UNIVERSITY OF MISSISSIPPI MEDICAL CENTER

2500 North State Street
Jackson, Mississippi 39216-4505

Institutional Review Board
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DHHS FWA # 00003630

Approval Notice
Renewal

January 9, 2009

Thomas H. Mosley, PhD
Medicine
University of Mississippi Medical Center
2500 North State Street
Jackson, MS 39216-4505

RE: IRB File # 1985-0122
Atherosclerosis Risk in Communities (ARIC) Study

Dear Dr. Mosley:

Your Renewal was reviewed and approved by the Expedited review process on January 9, 2009. You may continue this research.

Please note the following information about your approved research protocol:

- Protocol Approval period: January 30, 2009 - January 29, 2010
- Approved Enrollment #: 100000
- Participant Population: Adults - Healthy/Controls
- Performance Sites: UMC
- Expedited Category(ies): 45 CFR 46.110(b) and/or 21 CFR 56.110(b)(8a) - Continuing review of research originally approved by the convened IRB where (i) the research is permanently closed to the enrollment of new participants; (ii) all participants have completed all research-related interventions; and (iii) the research remains active only for long-term follow-up of participants.

Renewal Review History:

Receipt Date	Submission Type	Review Process	Review Date	Review Action
01/08/2009	Renewal	Expedited	01/09/2009	Approved

Please remember to:

→ Use **the IRB file number** (1985-0122) on all documents or correspondence with the IRB concerning your research protocol.

→ Review and comply with all requirements on the enclosure, UMC Investigator Responsibilities, Protection of Human Research Participants.

Please note that the UMC IRB has the prerogative and authority to ask additional questions, request further information, require additional revisions, and monitor the conduct of your research and the consent process.

As a condition for publication of study results, the International Committee of Medical Journal Editors (ICMJE) requires all clinical research studies that begin enrolling participants on or after July 1, 2005 to be entered in a public registry before enrollment begins. For additional information please go to http://dor.umc.edu/proposalprep/policies/RegisteringClinicalTrialinPublicRegistry.doc.

We wish you the best as you conduct your research. If you have questions or need additional information, please contact the Human Research Office at (601) 984-2815.

Sincerely,

Gailen D. Marshall, Jr., M.D., Ph.D.
Chairman, Institutional Review Board 2

GDM/kc

Enclosure(s): (1) Investigator Responsibilities, Protection of Human Research Participants