Battelle
The Business *of* Innovation
Centers for Public Health Research and Evaluation
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October 29, 2005

Diane R. Burkom, MA
Battelle CPHRE
6115 Falls Road, Suite 200
Baltimore, MD 21209

Dear Ms. Burkom:

As acting chair of the Battelle/CPHRE Institutional Review Board (IRB) I have reviewed the continuing review and modifications submission dated 10/24/2005 for the study entitled "Second Injury Control and Risk Survey (ICARIS-2)" (FG446622-02) and grant expedited approval to continue with this study. The modifications are minor and do not change the risk/benefits ratio.

As with all Battelle/CPHRE studies, this study will be subject to continuing review again next year. We will send you notification at the appropriate time. In the meantime, should any additional changes occur in your protocol or questionnaire, 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,

Christopher W. Lyu, MPA
Acting IRB Chair

cc: Brigette Brevard
Kevin Heaton
Jan Jaeger

Battelle/Centers for Public Health Research and Evaluation

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

INSTITUTIONAL REVIEW BOARD NOTICE OF APPROVAL

PROJECT DIRECTOR: Diane R. Burkom

PROJECT TITLE: Second Injury Control and Risk Survey (ICARIS-2)

CLIENT: CDC PROTOCOL DATE: **10/24/05**

BATTELLE PROJECT CODE: FG446622-02 or PROPOSAL NUMBER: _*(if preaward)*

NATURE OF REVIEW: *(check one)*

- [ ] FULL MEETING DATE: _____
- [X] EXPEDITED *(specify reason)*: minimal risk; minor change; 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.
- [X] AMENDMENT DATED 10/24/2005

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. Your next continuing review date should be on or before **10/29/06**.

IRB Chairperson

10/29/05
Date

Christopher W. Lyu, MPA
Print or Type Name

X Copy of approved Informed Consent on file.

cc: Project Director
IRB Administrator