

District of Columbia have developed, or are developing, a SACWIS with Federal financial participation.

45 CFR 1355.55 provides for continuing review, assessment and inspection of SACWIS. The purpose of this review is to determine whether the system, as described in the approved Advance Planning Document has been adequately completed and conforms to applicable regulations and policies.

To initiate a review, States complete and submit the SACWIS Assessment Review Guide (SARG) and other system documentation when they have

completed system development and the system is operational statewide. The SARG template provides a format for State description of system functionality, operation, and outputs such as reports. The additional materials submitted as part of this process, such as system design documentation, are typically readily available to the State as a result of good project management practices.

The information collected in the SACWIS Assessment Review Guide will allow Federal reviewers to determine if the State's SACWIS meets the

requirements for title IV–E Federal Financial Participation (FFP) defined at 45 CFR 1355.50, and that systems meet the goals and objectives of the approved Advance Planning Documents (APD) and conforms to the schedule, budget, and other conditions of their approved APDs. Additionally, other States may be able to use the documentation provided as part of their preparation for the review process of their own system development efforts.

Respondents: Title IV–E Agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
SACWIS Assessment Review Guide	3	1	250	750

Estimated Total Annual Burden Hours: 750.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection. *E-mail address:* infocollection@acf.hhs.gov.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–7285, *E-mail:* oir_submission@omb.eop.gov, Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2011–11995 Filed 5–16–11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; NCI Cancer Genetics Services Directory Web-Based Application Form and Update Mailer

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Cancer Institute (NCI), and the National Institutes of Health (NIH), have submitted to the Office of Management and Budget (OMB) a request to review and approve the information collection listed below. This proposed information collection was previously published in the **Federal Register** on March 15, 2011 (76 FR 14034) and allowed 60-days for public comment. No public comments were 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: NCI Cancer Genetics Services Directory Web-based Application Form and Update Mailer. *Type of Information Collection Request:* Existing Collection in Use Without an OMB Number. *Need and Use of Information Collection:* The purpose of the online application form and the Web-based update mailer is to collect information about genetics professionals to be included in the NCI Cancer Genetics Services Directory on NCI's Cancer.gov Web site. The information collected includes name, practice locations, professional qualifications, and areas of specialization. *Frequency of Response:* Information is collected once via the online application form, and then updated annually via the Web-based mailer. *Affected Public:* Individuals. *Type of Respondents:* Genetics professionals including nurses, physicians, genetic counselors, and other professionals who provide services related to cancer genetics. The annual reporting burden is estimated at 180 hours (see Table below). There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

TABLE 1—ESTIMATES OF ANNUAL BURDEN HOURS

Type of respondents	Tool	Number of respondents	Frequency of response	Average time per response minutes/hour (hours)	Annual burden hours
Genetics Professionals	Application Form	60	1	30/60 (.50)	30
	Web-based Update Mailer	600	1	15/60 (0.25)	150

TABLE 1—ESTIMATES OF ANNUAL BURDEN HOURS—Continued

Type of respondents	Tool	Number of respondents	Frequency of response	Average time per response minutes/hour (hours)	Annual burden hours
Totals	660	180

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.

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 Attention: NIH Desk Officer, Office of Management and Budget, at OIRA_submission@omb.eop.gov or by fax to 202-395-6974. To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Margaret Beckwith, Acting Branch Chief, International Cancer Research Databank Branch, Office of Cancer Content Management, Office of Communication and Education, National Cancer Institute, 6116 Executive Blvd., Rockville, MD 20852, or call non-toll-free number 301-496-9096 or e-mail your request, including your address to: mbeckwit@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: May 11, 2011.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2011-12047 Filed 5-16-11; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Arthritis and Musculoskeletal and Skin Diseases; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. App.), 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: Arthritis and Musculoskeletal and Skin Diseases Initial Review Group, Arthritis and Musculoskeletal and Skin Diseases Special Grants Review Committee.

Date: June 15–16, 2011.

Time: 7:30 p.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Hilton Washington/Rockville, 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Helen Lin, PhD, Scientific Review Officer, Scientific Review Branch, National Institute of Arthritis, Musculoskeletal and Skin Diseases, National Institutes of Health, 6701 Democracy Blvd., Suite 800, Bethesda, MD 20817, 301-594-4952, linh1@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.846, Arthritis, Musculoskeletal and Skin Diseases Research, National Institutes of Health, HHS)

Dated: May 10, 2011.

Jennifer S. Spaeth,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2011-12099 Filed 5-16-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; Notice of Closed Meetings

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

The meetings 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 materials, 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 Institute of Neurological Disorders and Stroke Special Emphasis Panel; Headache Clinical Trial.

Date: June 8, 2011.

Time: 9 a.m. to 11:30 a.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Richard D. Crosland, PhD, Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS/NIH/DHHS/Neuroscience Center, 6001 Executive Blvd., Suite 3208, MSC 9529, Bethesda, MD 20892-9529, 301-594-0635, Rc218u@nih.gov.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; Stroke Clinical Trial.

Date: June 17, 2011.

Time: 9 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Telephone Conference Call).

Contact Person: Richard D. Crosland, PhD, Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS/NIH/DHHS/Neuroscience Center, 6001 Executive Blvd., Suite 3208, MSC 9529, Bethesda, MD 20892-9529, 301-594-0635, Rc218u@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research