OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 822.

ESTIMATES OF ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Biobanks (Private Sector)	548	1	90/60	822

Dated: July 17, 2013.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2013–17642 Filed 7–22–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request: Application for the Postdoctoral Research Associate Program

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 Institute of General Medical Sciences (NIGMS), 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.

Written comments and/or suggestions from the public and affected agencies are invited on 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.

To Submit Comments and for Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing or request more information on the proposed project, contact Ms. Tammy Dean-Maxwell, NIGMS, NIH, Natcher Building, Room 3AN.44, 45 Center Drive, MSC 6200, Bethesda, MD 20892–6200, or call non-toll-free number 301–594–2755 or email your request, including your address to: *deanmat@mail.nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

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.

Proposed Collection: Application for the Postdoctoral Research Associate Program, (NIGMS) Extension of a currently approved collection, OMB No. 0925–0378, expiration date September 30, 2013. *Form Numbers:* NIH 2721–1, NIH 2721–2.

Need and Use of Information Collection: The Postdoctoral Research Associate (PRAT) Program will use the applicant and referee information to award opportunities for training and experience in laboratory or clinical investigation to individuals with an appropriate terminal degree who are seeking training in an NIGMS designated emerging area of science, through appointments as PRAT Fellows at the National Institutes of Health or the Food and Drug Administration. The goal of the program is to develop leaders in designated emerging areas of biomedical research for key positions in academic, industrial, and Federal research laboratories. *Frequency of Response:* Once a year. *Affected Public:* Individuals or households; businesses or other for-profit. *Type of Respondents:* Applicants and referees.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 331.

ESTIMATED ANNUALIZED BURDEN TABLE

Form name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
PRAT Primary Application (NIH 2721–1)	Applicants	25	1	8	200
PRAT Request for Evaluation Form (NIH 2721–2)	Referee	75	1	105/60	131

Dated: July 17, 2013.

Sally Lee,

Executive Officer, National Institute of General Medical Sciences, National Institutes of Health.

[FR Doc. 2013–17645 Filed 7–22–13; 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: National Cancer Institute (NCI) Cancer Nanotechnology Platform Partnership Scientific Progress Reports

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 13, Vol. 78, p. 27974 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 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.

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: To obtain a copy of the data collection plans and instruments or request more information on the proposed project contact: Dorothy Farrell, Center for Strategic Scientific Initiatives, Office of Cancer Nanotechnology Research, National Cancer Institute, 31 Center Drive, Bldg. 31 A, Rm. 10A52, Bethesda, MD 20892 or call non-toll-free number 301–496–5652 or Email your request, including your address to: *farrelld@mail.nih.gov*. Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: National Cancer Institute (NCI) Alliance for Nanotechnology in Cancer Platform Partnership Scientific Progress Reports, 0925–NEW, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: National Institutes of Health grantees are required to submit interim and final progress reports and other post-award documents associated with the monitoring, oversight, and closeout of an award. This submission represents a request for OMB to approve new program specific progress report guidelines for Cancer Nanotechnology Platform Partnerships (CNPP) awarded by the National Cancer Institute (NCI). The CNPPs are part of the Alliance for Nanotechnology in Cancer, a network of awards funded by NCI to promote the application of nanotechnology to cancer research and care. The proposed guidelines request information about award performance related to trans-Alliance collaboration, scientific milestones, progress towards clinical translation and technology commercialization, and education and outreach efforts. The report also gathers information on leveraged funding, patents and publications. The information is gathered every six months. This information is needed to monitor the performance of this special program within NCI, funded through Requests for Applications (RFA CA–09–013, released May 29, 2009) using the cooperative agreement mechanism (U01). The information will be used to monitor individual award performance and the effectiveness of the program as a whole. The respondents are the Principal Investigators of the awards, along with their institutional business officials. The awards are administered by and the reports reviewed by the Office of Cancer Nanotechnology Research (OCNR), part of the Center for Strategic Scientific Initiatives within NCI.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The estimated annualized burden hours are 72.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Principal Investigators	12	2	3	72

Dated: July 15, 2013.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2013–17644 Filed 7–22–13; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS–2013–0016]

Agency Information Collection Activities: Office of Biometric Identity Management (OBIM) Biometric Data Collection at the Ports of Entry

AGENCY: National Protection and Programs Directorate, DHS.

ACTION: 30-Day notice and request for comments; Extension, without change, of a currently approved collection.

SUMMARY: The Department of Homeland Security (DHS), National Protection and Programs Directorate (NPPD), Office of Biometric Identity Management (OBIM), formerly the United States Visitor and Immigrant Status Indicator Technology (US–VISIT) Program, will submit the following Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chapter 35). NPPD is soliciting comments