DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; Comment Request; Pretesting of NIAID's Biomedical HIV Prevention Research Communication Messages

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute of Allergy and Infectious Diseases (NIAID), the National Institutes of Health (NIH) has 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 November 17, 2010 (Volume 75, Number 221), page 70270–70271 and allowed 60-days for public comment. In response, NIAID received two requests for copies of the clearance package, which were provided. No additional requests, comments or suggestions 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: Pretesting of NIAID's Biomedical HIV Prevention Research Communication Messages. *Type of Information Collection Request:* Revision of a previously approved collection. *Need and Use of Information Collection:* This is a request for clearance to pretest messages, materials and program activities about biomedical HIV prevention research. The primary objectives of the pretests are to (1) assess audience knowledge, attitudes, behaviors and other characteristics for the planning/development of health messages, education products, communication strategies, and public information programs; and (2) pretest these health messages, products, strategies, and program components while they are in developmental form to assess audience comprehension, reactions, and perceptions. The information obtained from audience research and pretesting results in more effective messages, materials, and programmatic strategies. By maximizing the effectiveness of these messages and strategies for reaching targeted audiences, the frequency with which publications, products, and programs need to be modified is reduced. *Frequency of Response:* On occasion. *Affected Public:* Individuals. *Type of Respondents:* Adults at risk for HIV/AIDS; healthcare providers; representatives of organizations disseminating HIV-related messages or materials. The total reporting burden over the 3-year period is shown in the table below. There are no Capital Costs to report. There are no Operating or Maintenance Costs to report.

Note: The burden table below reflects what NIAID anticipates would be accomplished over the total 3-year life of the clearance. (Annual burden, therefore, is one-third of the total figures presented here.)

TABLE 1—ESTIMATES OF HOUR BURDEN BY ANTICIPATED DATA COLLECTION METHODS

	Total number of respondents	Frequency of response	Hours per response	Total hours
Individual In-Depth Interviews (in person or telephone)	228	1	1	228
	22 (Partners/Stakeholders)	2	1	44
Focus Group Interviews	864	1	2	1728
Intercept Interviews/Surveys	4500	1	.25	1125
Gatekeeper Reviews	150	1	.25	37.5
Self-Administered Questionnaires: Random selection from central location, online, etc.	1500	1	.25	375
Self-Administered Customer Satisfaction Surveys of Meetings and Conference Sessions.	2265	1	.2	453
	50 (Partners/Stakeholders)	3	.2	30
Self-Administered: Customer Satisfaction Surveys of Materials and Services.	50 (Partners)	3	.25	37.5
Self-Administered Customer Satisfaction Pop-up Surveys	900	1	.08	72
Telephone Surveys	1000	1	.25	250
Totals	11,529			4,380

(**Note:** On an annual basis, the total number of respondents is 3,843; and the total annual hours are 1,460)

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: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, 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 Katharine Kripke, Assistant Director, Vaccine Research Program, Division of AIDS, NIAID, NIH, 6700B Rockledge Dr., Bethesda, MD 20892–7628, or call non-toll-free number 301–402–0846, or e-mail your request, including your address to *kripkek@niaid.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: January 28, 2011.

William A. Gillen,

Acting Deputy Director for Science Management, National Institute of Allergy and Infectious Diseases, NIH/HHS.

[FR Doc. 2011–2546 Filed 2–3–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; Comment Request; Short Follow-Up Questionnaire for the National Institutes of Health (NIH)–AARP Diet and Health Study (NCI)

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 Cancer Institute (NCI), 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: Short Follow-Up Questionnaire for the National Institutes of Health (NIH)–AARP Diet and Health Study (NCI). *Type of Information Collection Request:* Extension. *Need and Use of Information Collection:* The purpose of this short 2-page questionnaire is to obtain information on 18 different medical conditions, several medical procedures, and lifestyle characteristics from 485,909 participants of the NIH–AARP Diet and Health Study. The questionnaire will support the ongoing examination between cancer and nutritional exposures. A pilot mailing to 1,600 randomly selected NIH–AARP Diet and Health study participants confirmed the feasibility of the methodology and willingness of respondents to participate in this data collection effort. This questionnaire adheres to The Public Health Service Act, Section 412 (42 U.S.C. 285a–1) and Section 413 (42 U.S.C. 285a–2), which authorizes the Division of Cancer Epidemiology and Genetics of the National Cancer Institute (NCI) to establish and support programs for the detection, diagnosis, prevention and treatment of cancer; and to collect, identify, analyze and disseminate information on cancer research, diagnosis, prevention and treatment. *Frequency of Response:* Once. *Affected Public:* Individuals. *Type of Respondents:* U.S. adults (persons aged 50–85). The annual reporting burden is displayed in the table below. There are no Capital Costs, Operating Costs, and/or Maintenance Costs to report.

Type of respondents	Number of respondents	Frequency of response	Average time per response (Minutes/Hour)	Annual hour burden
Senior Adults	485,909	1	4/60 (0.067)	32,394

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 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 CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact Yikyung Park, Sc.D., Staff Scientist, Nutritional Epidemiology Branch, Division of Cancer Epidemiology and Genetics, National Cancer Institute, NIH, DHHS, 6120 Executive Blvd., Rockville, MD 20852 or call non-toll-free number 301–594–6394 or e-mail your request, including your address to: *parkyik@mail.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: January 28, 2011.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2011–2540 Filed 2–3–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; 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 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 Initial Review Group; NST–2 Subcommittee.

Date: March 7–8, 2011.

Time: 8 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Willard InterContinental Washington, 1401 Pennsylvania Avenue, NW., Washington, DC 20004.

Contact Person: JoAnn McConnell, PhD, Scientific Review Officer, Scientific Review Branch, NINDS/NIH/DHHS, Neuroscience Center, 6001 Executive Blvd., Suite 3208, MSC 9529, Bethesda, MD 20892–9529, 301–496–5324, *mcconnej@ninds.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS)

Dated: January 31, 2011.

Anna P. Snouffer,

Deputy Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2011–2500 Filed 2–3–11; 8:45 am]

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