Form	Number of respondents	Responses per respondents	Total responses	Minutes per response	Total burden (in hours)	
EFN/FADHPS	80	1	80	10	13	

Send comments to Susan G. Queen, Ph.D., HRSA Reports Clearance Officer, Room 14–33, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857. Written comments should be received within 60 days of this notice.

Dated: November 7, 2000.

#### Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 00–29108 Filed 11–14–00; 8:45 am] **BILLING CODE 4160–15–U** 

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Health Resources and Services Administration

## Agency Information Collection Activities: Submission for OMB Review; 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, 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, 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: Social Support for Homeless Mothers: Implications for Best Practices and Program Design— New

The Health Care for the Homeless Clinicians' Network (HCHCN) of the National Health Care for the Homeless Council, Inc., through a cooperative agreement with the Bureau of Primary Health Care, Health Resources and Services Administration, proposes to

conduct a study on the social support available to homeless mothers, most of whom are parenting children alone. The study will be of adult homeless women and will be conducted by convening focus groups and administering a questionnaire to focus group members. The study is designed to look at clients' life events, histories of violence, medical and physical illness, social support, children's needs, and services use. The results will help to define best practices as they relate to social support processes and enable HCH programs to offer the appropriate mix of supports necessary to help mothers transition into permanent housing. The participants will be recruited from ten sites of the national Health Care for the Homeless program.

The estimated response burden is as follows:

Type of respondent	Number of respondents	Responses per respondent	Hours per response	Total hour bur- den
Focus Group (including survey)	100	1	1.5	150

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to: John Morrall, Human Resources and Housing Branch, Office of Management and Budget, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: November 7, 2000.

#### Jane M. Harrison,

Director, Division of Policy Review and Coordination.

[FR Doc. 00–29107 Filed 11–14–00;  $8:45~\mathrm{am}$ ] BILLING CODE 4160–15–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **National Institutes of Health**

National Heart, Lung, and Blood Institute; Proposed Collection; Comment Request, The Jackson Heart Study, Annual Follow-Up Component— Phase III

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 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: The Jackson Heart Study, Annual Follow-Up Component—Phase III. Type of Information Collection Request: Revision (OMB 0925–0464; expiration 04/30/2002). Need and use of

Information Collection: The Jackson Heart Study (JHS) Clinical Component will involve 6,500 African-American men and women aged 35-84. representative of African-American residents of Jackson, Mississippi. Family members are to be included in order to permit future studies of familial and genetic contributions to cardiovascular disease (CVD). The examination includes a series of questionnaires, physical assessments and laboratory measurements. Data collected in this study includes both conventional risk factors and new or emerging factors that may be related to CVD. Some of the newer areas of focus include early indicators of disease, genetics, sociocultural influences such as socioeconomic status and discrimination, and physiological relations between common disorders such as high blood pressure, obesity and diabetes and their influence on CVD and will take three years to complete. The JHS Clinical Component has received Clinical Exemption (CE-99-11-09) from the NIH Clinical Exemption Review

Committee. However, collection of follow information also involves third party individuals (next-of-kin decedents and physicians). This information is necessary for the interpretation and analysis of clinical findings and outcomes to ascertain the relationship between mortality and morbidity in the clinical study cohort. The information collected will be used by the public and

private sector for public health planning, medical education, other epidemiologic studies, and biomedical research. Frequency of Response: One-Time. Affected Public: Individuals or families; Businesses or other for profit; not-for-profit institutions. Type of Respondents: third party respondents (next-of-kin decedents and physicians). The annual reporting burden is as

follows: Estimated Number of Respondents: 480. Estimated Number of Responses per Respondent: 1. Average Burden Hours Per Response are shown in the table below; and Estimated Total Annual Burden Hours Requested: 160. The annualized cost to respondents is estimated at: \$3,600.

Estimates of the annual reporting burden to respondents.

Type of respondents	Estimated number of respondents	×	Estimated number of responses per respondent	×	Average bur- den hours per response	=	Estimated total annual burden hours re- quested
Morbidity and Mortality AFU 3rd party next-of-kin decedents	240		1		0.33		80
Morbidity and Mortality AFU 3rd party Physicians	240		1		0.33		80
Total	480						160

Note.—There are no Capital Costs, Operating Costs or Maintenance Cost for this study.

**REQUEST FOR COMMENTS: Written** comments and/or suggestions from the public and affected agencies should address one or more of the following points: (1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) evaluate 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) Enhance the quality, utility, and clarity of the information to be collected; and (4) 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.

FOR FURTHER INFORMATION: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Cheryl Nelson, Jackson Heart Study Project Officer, 6701 Rockledge Drive, Room 8152, MSC 7934, Rockville, MD 20892–7934, or call non-toll-free number (301) 435–0451 or E-mail your request, including your address to: cn80n@.nih.gov

**COMMENTS DUE DATE:** Comments regarding this information collection are best assured of having their full effect if received on or before January 16, 2001.

Dated: October 20, 2000.

#### Peter Savage,

Acting Director, Division of Epidemiology and Clinical Applications, National Heart, Lung, and Blood Institute.

[FR Doc. 00–29132 Filed 11–14–00; 8:45 am]

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **National Institute of Health**

# National Cancer Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel, Development of Novel Imaging Technologies. Date: December 6–7, 2000.

Time: 8:00 am to 5:00 pm.

Agenda: To review and evaluate grant applications.

*Place:* Hilton Gaithersburg, 620 Perry Parkway, Gaithersburg, MD 20877.

Contact Person: Timothy C. Meeker, MD, Scientific Review Administrator, Special Referral and Resources Branch, Division of Extramural Activities, National Cancer Institute, 6116 Executive Boulevard, Room 8088, Rockville, MD 20852, 301/594-1279. (Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: November 3, 2000.

### LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 00-29138 Filed 11-14-00; 8:45 am]

BILLING CODE 4140-01-M

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

#### **National Institutes of Health**

# National Cancer Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting.