information, please include the information request collection title for reference.

Information Collection Request Title: Faculty Loan Repayment Program OMB No. 0915–0150, Revision.

Abstract: Under the HRSA Faculty Loan Repayment Program, degree-trained health professionals from disadvantaged health backgrounds may enter into a contract under which the Department of Health and Human Services will make payments on eligible educational loans in exchange for a minimum of 2 years of service as a full-time or part-time faculty member of an accredited health professions college or university.

Need and Proposed Use of the Information: The Faculty Loan Repayment Program needs to collect data to determine an applicant's eligibility for the program. Information is collected from the applicants and/or the educational institutions which includes general applicant data, applicant educational loan history, employment status, and information regarding the educational institution which employs the applicant.

Likely Respondents: Faculty Loan Repayment Program applicants and institutions providing employment to the applicants.

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 Information Collection Request are summarized in the table below.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Eligible Applications	111	1	111	1	111
Institution/Loan Repayment Employment Form	*111	*1	111	1	111
Authorization to Release Information Form	111	1	111	.25	27.83
Total	222				249.83

*Respondent for this form is the institution for the applicant.

HRSA specifically requests comments on (1) the necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Jackie Painter,

Director, Division of the Executive Secretariat.

[FR Doc. 2015–07577 Filed 4–1–15; 8:45 am]

BILLING CODE CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request Electronic Prior Approval Submission System (ePASS) (NHLBI)

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 November 24, 2014 (79 FR 69865), and allowed 60-days for public comment. One public comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Heart, Lung and Blood Institute (NHLBI), 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: Ms. Ryan Lombardi, 6701 Rockledge, Office of Grants Management, National Heart, Lung, and Blood Institute, National Institutes of Health, 6701 Rockledge Dr, MSC 7926, Bethesda, MD 20892–7926, or call non-toll-free number 301–435–0166, or Email your request, including your address to *lombardr@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Electronic Prior Approval Submission System (ePASS), 0925—New, National Heart Lung and Blood Institute (NHLBI), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose and use of the information collection for this project is to collect and track certain requests (such as budget modifications or undertaking particular activities) from NIH grantees in an electronic format. This new electronic system, ePASS (electronic Prior Approval Submission System), will enable grantees to have a standard way to submit requests for their projects per NIH policy. The grantee will initiate a request for a certain action as required by NIH policy: Use of unobligated balances/carryover, change of PI, change of effort, Training

Grant (NRSA) waivers, significant rebudgeting, 2nd and 3rd no cost extensions, and change of scope. These are all prior approvals as required by the NIH Grants Policy, and need to be reviewed and approved by the NHLBI. ePASS will provide a template to ensure that all specific points are addressed and documented in the official grant file. All information is submitted via the internet, tracked in ePASS, and the documentation will automatically be forwarded to the official grant file. The system will ensure that individuals authorized by the grantee are submitting requests and that the appropriate NIH staff is receiving the requests. The requests will be template driven so that the grantee is including the minimally required information, thus eliminating the usual back and forth to obtain missing information. Forms will have automatic fill-in capability that will reduce typos in grant numbers and PI names, further reducing approval time. Reminders will be sent to NIH staff within ePASS based on roles to ensure timely responses to the grantee. The system will facilitate email communication with applicants by automatic notifications when applications are received and when NIH has made a determination regarding a request (approval issued or request denied with explanation for denial).

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 470.

ESTIMATES OF HOUR BURDEN

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual hour burden
NHLBI Grantees	940	1	30/60	470
Totals	940			470

Dated: March 17, 2015.

Lynn Susulske,

NHLBI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2015–07623 Filed 4–1–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee on Procedures Review, Advisory Board on Radiation and Worker Health (ABRWH or Advisory Board), National Institute for Occupational Safety and Health (NIOSH)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting for the aforementioned subcommittee:

Time and Date: 11:00 a.m.–5:00 p.m., Eastern Standard Time, April 28, 2015.

Place: Audio Conference Call via FTS Conferencing.

Status: Open to the public. The public is welcome to submit written comments in advance of the meeting, to the contact person below. Written comments received in advance of the meeting will be included in the official record of the meeting. The public is also welcome to listen to the meeting by joining the teleconference at the USA toll-free, dial-in number, 1–866–659–0537 and the passcode is 9933701.

Background: The ABRWH was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the compensation program. Key functions of the ABRWH include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort (SEC).

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2015.

Purpose: The ABRWH is charged with (a) providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, providing advice to the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is a reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee on Procedures Review was established to aid the ABRWH in carrying out its duty to advise the Secretary, HHS, on dose reconstructions. The Subcommittee on Procedures Review is responsible for overseeing, tracking, and participating in the reviews of all procedures used in the dose reconstruction process by the NIOSH Division of Compensation Analysis and Support (DCAS) and its dose reconstruction contractor (Oak Ridge Associated Universities—ORAU).

Matters for Discussion: The agenda for the Subcommittee meeting includes: discussion of procedures in the following ORAU and DCAS technical documents: Procedures for reconstructing dose associated with potential skin contamination, ORAU Team Technical Information Bulletin (OTIB) 0034 (''Internal Dose Coworker Data for X–10''), OTIB 0054 (''Fission and Activation Product Assignment for Internal Dose-Related Gross Beta and Gross Gamma Analyses''), OTIB 0082 (''Dose Reconstruction Method for Chronic Lymphocytic Leukemia''), Update on Review of ORAU Team Report 0053 (''Stratified Co-Worker Sets''); and a continuation of the comment-resolution process for other dose reconstruction procedures under review by the Subcommittee.

The agenda is subject to change as priorities dictate.