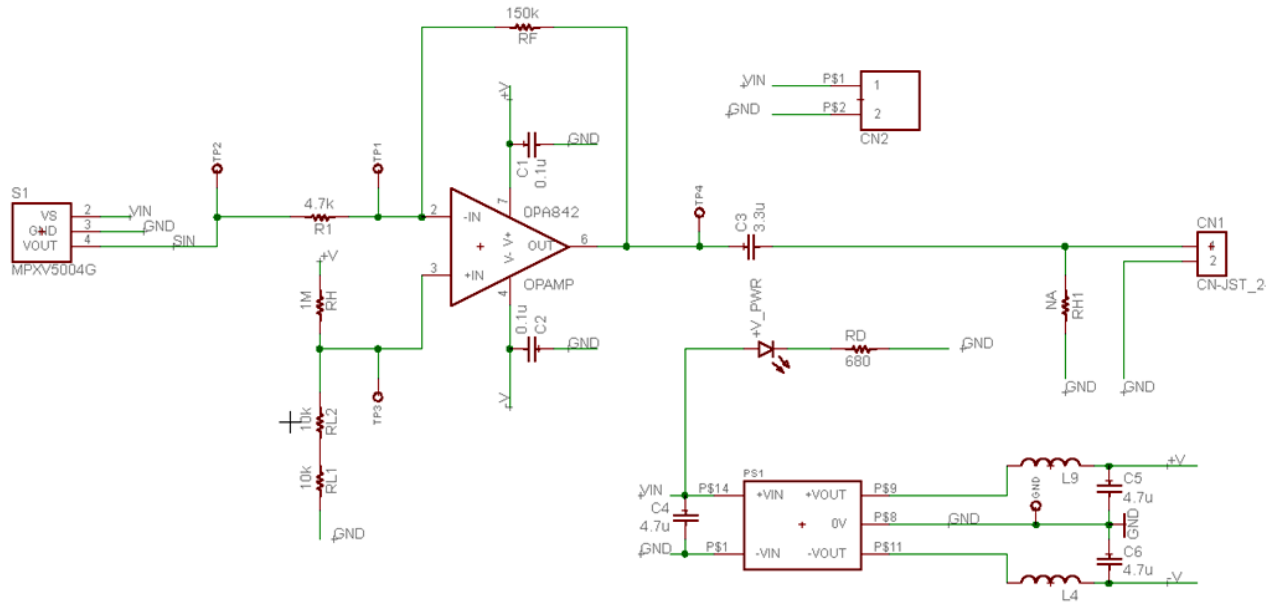
Along the same lines as the smart phone application, would be the implementation of this device in to a mobile workout monitor. Many of the electrode based workout monitors give accurate results for a short amount of time. The drawbacks begin when the ECG leads irritate the user's skin, or detract from the runner's usual stride, or the results are skewed by sweat and water buildup around the leads. A waterproof chest strap monitor that requires low levels of pressure against the runner's chest might be the answer to all of these issues. Being able to filter out all of the extraneous movement artifacts that occur during exercise would be an issue in implementing this device. However, understanding such an issue would also improve the original application of monitoring vital signs in a mass casualty situation, where indeed extraneous movement artifacts would also be present.

References

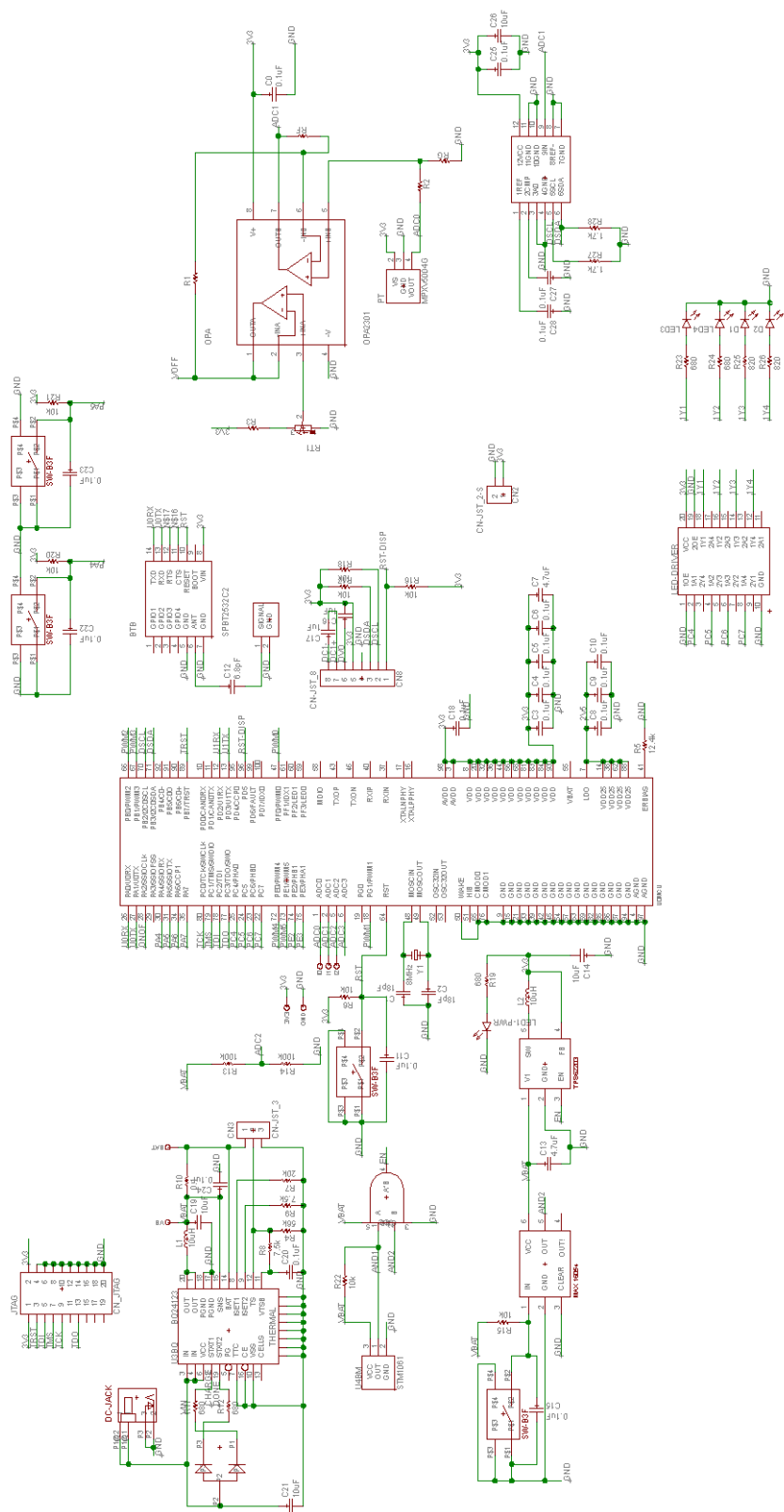
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Appendix A: Circuit Boards and Schematics

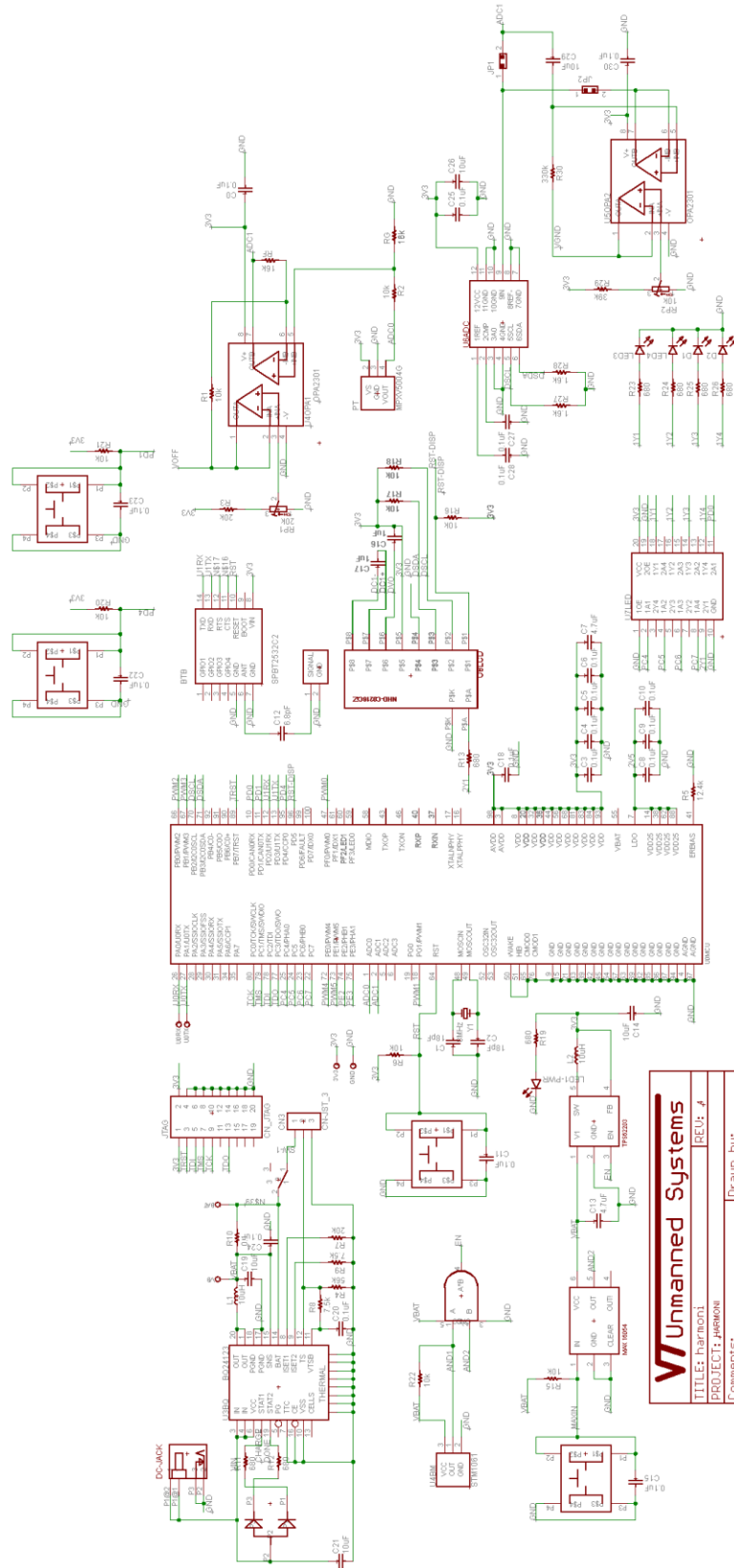
Rev 2, Simulation Board



Rev 3, First Full System Prototype Board



Rev 4, Second Full System Prototype Board



Unmanned Systems

TITLE: Harmoni REV: 1

PROJECT: HARMONI Drawn by: #harmoni B Downem

Comments: Date: not saved!

Sheet: 1/1

Appendix B: MATLAB Code

MATLAB Code for Feasibility Results

```
clc
clear all
close all

%% import the data
data = importdata('feasibility_5.txt');
null = importdata('feasibility_0.txt');

%% define a window, such as 30 seconds
x = data((length(data)-3100):(length(data)-100));
y = null((length(null)-3100):(length(null)-100));

%% normalize the data
x = x - mean(x);
y = y - mean(y);

%% define parameters, time and frequency domain
Fs = 100; % Sampling frequency
Nyq = Fs/2; % Nyquist frequency
T = 1/Fs; % Sample time

Lx = length(x); % Length of signal x
Ly = length(y); % Length of signal y

tx = 0:1/Fs:(length(x) - 1)/Fs; % Time vector
rx = (Fs)/Lx; % Frequency Resolution
fx = 0:rx:Fs - rx; % Frequency vector up to Fs

%% compute cross spectrums using averaging
m = 1;
k = 600;

gxx = 0; % auto spectrum x
gyy = 0; % auto spectrum y
gxy = 0; % cross spectrum x,y
gyx = 0; % cross spectrum y,x

while m < Lx - k;
    gxx = gxx + conj(fft(x(m:m+k-1))).*fft(x(m:m+k-1));
    gyy = gyy + conj(fft(y(m:m+k-1))).*fft(y(m:m+k-1));
    gxy = gxy + conj(fft(x(m:m+k-1))).*fft(y(m:m+k-1));
    gyx = gyx + conj(fft(y(m:m+k-1))).*fft(x(m:m+k-1));
    m = m+k;
end

gxx = (2*gxx)/k;
gyy = (2*gyy)/k;

pxy = unwrap(angle(gxy));
```

```

gxy = (abs(2*gxy)/k);
gyx = (abs(2*gyx)/k);

fa = 0:Fs/length(gxx):Fs - Fs/length(gxx);

%% plot results
figure()
plot(tx,x)
xlabel('time (s)')
ylabel('voltage (V)')
title('Feasibility Signal')

figure()
plot(fa,gxx/Lx)
axis([0 Nyq 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Feasibility Autospectrum')

figure()
plot(fa,gxx/Lx)
axis([0 4 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Feasibility Autospectrum Zoomed')

%% calculate and display the signal to noise ratios
[C,I] = max(gxx);
snr_r = sqrt(max(gxx(1:I-3)/max(gyy)));
snr_r_db = 20*(log10(snr_r));

snr_h = sqrt(max(gxx)/max(gyy));
snr_h_db = 20*(log10(snr_h));

fprintf('SNR for Respiration = %4.3f\n',snr_r);
fprintf('SNR for Respiration (dB) = %4.3f dB\n\n',snr_r_db);

fprintf('SNR for Heart Rate = %4.3f\n',snr_h);
fprintf('SNR for Heart Rate (dB) = %4.3f dB\n',snr_h_db);

```

MATLAB Code for Simulation Results

```
clc
clear all
close all

%% import the data
data = importdata('proof_13.txt');
null = importdata('proof_14.txt');

%% choose a window size
block = 30; % in seconds
block = block*1000;
x = data(100:block + 99);
y = null(100:block + 99);

%% normalize the data
x = x - mean(x);
y = y - mean(y);
original = x;
to = 0:1/1000:(length(original) - 1)/1000;

%% decimate the data
d = 10;
size = length(x);
n = 1;

while n < d + 1;
    a(n,:) = decimate(x(n:size - d + n),d);
    b(n,:) = decimate(y(n:size - d + n),d);
    n = n + 1;
end

x = a;
y = b;
Fs = 1000/d;

%% define parameters
Nyq = Fs/2; % Nyquist frequency
T = 1/Fs; % Sample time
Lx = length(x); % Length of signal
tx = 0:1/Fs:(length(x) - 1)/Fs; % Time vector
rx = (Fs)/Lx; % Frequency Resolution

Ly = length(y); % Length of signal
ty = 0:1/Fs:(length(y) - 1)/Fs; % Time vector
ry = (Fs)/Ly; % Frequency Resolution

fx = 0:rx:Fs - rx; % Frequency vector for x
fy = 0:ry:Fs - ry; % Frequency vector for y

%% compute cross spectrums using averaging
m = 1;
k = d;
```



```

gxx = 0; % auto spectrum x
gyy = 0; % auto spectrum y
gxy = 0; % cross spectrum x,y
gyx = 0; % cross spectrum y,x

fprintf('Averages = %4.3f\n',k);

while m < d;
    gxx = gxx + conj(fft(x(m,:))).*fft(x(m,:));
    gyy = gyy + conj(fft(y(m,:))).*fft(y(m,:));
    gxy = gxy + conj(fft(x(m,:))).*fft(y(m,:));
    gyx = gyx + conj(fft(y(m,:))).*fft(x(m,:));
    m = m+k;
end

gxx = (2*gxx)/(size/d);
gyy = (2*gyy)/(size/d);

pxy = unwrap(angle(gxy));
gxy = (abs(2*gxy)/(size/d));
gyx = (abs(2*gyx)/(size/d));

fa = 0:Fs/length(gxx):Fs - Fs/length(gxx);

%% plot results
figure()
plot(to,original)
xlabel('time (s)')
ylabel('voltage (V)')
title('Simulation Signal')

figure()
plot(fa,gxx/Lx)
axis([0 Nyq 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Simulation Autospectrum')

figure()
plot(fa,gxx/Lx)
axis([0 4 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Simulation Autospectrum Zoomed')

figure()
subplot(2,1,1)
plot(to,original)
xlabel('time (s)')
ylabel('voltage (V)')
title('Simulation Signal')

subplot(2,1,2)
plot(fa,gxx/Lx)

```

```
axis([0 4 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Simulation Autospectrum Zoomed')

%% calculate and display the signal to noise ratios
snr_r = sqrt(max(gxx)/mean(gyy));
snr_r_db = 20*(log10(snr_r));

fprintf('SNR for Respiration = %4.3f\n',snr_r);
fprintf('SNR for Respiration (dB) = %4.3f dB\n\n',snr_r_db);
```

MATLAB Code for Human Testing Results

```
clc
clear all
close all

%% import the data
x = importdata('full_6harmoni.txt');
y = importdata('full_6ecg.txt');

%% define a window
x = x((length(x)-1199):length(x));
y = y((length(y)-1199):length(y));

%% normalize the data
x = x - mean(x);
y = y - mean(y);

%% define parameters
fs = 20;
Nyq = fs/2;
Tx = length(x)/fs;
rx = 1/Tx;
Lx = length(x);
Ty = length(y)/fs;
ry = 1/Ty;
Ly = length(y);
t = 0:1/fs:(Lx - 1)/fs;
fx = 0:rx:fs - rx;
fy = 0:ry:fs - ry;

%% optional filter
% fL = .7/Nyq;
% fH = 2/Nyq;
% [b,a] = butter(4,fL,'high');
% x = filter(b,a,x);

%% spectral estimation
xdft = fft(x);
ydft = fft(y);

%% compute cross correlation
cxy = xcorr(x,y);
Lz = length(cxy);
lag = -(length(cxy)/2)/fs:1/fs:(length(cxy)/2 - 1)/fs;

%% compute cross spectrums using averaging
m = 1;
k = 200;
% k = 120, gives 6 second blocks of data, w/ fr = 0.1667 Hz, 10 averages
% k = 200, gives 10 second blocks of data, w/ fr = 0.1 Hz, 6 averages

gxx = 0; % auto spectrum x
gyy = 0; % auto spectrum y
```

```

gxy = 0; % cross spectrum x,y
gyx = 0; % cross spectrum y,x

while m < Lx - k;
    gxx = gxx + conj(fft(x(m:m+k-1))).*fft(x(m:m+k-1));
    gyy = gyy + conj(fft(y(m:m+k-1))).*fft(y(m:m+k-1));
    gxy = gxy + conj(fft(x(m:m+k-1))).*fft(y(m:m+k-1));
    gyx = gyx + conj(fft(y(m:m+k-1))).*fft(x(m:m+k-1));
    m = m+k;
end

gxx = (2*gxx)/k;
gyy = (2*gyy)/k;

pxy = unwrap(angle(gxy));
gxy = (abs(2*gxy)/k);
gyx = (abs(2*gyx)/k);

fa = 0:fs/length(gxx):fs - fs/length(gxx);

%% compute the coherence function
cf = (gxy.*gyx)./(gxx.*gyy);

%% plot results
figure()
subplot(2,1,1)
plot(t,x)
xlabel('time (sec)')
ylabel('voltage (V)')
title('Full System Prototype Signal')

subplot(2,1,2)
plot(fa,gxx/Lx)
axis([0 Nyq 0 1.15*max(gxx/Lx)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('Full System Prototype Autocorrelation')

figure()
subplot(2,1,1)
plot(t,y)
xlabel('time (sec)')
ylabel('voltage (V)')
title('ECG Signal')

subplot(2,1,2)
plot(fa,gyy/Ly)
axis([0 Nyq 0 1.15*max(gyy/Ly)])
xlabel('freq (Hz)')
ylabel('amplitude^2 / 2 (V^2)')
title('ECG Autocorrelation')

% plot cross correlation
figure()
plot(lag,cxy)

```

```

xlabel('time lag (sec)')
ylabel('Rxy')
title('Cross Correlation')

% plot cross spectrum in magnitude and phase
figure()
subplot(2,1,1)
plot(fa,gxy/Lx)
axis([0 Nyq 0 1.15*max(gxy/Lx)])
xlabel('freq (Hz)')
ylabel('magnitude (V^2)')
title('Cross Spectrum Magnitude')

subplot(2,1,2)
plot(fa,pxy)
axis([0 Nyq 1.15*min(pxy) 1.15*max(pxy)])
xlabel('freq (Hz)')
ylabel('phase (radians)')
title('Cross Spectrum Phase')

figure()
plot(fa,cf)
axis([0 Nyq 0 1])
xlabel('freq (Hz)')
ylabel('Coherence')
title('Coherence Function')

```

Battery Requirements for Full System Prototype

```
clear all
close all
clc

% define battery components and current draw
Im = 100*10^-3;
Ibt = 40*10^-3;
Iled = 20.5*10^-3;
Ipt = 10*10^-3;
Ioa = 9.5*10^-3;
Isd = 15*10^-6;
Iof = 7*10^-6;
Icm = 3.5*10^-6;
Ivd = 0.9*10^-6;
Iad = 3.5 *10^-3;
% I = [3.5*10^-6 100*10^-3 3.5*10^-3 7*10^-6 10*10^-3 20.5*10^-3 ...
%      9.5*10^-3 40*10^-3 0.9*10^-6 15*10^-6];

I = [Im Ibt Iled Ipt Ioa Isd Iof Icm Iad Ivd];
Vs = 3.3;

% battery specs
Vbat = 3.7;
% efficiency
e = 0.90;

% calculate the 24 hour battery life requirement
b = (24*sum(I)*Vs)/(e*Vbat);

% 24 hour operation
note1 = sprintf('Required for 24 hour operation %.4f mAhrs /n ',1000*b);
% 16 hour operation
note2 = sprintf('Required for 16 hour operation %.4f mAhrs',1000*b*(16/24));
% 8 hour operation
note3 = sprintf('Required for 8 hour operation %.4f mAhrs',1000*b*(8/24));

% 24, 16, and 8 hour operation
fprintf('Required for 24 hour operation %.4f mAhrs \nRequired for 16 hour
operation %.4f mAhrs \nRequired for 8 hour operation %.4f mAhrs \n\n'...
,1000*b,1000*b*(16/24),1000*b*(8/24))

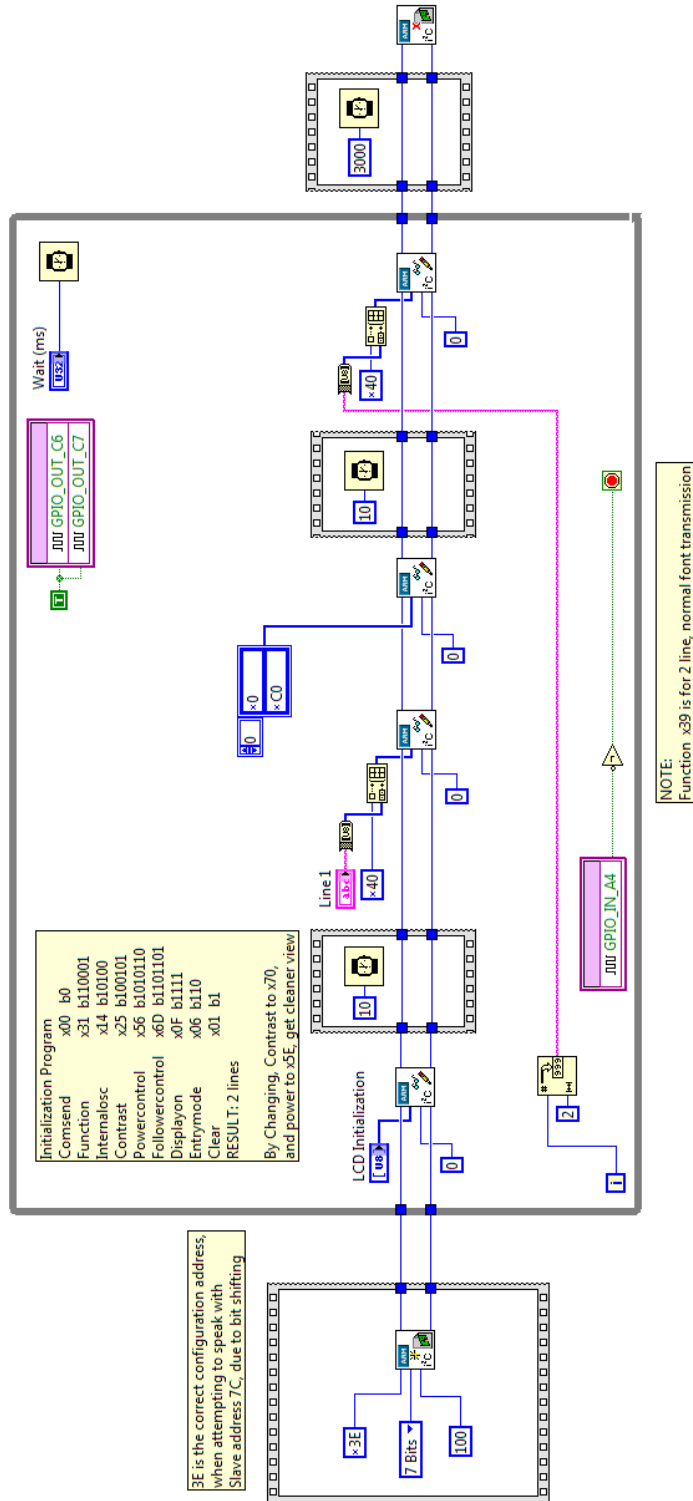
% Choose Which Battery Life
% b = b;          for 24 hrs
% b = b*(16/24); for 16 hrs
% b = b*(8/24);  for 8 hrs

% percent of time lost
xof = ((Iof*24)/(b));
yof = (xof*24*60*60);
fprintf('The On/Off Controller (MAX16054) Quiescent Current Subtracts %.8f
percent of usage per day \n',xof)
fprintf('or %.4f seconds lost per day \n',yof)
```

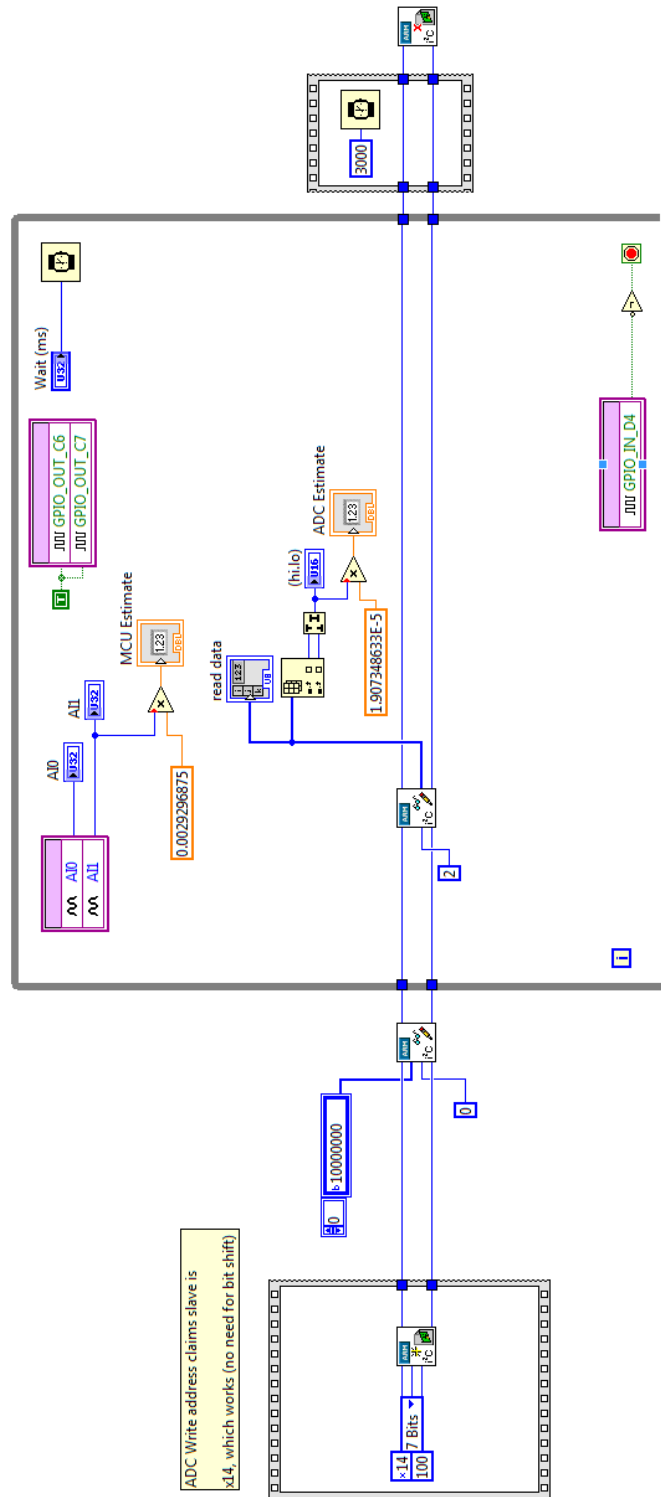
```
xsd = ((Isd*24)/(b));  
ysd = (xsd*24*60*60);  
fprintf('The Step Down (TPS62203) Quiescent Current Subtracts %0.8f percent  
of usage per day \n',xsd)  
fprintf('or %0.4f seconds lost per day \n',ysd)
```

Appendix C: LabVIEW Code

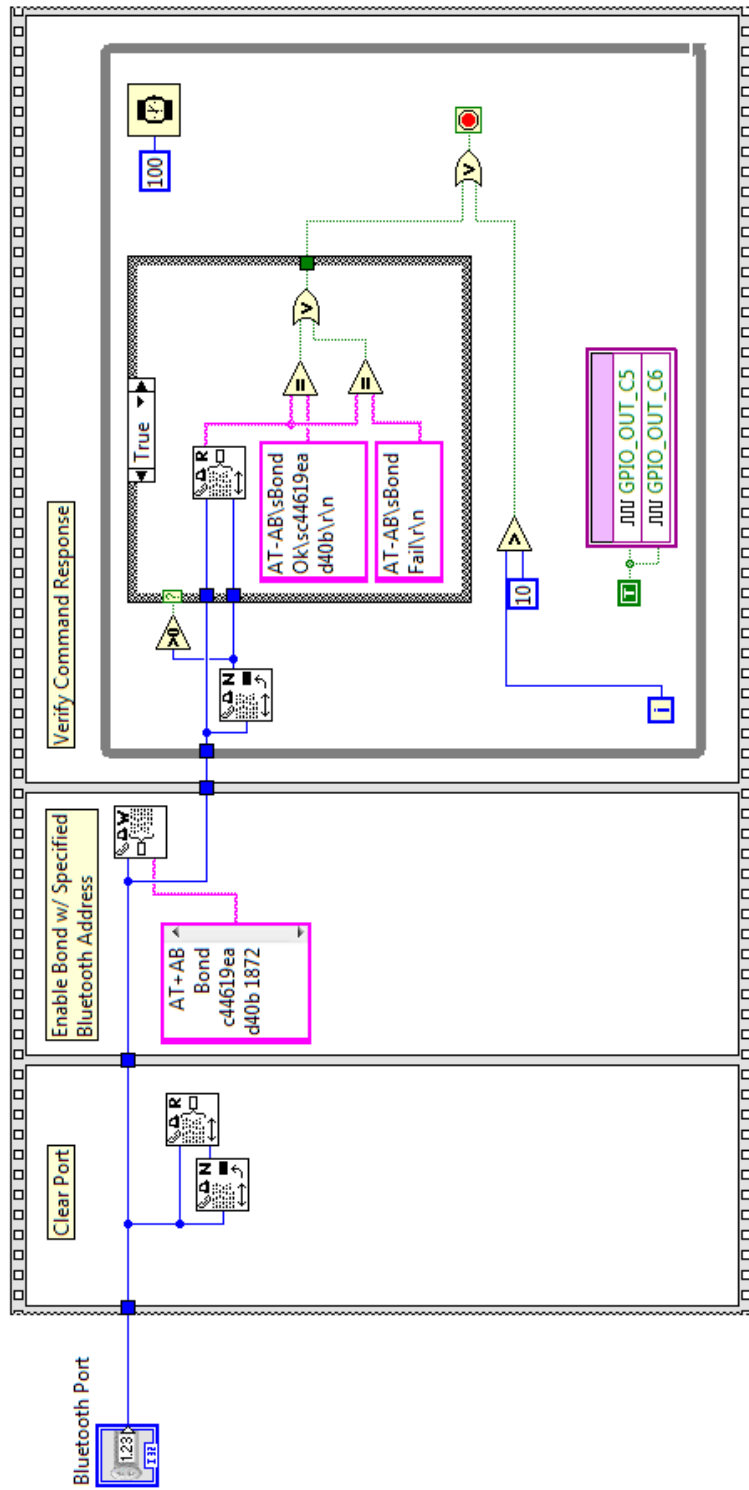
Example for the LCD Configuration:



Example for the ADC Configuration:



Example for the Bluetooth Configuration



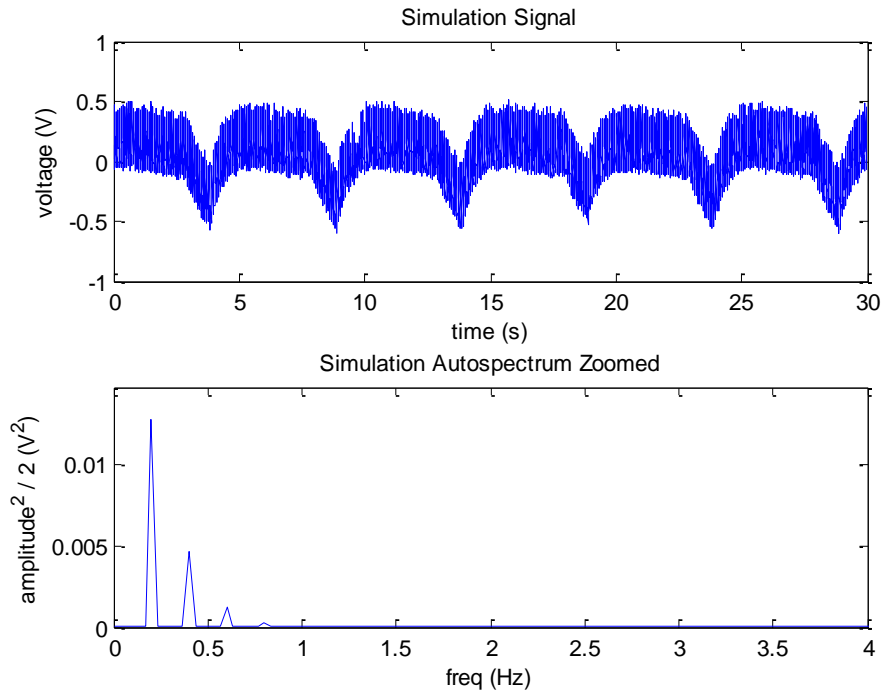
Appendix D: IRB Forms

Explorative Survey

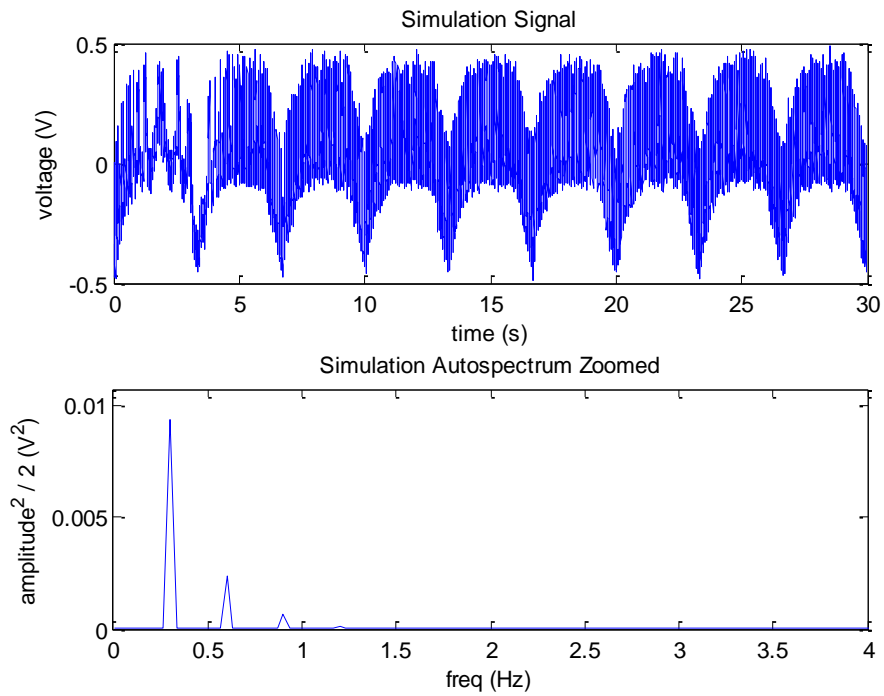
Survey No.	Date:
Interviewer.	Time:
Survey Questionnaire	
<hr/>	
<p>Have you been trained as an EMT, a first responder, nurse, or a doctor?</p>	
<p>How long have you been in this profession?</p>	
<p>Have you ever been involved in mass casualty situation? If yes, explain...</p>	
<p>By what means did you take a person's pulse or respiratory rate in an emergency situation?</p>	
<p>About how much time does it take to determine the pulse or respiratory rate?</p>	
<p>Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...</p>	
<p>Have you ever needed to monitor multiple patients at the same time?</p>	
<p>What are some of the most common types of patients that you have responded to?</p>	
<p>If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...</p>	

Appendix E: Results

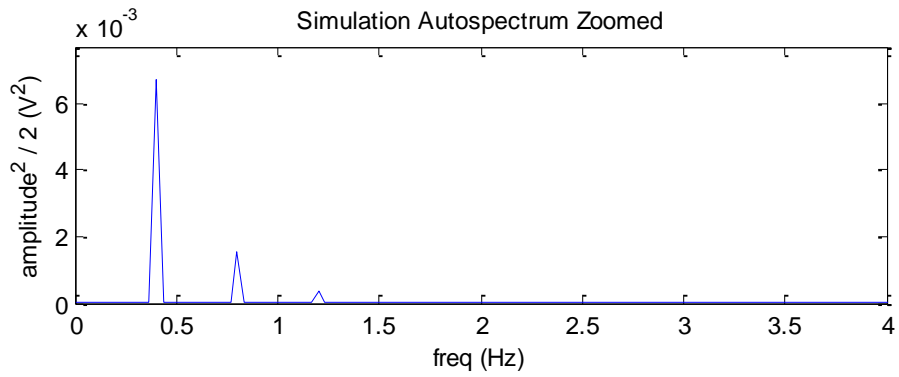
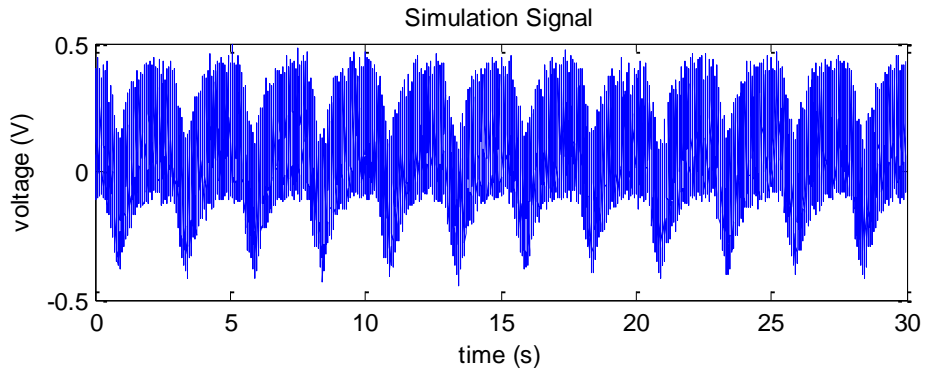
Simulation Testing Results



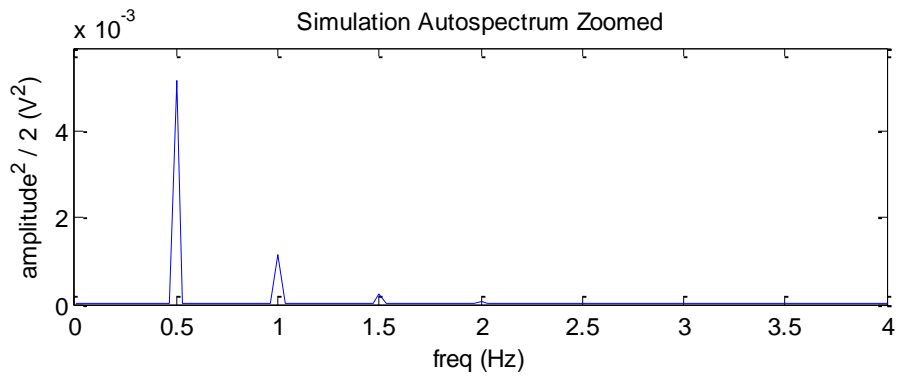
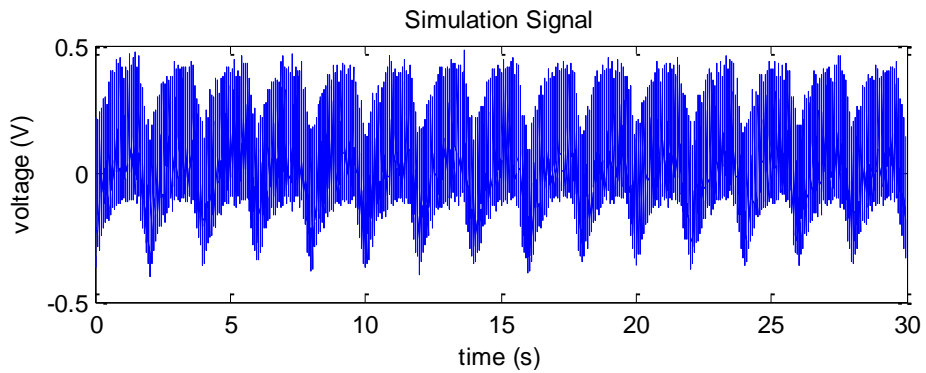
12 breaths per minute time domain and autocorrelation results.



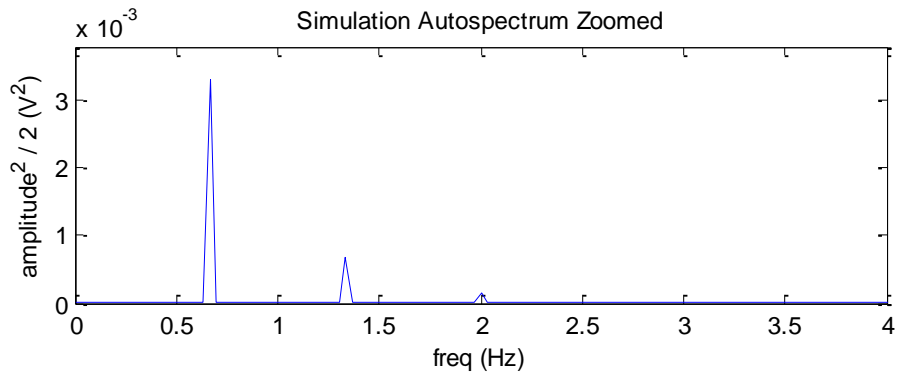
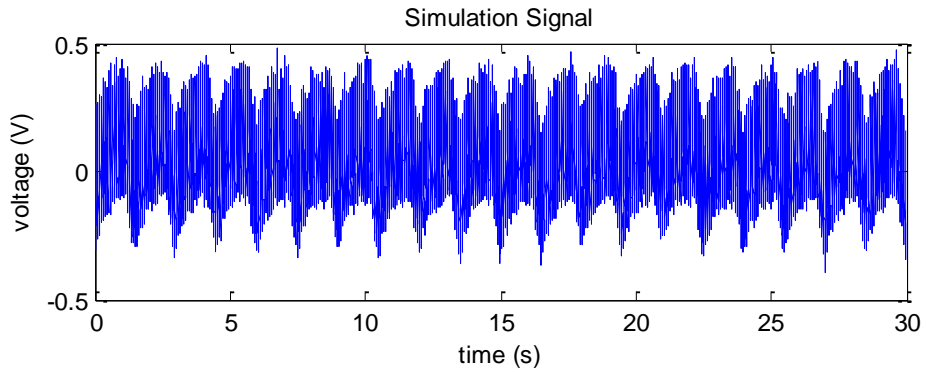
18 breaths per minute time and frequency domain results.



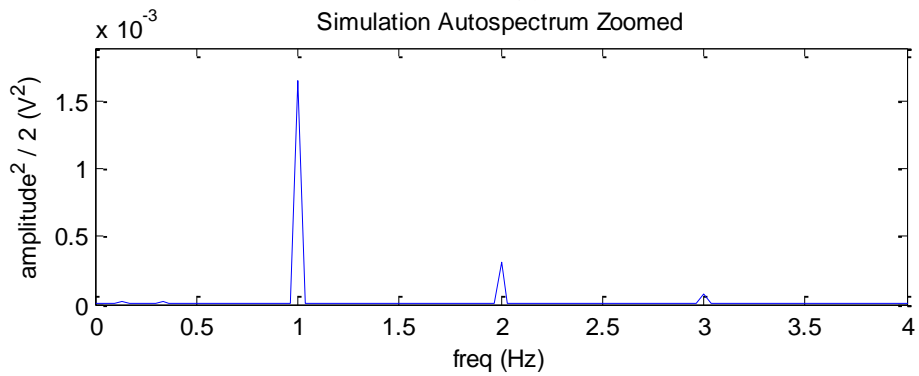
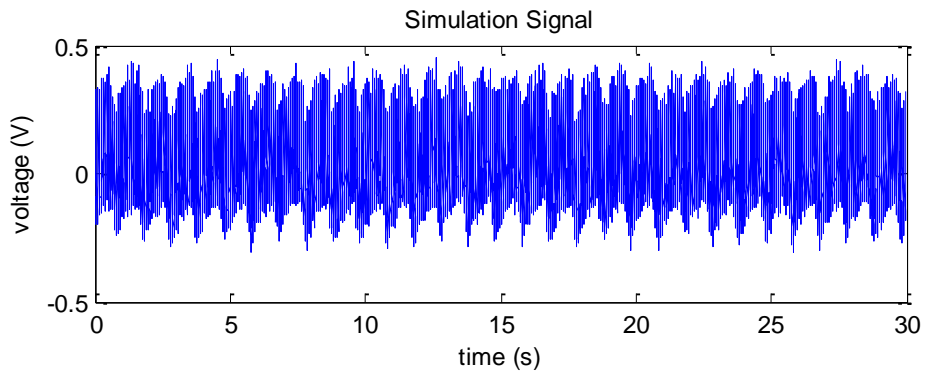
24 breaths per minute time and frequency domain results.



30 breaths per minute time and frequency domain results.

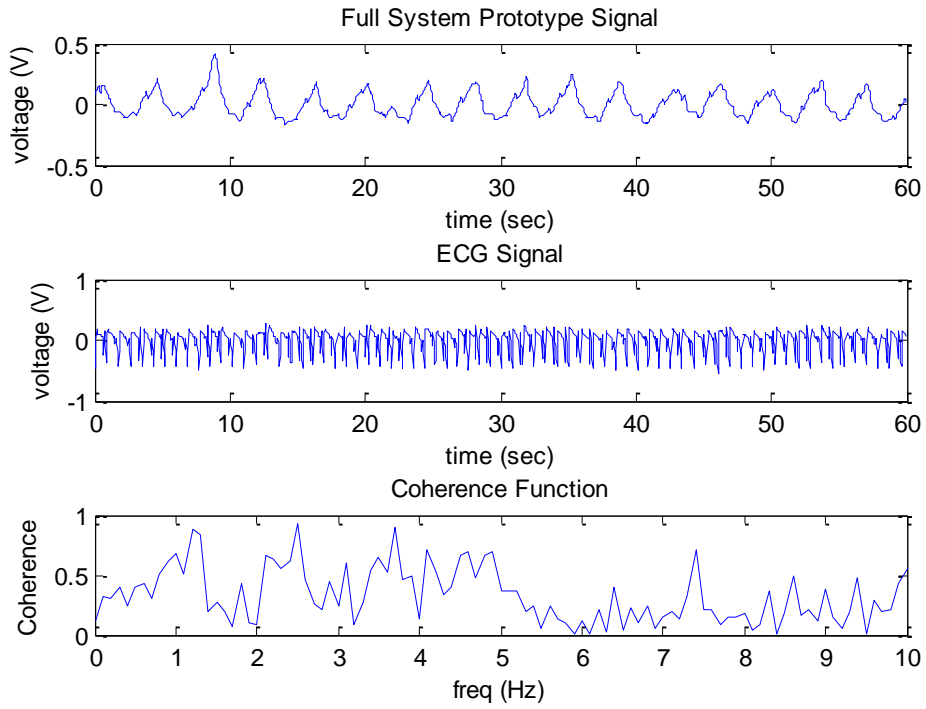


40 breaths per minute time and frequency domain results.

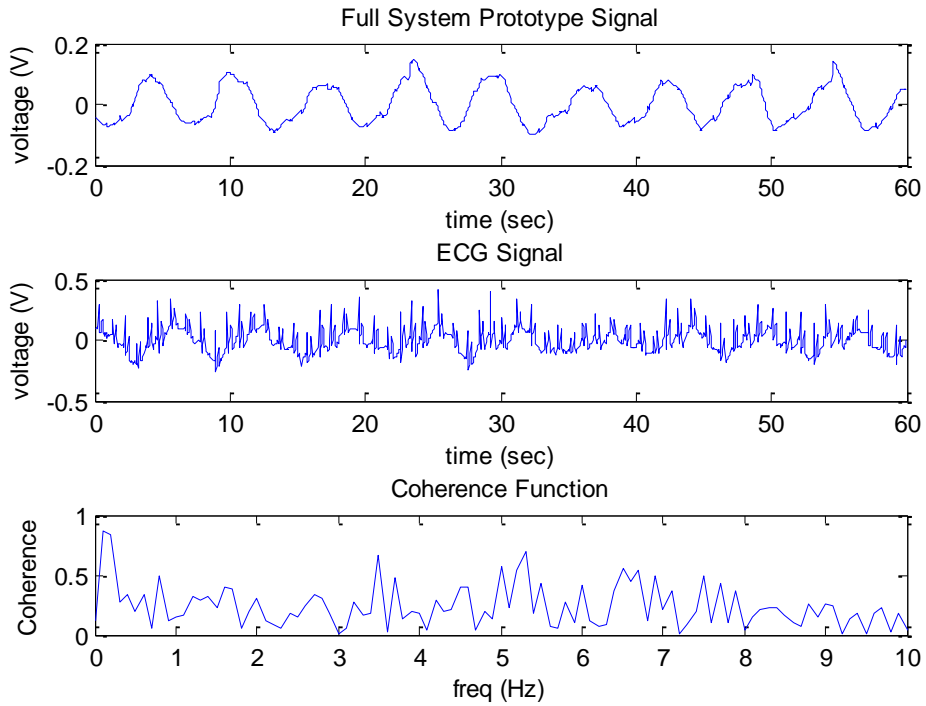


60 breaths per minute time and frequency domain results.

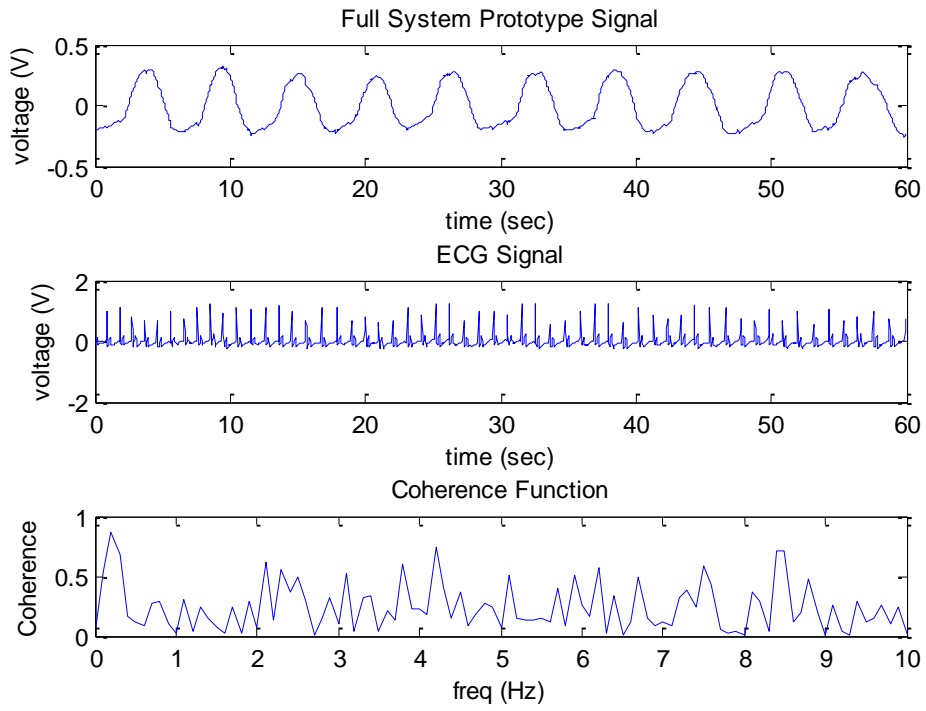
IRB Benchmark Testing Results



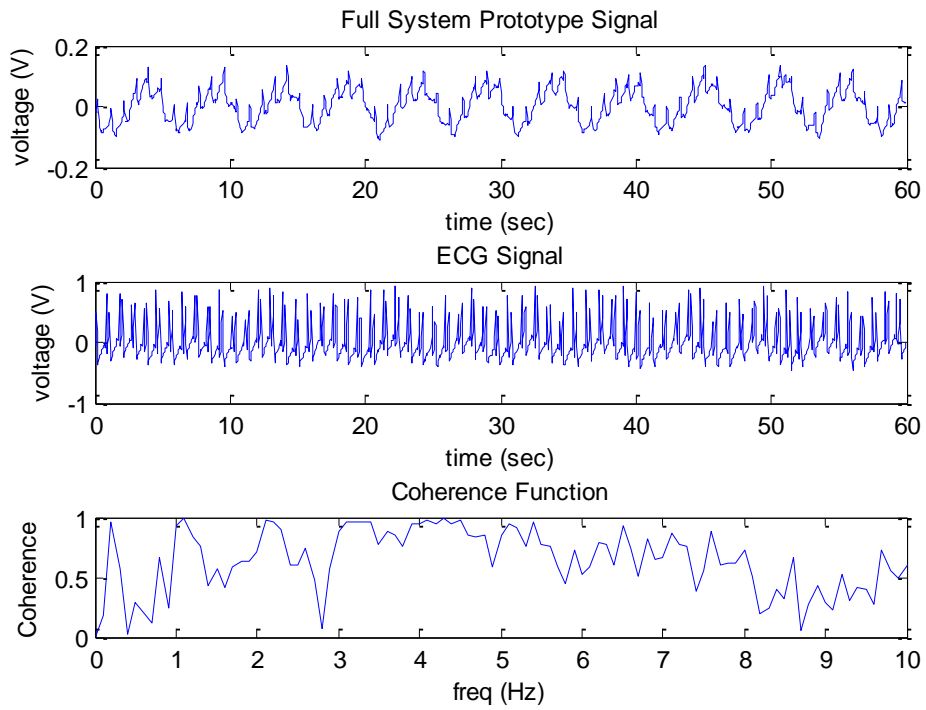
Subject 1 results.



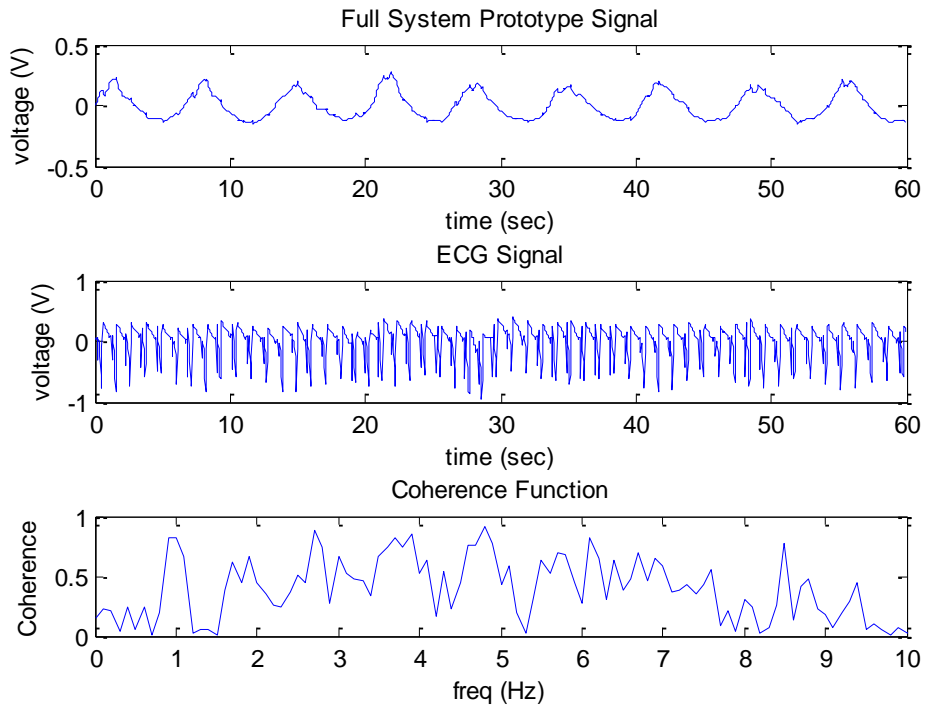
Subject 2 results.



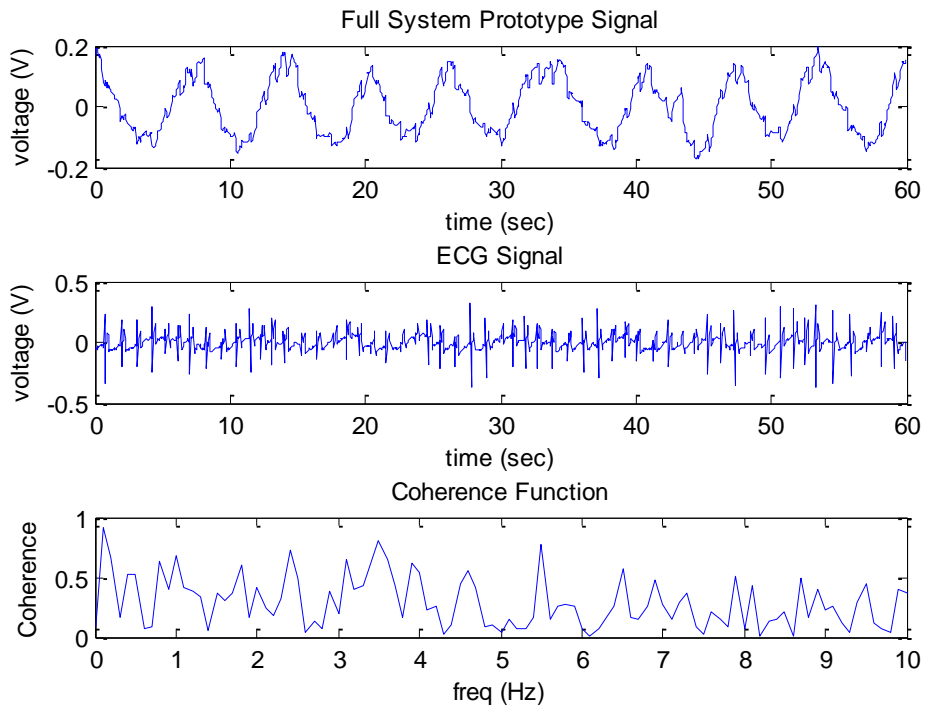
Subject 3 results.



Subject 4 results.



Subject 5 results.



Subject 6 results.

IRB Survey Results

SURVEY 1

Have you been trained as an EMT, a first responder, nurse, or a doctor?

EMT

How long have you been in this profession?

8 Years

Have you ever been involved in mass casualty situation? If yes, explain...

Yes. April 16th

By what means did you take a person's pulse or respiratory rate in an emergency situation?

Depends. If awake, radial pulse, if unconscious, carotid pulse. Respiratory rate based on rise and fall of shoulders or chest.

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds to be accurate and determine any anomalies in rhythm.

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes. When heart rate and blood pressure are low, extremity pulses are hard to find, and you have to return to the carotid.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Respiratory rate is the most estimated vital sign. You can typically tell if they are breathing fast, slow or normally.

Have you ever needed to monitor multiple patients at the same time?

Yes. First aid room at football games.

What are some of the most common types of patients that you have responded to?

Musculoskeletal injuries, alcohol related injuries. Transports from Schiffert Health Center

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Only worth while in controlled situations such as first aid rooms, but this means someone would be dedicated to watch a screen. Depends on overall patient load. Low pt load would work better, but not worth it in hectic situations.

SURVEY 2

Have you been trained as an EMT, a first responder, nurse, or a doctor?

EMT

How long have you been in this profession?

8 Years

Have you ever been involved in mass casualty situation? If yes, explain...

Yes. April 16th

By what means did you take a person's pulse or respiratory rate in an emergency situation?

Depends. If awake, radial pulse, if unconscious, carotid pulse. Respiratory rate based on rise and fall of shoulders or chest.

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds to be accurate and determine any anomalies in rhythm.

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes. When heart rate and blood pressure are low, extremity pulses are hard to find, and you have to return to the carotid.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Respiratory rate is the most estimated vital sign. You can typically tell if they are breathing fast, slow or normally.

Have you ever needed to monitor multiple patients at the same time?

Yes. First aid room at football games.

What are some of the most common types of patients that you have responded to?

Musculoskeletal injuries, alcohol related injuries. Transports from Schiffert Health Center

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Only worth while in controlled situations such as first aid rooms, but this means someone would be dedicated to watch a screen. Depends on overall patient load. Low pt load would work better, but not worth it in hectic situations.

SURVEY 3

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes. And, though you didn't ask this, and EMT-I

How long have you been in this profession?

5 years

Have you ever been involved in mass casualty situation? If yes, explain...

yes, April 16, 2007

By what means did you take a person's pulse or respiratory rate in an emergency situation?

pulse by palp or pulse-ox resps by counting over 15 sec and estimating

About how much time does it take up to estimate the pulse or respiratory rate?

less than a minute

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

yes, BP can be difficult.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Estimate? I'll ball park it. You can tell by a quick glance if the patient is breathing at an abnormal rate. So, yes, I might give a number in the normal range without spending much time on it. BP and pulse, no. If I can't get a BP, I'll say so.

Have you ever needed to monitor multiple patients at the same time?

Yes. Anyone who has been stationed in a first aid room at Lane Stadium during a football game has had to keep track of multiple patients at one time.

What are some of the most common types of patients that you have responded to?

Pretty much everything. Heat stroke, intoxication, gun shot wounds, altered mental status, diabetic emergency, heart issues, asthma, difficulty breathing, broken bones...

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Yes, if it were reliable and easy to deploy. I'm assuming you mean a chest strap that would go all the way around a patient. It would be very difficult to put something like that on many patients. For example, trauma patients are immediately backboarded. Bariatric patients are nearly impossible to lift. Would it interfere with cardiac monitoring (12-leads)? Would it require patients to be exposed in the chest area, leading to privacy issues? So, maybe in a perfect world it's a good idea, but it would depend on the device and the circumstances.

SURVEY 4

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes (EMT)

How long have you been in this profession?

2 years

Have you ever been involved in mass casualty situation? If yes, explain...

No

By what means did you take a person's pulse or respiratory rate in an emergency situation?

I have taken radial pulses every time I had to assess pulse rate/quality and I observed the rise and fall of the chest to assess respiratory rate.

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds to 1 minute total

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes. When patients have low blood pressure or a weak heart beat or small sunken veins it can be difficult. I've found the hardest pulses to locate are usually on small women.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Never. I don't estimate or make up vital signs-that's dangerous. I take the time to get it right, or I get another crew member to try.

Have you ever needed to monitor multiple patients at the same time?

Yes.

What are some of the most common types of patients that you have responded to?

Flu-like symptoms, sports injuries, and drunks.

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

It seems like a good idea, but with something that invasive (particularly for women who don't want to show off the pups) it might not always be warranted. In a true emergency, why not just use the heart monitors we already have? The only real advantages I could see over a heart monitor are 1) wireless technology may be easier to deploy, especially with multiple patients and 2) our entire agency only has 10 or so heart monitors, and they aren't cheap. If we had a 20+ MCI, throwing in a dozen of your chest strap devices might save us time, money, and effort.

SURVEY 5

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Currently a paramedic and also in medical school

How long have you been in this profession?

Began EMS in 2005 with initial certification in 2006. Began medical school in 2005

Have you ever been involved in mass casualty situation? If yes, explain...

Yes, 4/16 at VT

By what means did you take a person's pulse or respiratory rate in an emergency situation?

Radial pulse x 15 seconds and multiply by four. Observe chest rise and fall x 15 seconds and multiply by four. Though sometimes in an emergency setting I just check for quality and approximate rate of both pulse and RR during quick patient interactions such as triage

About how much time does it take up to estimate the pulse or respiratory rate?

As stated above, about 15 seconds for a count. Can take shorter time if you just want an approximation. If there is a non emergency situation, pulse and RR are measured by counting for 30 seconds and multiplying by two, in theory providing a more accurate set of vital signs.

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Sometimes can be challenging to locate peripheral pulses on trauma patients secondary to hypotension

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Only in emergent, life threatening situations.

Have you ever needed to monitor multiple patients at the same time?

Not really in the out of hospital setting

What are some of the most common types of patients that you have responded to?

Minor trauma, breathing difficulty, abdominal pain, altered level of consciousness

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

It sounds great but the cost of implementing such a device would likely be restrictive

SURVEY 6

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes.

How long have you been in this profession?

8 months

Have you ever been involved in mass casualty situation? If yes, explain...

No.f

By what means did you take a person's pulse or respiratory rate in an emergency situation?

I take pulses and respiratory rates on a daily basis when on the scene of a 911 call.

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds for each.

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes, bariatric patients can have layers of fat covering their arteries and therefore it can be difficult to initially locate a pulse.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Never.

Have you ever needed to monitor multiple patients at the same time?

Yes.

What are some of the most common types of patients that you have responded to?

Sports related injuries.

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Yes and no. It would depend what the device is measuring. For vitals such as pulse, a machine can calculate the vital but cannot calculate the quality of the pulse (strong, weak, thready). You would have to make sure that the technology could completely detect any vital signs that may be taken.

SURVEY 7

Have you been trained as an EMT, a first responder, nurse, or a doctor?

2.5 years

How long have you been in this profession?

3 years

Have you ever been involved in mass casualty situation? If yes, explain...

Yes.... Car accidents

By what means did you take a person's pulse or respiratory rate in an emergency situation?

By palpation

About how much time does it take up to estimate the pulse or respiratory rate?

15 seconds

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes. Some people have wear pulses or very shallow respirations

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Very rarely

Have you ever needed to monitor multiple patients at the same time?

Yes

What are some of the most common types of patients that you have responded to?

Medical, sports injuries

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Yea. It would allow me to focus on pt care and assessing them rather than getting vitals continuously.

SURVEY 8

Have you been trained as an EMT, a first responder, nurse, or a doctor?

yes

How long have you been in this profession?

less than a year

Have you ever been involved in mass casualty situation? If yes, explain...

no

By what means did you take a person's pulse or respiratory rate in an emergency situation?

pulse by palpation either radial or carotid we also use a pulse-ox Respirations by visually counting number of breaths

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds for both

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

yes, on some patients especially elderly it can be difficult to hear a blood pressure

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Never

Have you ever needed to monitor multiple patients at the same time?

no

What are some of the most common types of patients that you have responded to?

sport related injuries, ETOH, respiratory difficulties

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

yes and no, if there are multiple patients then yes also if the device can accurately read respiration, however on a normal call we almost always like to take manual vital signs for example in the case of taking a pulse if take it manually you can feel rate, strength and regularity, if it is taken manually you only get a rate

SURVEY 9

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes

How long have you been in this profession?

Two Years

Have you ever been involved in mass casualty situation? If yes, explain...

No

By what means did you take a person's pulse or respiratory rate in an emergency situation?

feeling and counting a pulse and observing respirations

About how much time does it take up to estimate the pulse or respiratory rate?

no longer than 15 seconds

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

yes. Sometimes the pulse is very weak, which can mean finding a blood pressure is hard as well. It typically means you just try a second time.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Often I estimate the respiratory rate, but none other

Have you ever needed to monitor multiple patients at the same time?

Yes

What are some of the most common types of patients that you have responded to?

Sports injury (ankle, knee, wrist) and abdominal pain

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

yes. If there is one person in the back of an ambulance or multiple patients, it becomes difficult to obtain a set of vitals as needed while determining required information about the patients condition and demographics.

SURVEY 10

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes

How long have you been in this profession?

9 months

Have you ever been involved in mass casualty situation? If yes, explain...

I participated in an MCI drill during fall 2011 semester. Just a drill.

By what means did you take a person's pulse or respiratory rate in an emergency situation?

Manually; radial pulse

About how much time does it take up to estimate the pulse or respiratory rate?

Pulse- 30 seconds Respiratory rate- 30 seconds

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes, some individuals have difficult radial pulses to find; sometimes it is difficult to count respirations when they are shallow

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Never.

Have you ever needed to monitor multiple patients at the same time?

Only during the MCI drill.

What are some of the most common types of patients that you have responded to?

Athletic injuries; syncopal episodes; difficulty breathing; ETOH alcohol involved

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

No. No technology compares to manual readings

SURVEY 11

Have you been trained as an EMT, a first responder, nurse, or a doctor?

EMT

How long have you been in this profession?

1 year

Have you ever been involved in mass casualty situation? If yes, explain...

no

By what means did you take a person's pulse or respiratory rate in an emergency situation?

manual

About how much time does it take up to estimate the pulse or respiratory rate?

15-30 seconds

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

yes, it can be difficult in the truck especially if the person has weak pulses. it is also difficult to listen for a blood pressure and lung sounds because you can hear the engine of the truck

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

I never make up vitals, if you can't get them you include that in the report

Have you ever needed to monitor multiple patients at the same time?

yes

What are some of the most common types of patients that you have responded to?

sports injuries, illness, falls

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

Not for what we do. If I did long transports, then maybe. It isn't too hard to monitor a patient and I don't know how much I would trust a chest strap in the truck

SURVEY 12

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Yes

How long have you been in this profession?

Since Fall 2011

Have you ever been involved in mass casualty situation? If yes, explain...

No

By what means did you take a person's pulse or respiratory rate in an emergency situation?

I watch the chest for movement for a respiration rate and check the radial pulse

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes, radial pulses can be difficult to find

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

Never

Have you ever needed to monitor multiple patients at the same time?

No

What are some of the most common types of patients that you have responded to?

Drunks and sports injuries

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

I am not trained to use cardiac monitors to monitor the heart

SURVEY 13

Have you been trained as an EMT, a first responder, nurse, or a doctor?

EMT

How long have you been in this profession?

5 years

Have you ever been involved in mass casualty situation? If yes, explain...

Yes. Multiple from bus accidents to multi car accidents, as well as countless trainings.

By what means did you take a person's pulse or respiratory rate in an emergency situation?

Pulse: I use my index and middle finger to feel the radial pulse. If they are unconscious, i use the same fingers and feel their carotid. Respirations: look at the chest and count...

About how much time does it take up to estimate the pulse or respiratory rate?

You never estimate a vital sign.....thats piss poor patient care. When i assess their vitals, it takes 30 seconds for pulse and an additional 30 seconds for respirations.

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

Yes, especially on geriatrics. If it is difficult, you should note the problem and NEVER 'estimate' vitals.

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

NEVER ESTIMATE VITAL SIGNS!!!! The job of an EMT aka a medical professional is to obtain vitals, thats you job...dont estimate, because when you do that, and you get it wrong you could kill someone. If you cant get an accurate reading, note that, dont estimate or a better way yo put it....lie.

Have you ever needed to monitor multiple patients at the same time?

Yes.

What are some of the most common types of patients that you have responded to?

Sports injuries, general illness, ETOH abuse.

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

For the right price, yes. It would help in the circumstances when u have multiple patients and few providers, but that rarely happens, so the price would have to be right to be worth it.

SURVEY 14

Have you been trained as an EMT, a first responder, nurse, or a doctor?

Trained as an EMT

How long have you been in this profession?

8 months

Have you ever been involved in mass casualty situation? If yes, explain...

no

By what means did you take a person's pulse or respiratory rate in an emergency situation?

pulse be radial pulse and respirations by observation/auscultation.

About how much time does it take up to estimate the pulse or respiratory rate?

30 seconds for each = 1 minute total

Have you ever had a difficult time finding the vital signs on a patient? If yes, explain...

yes- I had an elderly patient who had recent surgery on their left hand. it was difficult to find a radial pulse in that hand

How often do you make an estimate of vital signs, because you couldn't get an accurate reading, or didn't have enough time?

when it comes to a noncritical patient, say a minor lac or superficial abrasion, I have estimated respirations/

Have you ever needed to monitor multiple patients at the same time?

yes- football games

What are some of the most common types of patients that you have responded to?

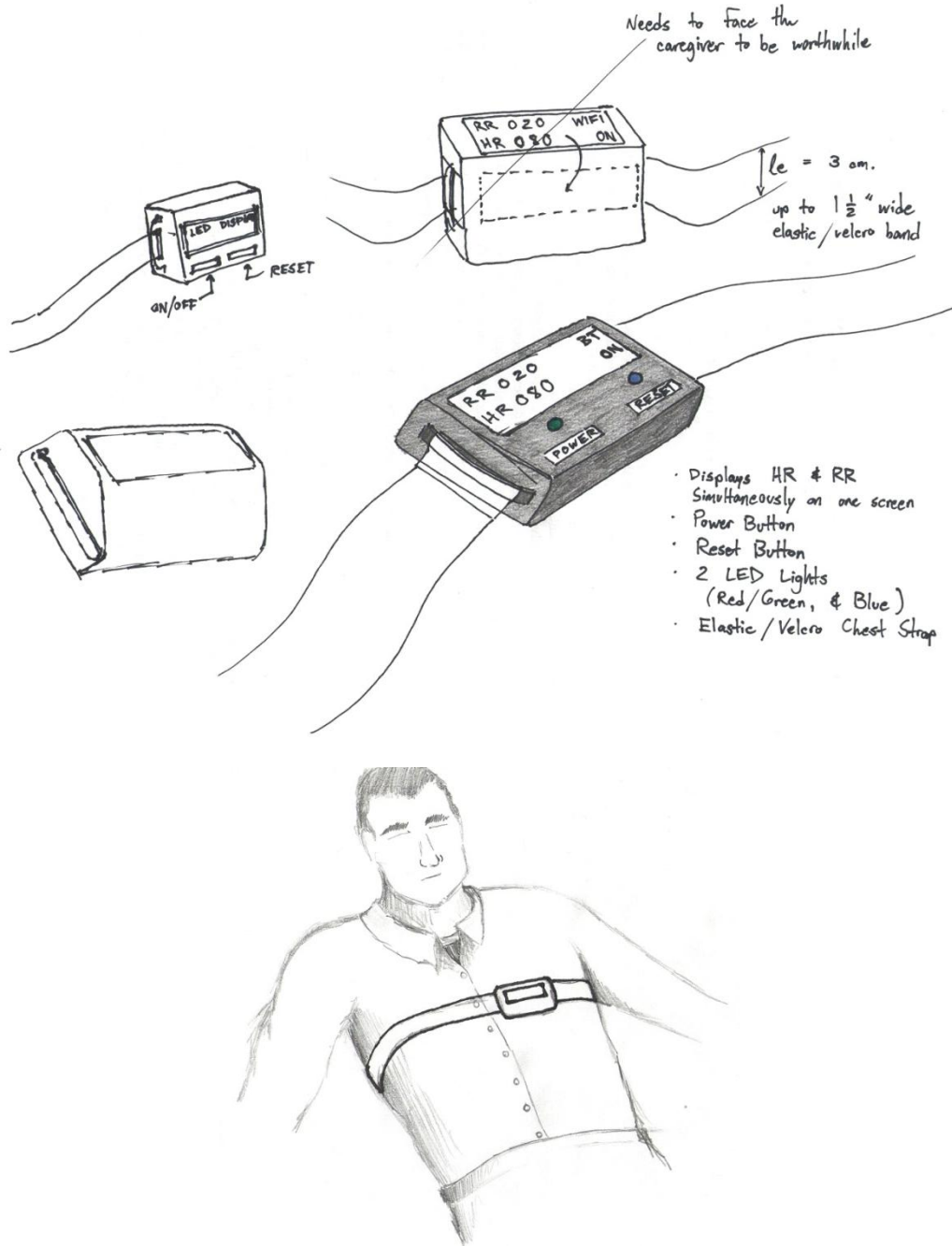
Medical issues- chest pain, nausea, ETOH, etc.

If you could deploy a chest strap on to patients and wirelessly monitor multiple patients simultaneously, would you consider it to be a worthwhile technology? Explain...

it seems expensive... plus we are taught to treat the patient, not the machine. peoples "normal range" vital signs vary

Appendix F: Miscellaneous

Prototype Design Drawings



All drawings in this section are created and owned by Matthew R. B. Dowden

Datasheets

Product Description	Product Number	Datasheet URL
Analog to Digital Converter	LTC2471	http://cds.linear.com/docs/Datasheet/24713f.pdf
And Gate	SN74AHC1G08	http://www.ti.com/lit/ds/symlink/sn74ahc1g08.pdf
BQ Charge Controller	BQ24123	
Bluetooth Antenna	Rufa2v4 3030A58	http://www.antenova.com/Product%20Specs/AE020157-N-Product-Specification-Rufa.pdf
Bluetooth Module	SPBT2532C2-AT	http://www.st.com/internet/com/TECHNICAL_RESOURCES/TECHNICAL_LITERATURE/DATASHEET/CD00243909.pdf
LCD Screen	NHD-C0216CiZ-FSW-FBW-3V3	http://www.newhavendisplay.com/specs/NHD-C0216CiZ-FSW-FBW-3V3.pdf
LCD Screen User Manual	ST7032	http://www.newhavendisplay.com/app_notes/ST7032.pdf
LED Driver	sn74lvc244a	http://www.ti.com/lit/ds/symlink/sn74lvc244a.pdf
Low Power Voltage Detector	STM1061	http://www.st.com/internet/com/TECHNICAL_RESOURCES/TECHNICAL_LITERATURE/DATASHEET/CD00065467.pdf
Microcontroller	LM3S8962	http://www.ti.com/lit/ds/symlink/lm3s8962.pdf
On/Off Controller	MAX16054	http://datasheets.maxim-ic.com/en/ds/MAX16054.pdf
Op-amp	OPA2301	http://www.ti.com/lit/ds/symlink/opa2301.pdf
Project Box		http://www.pactecenclosures.com/pdfs/drw_HM.pdf
Power Jack	PJ-053D	http://www.cui.com/Product/Resource/DigiKeyPDF/PJ-053D.pdf
Pressure Transducer	MP3V5004G	http://www.freescale.com/files/sensors/doc/data_sheet/MP3V5004G.pdf
Step Down Converter	TPS62203	http://www.ti.com/general/docs/lit/getliterature.tsp?genericPartNumber=tps62203&fileType=pdf
Switch, SPDT	SLS121RA04	http://documents.tycoelectronics.com/commerce/DocumentDelivery/DDEController?Action=srchrtv&DocNm=1825116&DocType=CD&DocLang=EN
Switch, Momentary	TL1105BF160Q	http://spec.e-switch.com/P-J/P010003C.pdf