



Office of Research Compliance
 Institutional Review Board
 1880 Pratt Drive (0497)
 Blacksburg, Virginia 24061
 540/231-4991 Fax: 540/231-0959
 E-mail: moored@vt.edu
 www.irb.vt.edu


FWA00000572(expires 7/20/07)
 IRB # is IRB00000667.

DATE: March 24, 2006

MEMORANDUM

TO: Christopher L. North
 Andrew Sabri
 Alain Fabian

Approval date: 3/23/2006
 Continuing Review Due Date:3/8/2007
 Expiration Date: 3/22/2007

FROM: David M. Moore 

SUBJECT: **IRB Expedited Approval:** "Notifications on Large High Resolution Displays" , IRB # 06-204

This memo is regarding the above-mentioned protocol. The proposed research is eligible for expedited review according to the specifications authorized by 45 CFR 46.110 and 21 CFR 56.110. As Chair of the Virginia Tech Institutional Review Board, I have granted approval to the study for a period of 12 months, effective March 23, 2006.

As an investigator of human subjects, your responsibilities include the following:

1. Report promptly proposed changes in previously approved human subject research activities to the IRB, including changes to your study forms, procedures and investigators, regardless of how minor. The proposed changes must not be initiated without IRB review and approval, except where necessary to eliminate apparent immediate hazards to the subjects.
2. Report promptly to the IRB any injuries or other unanticipated or adverse events involving risks or harms to human research subjects or others.
3. Report promptly to the IRB of the study's closing (i.e., data collecting and data analysis complete at Virginia Tech). If the study is to continue past the expiration date (listed above), investigators must submit a request for continuing review prior to the continuing review due date (listed above). It is the researcher's responsibility to obtain re-approval from the IRB before the study's expiration date.
4. If re-approval is not obtained (unless the study has been reported to the IRB as closed) prior to the expiration date, all activities involving human subjects and data analysis must cease immediately, except where necessary to eliminate apparent immediate hazards to the subjects.

Important:

If you are conducting **federally funded non-exempt research**, this approval letter must state that the IRB has compared the OSP grant application and IRB application and found the documents to be consistent. Otherwise, this approval letter is invalid for OSP to release funds. Visit our website at <http://www.irb.vt.edu/pages/newstudy.htm#OSP> for further information.

cc: File
 Department Reviewer:Donald S. McCrickard

Invent the Future

