

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 those who are able 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 Veronica Chollette, RN, MS Program Director, Applied Cancer Screening Research Branch, Behavioral Research Program Division of Cancer Control and Population Sciences, National Cancer Institute, 6130 Executive Blvd., Room 4100, Rockville, MD 20852 or call non-toll free number 301-435-2837 or e-mail your request to: vc24a@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: May 9, 2005.

Rachelle Ragland-Greene,

NCI Project Clearance Liaison, National Institutes of Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Responsibility of Applicants for Promoting Objectivity in Research for Which Public Health Service Funding Is Sought and Responsible Prospective Contractors—42 CFR Part 50, Subpart F

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 Office of the Director (OD), 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: Responsibility of Applicants for Promoting Objectivity in Research for Which Public Health Service Funding is Sought and Responsible Prospective Contractors—42 CFR Part 50, Subpart F. *Type of Information Collection Request:* Revision of OMB No. 0925-0417, expiration date 09/31/2005. *Need and Use of Information Collections:* This is a request for OMB approval for the information collection and recordkeeping requirements contained in the final rule 42 CFR part 50, subpart F and Responsible Prospective Contractors: 45 CFR part 94. The purpose of the regulations is to promote objectivity in research by requiring institutions to establish standards which ensure that there is no reasonable expectation that the design, conduct, or reporting of research will be biased by a conflicting financial interest of an investigator. *Frequency of Response:* On occasion. *Affected Public:* Individuals or households; business or other for-profit; not-for-profit institutions; State local or tribal government. *Type of Respondents:* Any public or private entity or organization. The annual reporting burden is as follows: *Estimated Number of Respondents:* 42,800; *Estimated Number of Responses per Respondent:* 1.60; *Average Burden Hours per Response:* 3.40; and *Estimated Total Annual Burden Hours Requested:* 232,000. The annualized costs to respondents is estimated at: \$8,120,000. Operating costs and/or Maintenance costs are \$4,633.

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 function 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 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 Contact: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Mikia Currie, Assistant Project Clearance Officer, Office of Extramural Research (OER) Office of Policy for Extramural Research Administration (OPERA), 6705 Rockledge Drive, Room 1198, Bethesda, MD 20892-7974 or call non-toll-free number (301) 435-0941, e-mail your request including your address to: curriem@od.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: May 9, 2005.

Joe Ellis,

Acting Director, Office of Policy for Extramural Research Administration, National Institutes of Health.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, National Institutes of Health; Notice of 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 meeting of the Advisory Committee to the Director, National Institutes of Health (NIH).

The meeting will be open to the public as indicated below, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting.