seconded by Director John E. Bowman (Acting Director, Office of Thrift Supervision), concurred in by Director Thomas J. Curry (Appointive), and Chairman Sheila C. Bair, that Corporation business required its consideration of the matters which were to be the subject of this meeting on less than seven days' notice to the public; that no earlier notice of the meeting was practicable; that the public interest did not require consideration of the matters in a meeting open to public observation; and that the matters could be considered in a closed meeting by authority of subsections (c)(2), (c)(4), (c)(6), (c)(8), (c)(9)(A)(ii), (c)(9)(B), and (c)(10) of the "Government in the Sunshine Act" (5 U.S.C. 552b(c)(2), (c)(4), (c)(6), (c)(8), (c)(9)(A)(ii), (c)(9)(B), and (c)(10)). The meeting was held in the Board Room of the FDIC Building located at 550 - 7th Street, NW., Washington, DC.

Dated: February 17, 2010.

Federal Deposit Insurance Corporation.

By:

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2010–3391 Filed 2–17–10; 4:15 pm]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Secretary's Advisory Committee on Human Research Protections

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science.

ACTION: Notice.

SUMMARY: Pursuant to Section 10(a) of the Federal Advisory Committee Act, U.S.C. Appendix 2, notice is hereby given that the Secretary's Advisory Committee on Human Research Protections (SACHRP) will hold its twenty-second meeting. The meeting will be open to the public.

DATE: The meeting will be held on Tuesday, March 9, 2010 from 8:30 a.m. until 5 p.m. and Wednesday, March 10, 2010 from 8:30 a.m. until 5 p.m.

ADDRESSES: U.S. Department of Health & Human Services, 200 Independence Avenue, SW., Hubert H. Humphrey Building, Room 800, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Jerry Menikoff, M.D., J.D., Director, Office for Human Research Protections (OHRP), or Julia Gorey, J.D., Executive Director, SACHRP; U.S. Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852; 240–453–8141; fax: 240–453–6909; e-mail address: *sachrp@osophs.dhhs.gov*.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, SACHRP was established to provide expert advice and recommendations to the Secretary of Health and Human Services and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

On March 9, 2010, OHRP staff will provide a summary of public comments received on two recent draft guidance documents: Institutional Review Board (IRB) Continuing Review of Research and IRB Approval of Research with Conditions. Following this presentation, there will be a panel that will examine the context for resolution of regulatory harmonization issues through the Clinical Trials Transformation Initiative and the International Council on Harmonization and Good Clinical Practice. After lunch, the day will conclude with a report from the Subpart A Subcommittee (SAS) focusing on issues surrounding consent for future use of specimens or data. SAS is charged with developing recommendations for consideration by SACHRP about the application of subpart A of 45 CFR part 46 in the current research environment. This subcommittee was established by SACHRP at its October 2006 meeting.

On March 10, 2010, co-chairs of the Subcommittee on Harmonization (SOH) will discuss the charge, initial steps, and membership of this new group. The SOH was established by SACHRP at its July 2009 meeting and is charged with identifying and prioritizing areas in which regulations and/or guidelines for human subjects research adopted by various agencies or offices within HHS would benefit from harmonization, consistency, clarity, simplification, and/or coordination. The remainder of March 10 will be devoted to continuing the previous day's focus on the work of the Subpart A Subcommittee. Public comment will be heard on both days.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact persons. Members of the public will have the opportunity to provide comments on both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Executive Director, SACHRP, prior to the close of business Thursday, March 4, 2010. Information about SACHRP and the draft meeting agenda will be posted on the SACHRP Web site at: *http://www.hhs.gov/ohrp/sachrp/index.html*.

Dated: February 16, 2010.

Jerry Menikoff,

Director, Office for Human Research Protections, Executive Secretary, Secretary's Advisory Committee on Human Research Protections.

[FR Doc. 2010–3271 Filed 2–18–10; 8:45 am]

BILLING CODE 4150–36–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, e-mail *paperwork@hrsa.gov* or call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: The Health Education Assistance Loan (HEAL) Program Regulations (OMB No. 0915–0108) Extension

The Health Education Assistance Loan (HEAL) Program has regulations that contain notification, reporting and recordkeeping requirements to ensure that the lenders, holders and schools participating in the HEAL program follow sound management procedures in the administration of federally-insured student loans. While the regulatory requirements are approved under the OMB number referenced above, much of the burden associated with the submission of required HEAL forms and certain reporting requirements is approved under

separate OMB numbers. The table below provides the estimate of burden for the remaining regulations.

The estimates of burden are as follows:

REPORTING REQUIREMENTS

Number of respondents	Number of transactions	Total transactions	Hours per response (min)	Total burden hours
17 Holders	5	78	12	16
190 Schools	.4	78	10	13
Total Reporting				29

NOTIFICATION REQUIREMENTS

Number of respondents	Number of transactions	Total transactions	Hours per response (min)	Total burden hours
7,930 Borrowers	1	7,930	10	1,322
17 Holders	7,910	134,470	10	22,412
190 Schools	.89	170	14	40
Total Notification				23,774

RECORDKEEPING REQUIREMENTS

Number of respondents	Number of transactions	Total transactions	Hours per response (min)	Total burden hours
17 Holders	3,568	60,657	14	14,153
190 Schools	257	48,822	15	12,206
Total Recordkeeping				26,359
Total				50,162

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the "attention of the desk officer for HRSA."

Dated: February 5, 2010.

Sahira Rafiullah,

Director, Division of Policy Review and Coordination.

[FR Doc. 2010–3167 Filed 2–18–10; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; The Atherosclerosis Risk in Communities Study (ARIC)

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval the information collection listed below. This proposed information collection was previously published in the **Federal Register** on November 16, 2009, page 58962, and allowed 60 days for public comment. Only one comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

Proposed Collection

Title: The Atherosclerosis Risk in Communities Study (ARIC). *Type of Information Collection Request:* Extension of a currently approved collection (OMB No. 0925–0281). *Need and Use of Information Collection:* This project involves annual follow-up by telephone of participants in the ARIC study, review of their medical records, and interviews with doctors and family to identify disease occurrence. Interviewers will contact doctors and hospitals to ascertain participants' cardiovascular events. Information gathered will be used to further describe the risk factors, occurrence rates, and consequences of cardiovascular disease in middle aged and older men and women. *Frequency of Response:* The participants will be contacted annually. *Affected Public:* Individuals or households; Businesses or other for profit; Small businesses or organizations. *Type of Respondents:* Individuals or households; doctors and staff of hospitals and nursing homes. The annual reporting burden is as follows: *Estimated Number of Respondents:* 11,992; *Estimated Number of Responses per Respondent:* 1.0; *Average Burden Hours per Response:* 0.2399; and *Estimated Total Annual Burden Hours Requested:* 2,877.4. The annualized cost to respondents is estimated at $54,583, assuming respondents' time at the rate of $17.5