National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: Multi-Ethnic Study of Atherosclerosis (MESA) Event Surveillance. *Type of Information Request:* Renewal (OMB No. 0925–0493). *Need and Use of Information Collection:* The study, MESA, is identifying and quantifying factors associated with the presence and progression of subclinical cardiovascular disease (CVD)—that is, atherosclerosis and other forms of CVD that have not produced signs and symptoms. The findings provide important information on subclinical CVD in individuals of different ethnic backgrounds and provide information for studies on new interventions to prevent CVD. The aspects of the study that concern direct participant evaluation received a clinical exemption from OMB clearance (CE–99–11–08) in April 2000. OMB clearance is being sought for the contact of physicians and participant proxies to obtain information about clinical CVD events that participants experience during the follow-up period. *Frequency of response:* Once per CVD event. *Affected public:* Individuals. *Types of Respondents:* Physicians and selected proxies of individuals recruited for MESA. The annual reporting burden is as follows: *Estimated Number of Respondents:* 74; *Estimated Number of Responses per respondent:* 1.0; *Average Burden Hours Per Response:* 0.20; *and Estimated Total Annual Burden Hours Requested:* 14.7.

There are no capital, operating, or maintenance costs to report.

Type of respondents	Estimated number of respondents	Estimated number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
Physicians	17	1.0	0.20	3.4
Proxies	57	1.0	0.20	11.3
Total	74	1.0	0.20	14.7

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information will have practical utility; (2) The accuracy of the agency's estimate of burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

For Further Information: To request more information on the proposed project or to obtain a copy of data collection plans and instruments, contact Dr. Diane Bild, Division of Cardiovascular Sciences, NHLBI, NIH, II Rockledge Centre, 6701 Rockledge Drive, Suite 10122, MSC # 7936, Bethesda, MD 20892–7936, or call non-toll-free number (301) 435–0457, or e-mail your request, including your address to: *bildd@nhlbi.nih.gov.*

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Dated: July 27, 2010.

Suzanne Freeman,

NHLBI Project Clearance Liaison, National Institutes of Health.

Michael Lauer,

Director, DCVS, National Institutes of Health.

[FR Doc. 2010–19164 Filed 8–3–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Assessing the Long-Term Impacts of the John E. Fogarty International Center's Research and Training Programs

Summary: In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the John E. Fogarty International Center, the National Institutes of Health (NIH), will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection

Title: Assessing the Long-Term Impacts of the John E. Fogarty International Center's Research and Training Programs.

Type of Information Collection Request: New collection.

Need and Use of Information Collection: This study will inform investment decisions and strategies employed by the Fogarty International Center for the purpose of strengthening biomedical research capacity in low and middle income countries. The primary objective of the study is to develop detailed case studies of the long-term impacts of Fogarty's research and training programs on educational institutions located in low and middle income countries. The findings will provide valuable information concerning return on the Center's investments over the past twenty years and effective strategies for promoting research capacity development in the future.

Frequency of Response: Once.

Affected Public: Individuals.

Type of Respondents: Current and former NIH grantees; Current and former NIH trainees in countries of interest; Leaders and administrators at institutions of interest; Policy-makers and scientific leaders in countries of interest.

Estimated Number of Respondents: 105 per institution; total of 10 institutions over five years.

Estimated Number of Responses per Respondent: 1.

Average Burden Hours per Response: 1 hour for interview participants; 2 hours for focus group participants.

Estimated Total Annual Burden Hours Requested: 290, and the annualized cost to respondents is estimated at $4,841.

There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

	Number of respondents/ participants per institution	Number of institutions per year	Number of responses per respondent	Average burden hours per response	Estimated total annual burden hours requested
Interviews with US-based principal investigators	20	2	1	1	40
Focus groups with selected trainees and follow-on survey	40	2	1	2	160
Interviews with university leadership	4	2	1	1	8
Interviews with trainees	13	2	1	1	26
Interviews with foreign grantees	20	2	1	1	40
Interviews with foreign policy-makers/scientific leaders	8	2	1	1	16
Total	105				290

Request for Comments: Written comments and/or suggestions from the public and affected agencies are invited on one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (2) the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Regulatory Affairs, New Executive Office Building, Room 10235, Washington, DC 20503, Attention: Desk Officer for NIH. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Dr. Linda Kupfer, Fogarty International Center, National Institutes of Health, 16 Center Drive, Bethesda, MD 20892, or call non-toll-free number 301–496–3288, or e-mail your request, including your address to: *kupferl@mail.nih.gov.*

Comments Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30 days of the date of this publication.

Dated: July 22, 2010.

Timothy J. Tosten,

Executive Officer, John E. Fogarty International Center, National Institutes of Health.

[FR Doc. 2010–19160 Filed 8–3–10; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Secretary's Advisory Committee on Heritable Disorders in Newborns and Children; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), notice is hereby given of the following meeting:

Name: Secretary's Advisory Committee on Heritable Disorders in Newborns and Children.

Dates and Times: September 16, 2010, 8:30 a.m. to 5 p.m., September 17, 2010, 8:30 a.m. to 3:30 p.m.

Place: Marriott Washington at Metro Center, 775 12th Street, NW., Washington, DC 20005.

Status: The meeting will be open to the public with attendance limited to space availability. Participants are asked to register for the meeting by going to the registration Web site at *http://altarum.cvent.com/event/achdnc2010.* The registration deadline is Tuesday, September 14, 2010. Individuals who need special assistance, such as sign language interpretation or other reasonable accommodations should indicate their needs on the registration Web site. The deadline for special accommodation requests is Friday, September 10, 2010. If there are technical problems gaining access to the Web site, please contact Maureen Ball, Meetings Coordinator at *conferences@altarum.org.*

Purpose: The Secretary's Advisory Committee on Heritable Disorders in Newborns and Children (Advisory Committee) was established to advise and guide the Secretary regarding the most appropriate application of universal newborn screening tests, technologies, policies, guidelines and programs for effectively reducing morbidity and mortality in newborns and children having or at risk for heritable disorders. The Advisory Committee also provides advice and recommendations concerning the grants and projects authorized under the Public Health Service Act, 42 U.S.C. 300b–10, (Heritable Disorders Program) as amended in the Newborn Screening Saves Lives Act of 2008.

Agenda: The meeting will include: (1) A presentation of the External Review Workgroup's final report on the nomination of Critical Cyanotic Congenital Heart Disease and draft report on the nomination of Hyperbilirubinemia to the Advisory Committee's recommended uniform screening panel; (2) a discussion of the Advisory Committee's final draft of the report on the use and storage of newborn screening Residual Blood Spots; (3) an update on the report being developed by the Sickle Cell Disease Carrier Screening workgroup; and (4) presentations on the continued work and reports of the Advisory Committee's subcommittees on laboratory standards and procedures, follow-up and treatment, and education and training. Proposed Agenda items are subject to change as priorities dictate. You can locate the Agenda, Committee Roster and Charter, presentations, and meeting materials at the home page of the Advisory Committee's Web site at *http://www.hrsa.gov/heritabledisorderscommittee/.*

Public Comments: Members of the public can present oral comments during the public comment periods of the meeting, which are scheduled for both days of the meeting. Those individuals who want to make a comment are requested to register online by Tuesday, September 14, 2010 at *http://altarum.cvent.com/event/achdnc2010.* Requests will contain the name, address, telephone number, and any professional or business affiliation of the person desiring to make an oral presentation. Groups having similar interests are requested to combine their comments and present them through a single representative. The list of public comment participants will be posted on the Web site. Written comments should be e-mailed no later than Tuesday, September 14, 2010 for consideration. Comments should be submitted to Maureen Ball, Meetings