formulate improvement plans and take action when necessary.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 2,485.

Estimated Annualized Burden Hours

QUANTITATIVE SURVEY

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Science professionals, applicants, reviewers, Institutional Officials	3,820	1	15/60	955
Adult Science Trainees	2,000	1	15/60	500
General Public	4,000	1	15/60	1,000

QUALITATIVE SURVEY

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Science professionals, applicants, reviewers, Institutional Officials	12	1	1	12
Adult Science Trainees	6	1	1	6
General Public	12	1	1	12

Dated: November 13, 2013.

Seleda Perryman,

Chief, Project Clearance Officer, Office of Policy for Extramural Research Administration, National Institutes of Health.

[FR Doc. 2013–27965 Filed 11–21–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases; 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 of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; NIDDK Career Awards Review.

Date: December 4, 2013.

Time: 4:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Carol J. Goter-Robinson, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 748, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, (301) 594–7791, *goterrobinsonc@extra.niddk.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 of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Time-Sensitive Obesity.

Date: December 11, 2013.

Time: 3:30 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Michele L. Barnard, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 753, 6707 Democracy Boulevard, Bethesda, MD 20892–2542, (301) 594–8898, *barnardm@extra.niddk.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: November 18, 2013.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–28003 Filed 11–21–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; 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 a meeting of the Board of Scientific Counselors, National Human Genome Research Institute.

The meeting will be closed to the public as indicated below in accordance with the provisions set forth in section 552b(c)(6), Title 5 U.S.C., as amended for the review, discussion, and evaluation of individual intramural programs and projects conducted by the NATIONAL HUMAN GENOME RESEARCH INSTITUTE, including consideration of personnel qualifications and performance, and the competence of individual investigators, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Board of Scientific Counselors, National Human Genome Research Institute.

Date: December 11–13, 2013.

Time: 6:00 p.m. to 1:00 p.m.

Agenda: To review and evaluate personal qualifications and performance, and competence of individual investigators.

Place: National Institutes of Health, Building 45, Room D, 9000 Rockville Pike, Rockville, MD 20892.

Contact Person: Monica Berger, Executive Secretary, Office of the Scientific Director, National Human Genome Research Institute, 50 South Drive, Bldg. 50, Rm. 5222,

Bethesda, MD 20892, 301–294–6873, *bergerm@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.172, Human Genome Research, National Institutes of Health, HHS)

Dated: November 18, 2013.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2013–28005 Filed 11–21–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-day Comment Request: Generic Clearance To Support Programs and Administrative Operations at the National Cancer Institute (NCI)

SUMMARY: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the 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 May 23, 2013, Volume 78, p. 30930 and allowed 60-days for public comment. One public comment was received on May 24, 2013 stating that the agency should spend more money on funding prevention research. An email response was sent on May 28, 2013 stating, ''Your comments were received and they will be taken into consideration.'' The purpose of this notice is to allow an additional 30 days for public comment. The National Cancer Institute (NCI), 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.

DATES: *Comment 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: Vivian Horovitch-Kelley, PRA/OMB Project Clearance Liaison, Office of Management Policy and Compliance (OMPC), National Cancer Institute, 11400 Rockville Pike, Room 707, Rockville, MD 20852 or call non-toll-free number 301–480–0541 or Email your request, including your address to: *Horovitchkellv@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Generic Clearance to Support Programs and Administrative Operations At the National Cancer Institute (NCI), NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: This is a request for a generic submission that would be used for administrative and program-related submissions. Administrative submissions are defined as information collections (ICs) wherein the primary content is used for administrative purposes (e.g., an application) or to monitor, measure, manage or improve a program. These ICs may involve little if any, subsequent analysis and/or the use of descriptive statistics. Some ICs are forms used to source and aggregate contact information, history, preferences, opinions, and/or other data that does not necessitate further inquiry but allow the respondents to maintain contact, indicate preferences, and respond to data calls of information that has not already been collected. Other ICs may be program-related requests for the purpose of program monitoring, performance measurement, and improving or assessing the effectiveness of the program. This submission is the result of a year worth of analysis at the National Cancer Institute (NCI) which has demonstrated that more often than not, the potential and actual Paperwork Reduction Act (PRA) bootlegs that occur are administrative in nature, not research based. Additionally, NCI program staff who have submitted sub-projects that have been reviewed and returned by OMB, have contributed ideas and comments to this request. And finally, input and collaborations have been sought regarding this submission with program staff from different divisions and offices at NCI and PRA Liaisions at a variety of other National Institutes of Health (NIH) Institutes. Along with the analysis, NCI's ongoing education and outreach effort has increased the awareness and the need for a generic submission that covers administrative and program-related information collections. NCI's current scope for administrative generic sub-projects is non-existent and this submission would fill that gap. Subsequently to publishing the 60-day **Federal Register** Notice for this project, the program staff realized that the need was understated and thus increased the requested burden hours from 6,000 hours to 16,667 hours over the three-year information collection period.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 16,667.

ESTIMATED BURDEN HOURS OVER THREE YEARS

Category of respondents	Number of respondents	Frequency of responses	Average time per response (in hours)	Total burden hours
Individuals, Households, Private Sector, State Government, Local Government, Tribal Government, or Federal Government	20,000	1	50/60	16,667