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.