Battelle
The Business *of* Innovation

Centers for Public Health Research and Evaluation
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September 7, 2006

Talmage Holmes, PhD MPH
Battelle CPHRE
2971 Flowers Road South
Suite 233
Atlanta, GA 30341-5404

Dear Dr. Holmes:

As Chair of the Battelle/CPHRE Institutional Review Board (IRB) I have reviewed the continuing review and modifications submission dated 8/18/2006 for the study entitled "Human Behavior Fire Study" (FG487102-10) and grant expedited approval to continue with this study. The changes are minor and do not affect the risk/benefits ratio.

As with all Battelle/CPHRE studies, this study will be subject to continuing review again next year. Your current approval expires 9/6/2007. We will send you notification at the appropriate time. In the meantime, should any additional change occur in your protocol, please inform the IRB and submit the changes for review. Similarly, the IRB needs to be notified in the event of any injury or unexpected outcome arising from this study.

Sincerely,

Margaret Pennybacker

Margaret Pennybacker, PhD, CIP
IRB Chair

cc: Brigette Brevard
Kevin Heaton
Jan Jaeger
Anthony Santella

Battelle/Centers for Public Health Research and Evaluation

100 Capitola Drive, Suite 301
Durham, NC 27713
Federal-wide Assurance No. FWA00004696 (IRB No. 284)

INSTITUTIONAL REVIEW BOARD NOTICE OF APPROVAL

PROJECT DIRECTOR: Talmage Holmes

PROJECT TITLE: Human Behavior Fire Study

CLIENT: CDC PROTOCOL DATE: **8/18/06**

BATTELLE PROJECT CODE: FG487102-10 or PROPOSAL NUMBER: _*(if preaward)*

NATURE OF REVIEW: *(check one)*

- [ ] FULL MEETING DATE: _____
- [X] EXPEDITED *(specify reason)*: no problems reported; minor changes; no change to risk/benefits
- [ ] EXEMPT *(specify reason)*: _____

TYPE OF APPROVAL: *(check one)*

- [ ] PRELIMINARY. SCHEDULE NEXT REVIEW PRIOR TO INVOLVEMENT OF HUMAN SUBJECTS.
- [ ] PRETEST/PILOT TEST. SCHEDULE NEXT REVIEW PRIOR TO FULL IMPLEMENTATION.
- [ ] FULL IMPLEMENTATION.
- [X] RENEWAL/CONTINUING REVIEW.
- [X] AMENDMENT DATED 8/18/2006

Please note the following requirements:

PROBLEMS OR ADVERSE REACTIONS: If any problems in treatment of human subjects or unexpected adverse reactions occur as a result of this study, you must notify the IRB Chairperson immediately, then complete an Adverse Event/Incident Report and forward it to the CPHRE IRB Administrator.

CHANGES IN PROTOCOL: If there are any changes in procedures or study protocol, you must notify the IRB Chairperson and submit the revisions for review before they are implemented.

RENEWAL: You are required to apply for renewal of approval at least annually for as long as the study is active unless the Board finds it necessary to require more frequent reviews. Your next continuing review date should be on or before / / .

Margaret Pennybacker

IRB Chairperson

9,7,06

Date

Margaret R. Pennybacker, PhD, CIP
Print or Type Name

X Copy of approved Informed Consent on file.

cc: Project Director
IRB Administrator