order to perform the functions related to hematopoietic stem cell transplantation under contract to the U.S. Department of Health and Human Services (HHS). The Act requires the Secretary to contract for the establishment and maintenance of information related to patients who have received stem cell therapeutic products and to do so using a standardized, electronic format. Data is collected from transplant centers by the Center for International Blood and Marrow Transplant Research and is used for ongoing analysis of transplant outcomes. HRSA uses the information in order to carry out its statutory responsibilities. Information is needed to monitor the clinical status of transplantation and to provide the Secretary of HHS with an annual report of transplant center-specific survival data. The increase in burden, as reflected in this revised submission request, is due to an increase in the annual number of transplants and increasing survivorship after transplantation.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form name	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Baseline Pre-TED (Transplant Essential Data)	200	38	7,600	1	7,600
Product Form (includes Infusion, HLA, and Infectious Disease Marker inserts)	200	29	5,800	1	5,800
100-Day Post-TED	200	38	7,600	0.85	6,460
6-Month Post-TED	200	31	6,200	1	6,200
12-Month Post-TED	200	27	5,400	1	5,400
Annual Post-TED	200	104	20,800	1	20,800
Total	200		53,400		52,260

Dated: June 5, 2013.

Bahar Niakan,

Director, Division of Policy and Information Coordination.

[FR Doc. 2013–13790 Filed 6–10–13; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-day Comment Request; Web-Based Media Literacy Parent Training for Substance Use Prevention in Rural Locations

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute on Drug Abuse, National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below. This proposed information collection was previously published in the **Federal Register** on March 27, 2013, pages 18612–18613 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 Institute on Drug Abuse (NIDA), 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.

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: NIH Desk Officer.

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.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project contact: Dr. Augie Diana, Health Scientist Administrator, Prevention Research Branch, Division of Epidemiology, Services, and Prevention Research, NIDA, NIH, 6001 Executive Boulevard, Room 5163, Bethesda, MD 20892, or call non-toll-free number (301) 443–1942 or Email your request, including your address to: *dianaa@nida.nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Web-based Media Literacy Parent Training for Substance Use Prevention in Rural Locations, 0925-New, National Institute on Drug Abuse (NIDA), National Institutes of Health (NIH).

Need and Use of Information Collection: This study will develop a web-based media literacy substance use prevention intervention for use with parents and their elementary school children (approximately ages 7–12), and will evaluate the program in a randomized controlled trial to establish program efficacy in six rural communities in North Carolina and Texas. The primary objectives of the study are to assess the efficacy of a media literacy education program that is specifically designed to overcome barriers to prevention efforts in rural communities, and to provide the scientific basis for establishing the program, Media Detective Family, as an evidence-based substance use prevention curriculum. The findings will provide valuable information concerning: (1) The appropriateness of using technology for substance use prevention programming (i.e., internet, Smartphone, or tablet-based applications) to reach rural families with elementary school-aged children;

(2) improvements in parents' and children's critical thinking skills associated with intervention exposure; (3) improvements in parent-child communication about substances and the media associated with intervention exposure; and (4) reductions in children's behavioral intentions to use substances associated with intervention exposure.

OMB approval is requested for two years. There are no costs to respondents other than their time. There are no capital, operating, and/or maintenance costs. The total estimated annualized burden hours are 1067.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Number of respondents	Frequency of response	Average time per response	Annual hour burden
Adults:				
Permission & Consent	200	1	10/60	33.33
Pretest		1	50/60	166.67
Posttest		1	45/60	150.00
Follow-up		1	45/60	150.00
Usage Log		2	10/60	67.00
Children:				
Assent	200	1	10/60	33.33
Pretest		1	50/60	166.67
Posttest		1	45/60	150.00
Follow-up		1	45/60	150.00

Dated: June 5, 2013.

Glenda J. Conroy,

Executive Officer, (OM Director), National Institute on Drug Abuse, NIH.

[FR Doc. 2013–13795 Filed 6–10–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; 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 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 Institute on Aging Special Emphasis Panel SWAN.

Date: June 27, 2013.

Time: 9:30 a.m. to 12:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Aging, Gateway Building, Suite 2C212, 7201 Wisconsin Avenue, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Rebecca Jo Ferrell, Ph.D., Scientific Review Branch, National Institute on Aging, Gateway Building, 7201 Wisconsin Avenue, Suite 2C212, Bethesda, MD 20892, 301–402–7703, *rebecca.ferrell@nih.gov.*

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

Name of Committee: National Institute on Aging Special Emphasis Panel Member Conflict.

Date: July 29, 2013.

Time: 10:00 a.m. to 11:30 a.m.

Agenda: To review and evaluate grant applications.

Place: National Institute on Aging, Gateway Building, Suite 2C212, 7201 Wisconsin Avenue, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Ramesh Vemuri, Ph.D., Scientific Review Branch, National Institute on Aging, Gateway Building, 7201 Wisconsin Avenue, Suite 2C212, Bethesda, MD 20892, 301–402–7700, *rv23r@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS)

Dated: June 4, 2013.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–13620 Filed 6–10–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Biomedical Imaging and Bioengineering; 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 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 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 Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel; NIBIB K Training Meeting.

Date: July 15, 2013.

Time: 12:30 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, DEM II, 6707 Democracy Blvd., Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Ruth Grossman, DDS, Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, 6707 Democracy Boulevard, Room 960, Bethesda, MD 20892, 301–496–8775, *grossmanrs@mail.nih.gov.*

Dated: June 5, 2013.

Melanie J. Gray,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–13743 Filed 6–10–13; 8:45 am]

BILLING CODE 4140–01–P