Further information regarding the Webcast, including the Web address for the Webcast, will be made available at least 2 days in advance of the meeting at the following Web site: *http://www.fda.gov/AdvisoryCommittees/CommitteesMeetingMaterials/MedicalDevices/MedicalDevicesAdvisoryCommittee/Gastroenterology-UrologyDevicesPanel/default.htm.* Select the link for 2015 Meeting Materials.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before April 30, 2015. Oral presentations from the public will be scheduled between approximately 1 p.m. and 2 p.m. on May 14 and between approximately 9 a.m. and 10 a.m. on May 15. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before April 15, 2015. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by April 20, 2015.

FDA is opening a docket for public comment on this document. The Docket No. is FDA–2015–N–0722. The docket will close on May 28, 2015. Interested persons are encouraged to use the docket to submit electronic or written comments regarding this meeting. Comments received on or before April 30, 2015, will be provided to the committee. Comments received after that date will be taken into consideration by the Agency. Submit electronic comments to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management, Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852. Submit a single copy of electronic comments or two paper copies of any mailed comments. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Divisions of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

For press inquiries, please contact the Office of Media Affairs at *fdaoma@fda.hhs.gov* or 301–796–4540.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, at *AnnMarie.Williams@fda.hhs.gov* or 301–796–5966 at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at *http://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: March 9, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–05710 Filed 3–12–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60 Day Comment Request; Assessment of NHLBI's Global Health Initiative Collaborating Centers of Excellence (NHLBI)

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 Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH), will publish periodic summaries of proposed projects 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) 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.

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: Deshiree Belis, 6705 Rockledge Drive, Suite 6070, Bethesda, MD 20892, or call non-toll-free number (301)-435–1032, or Email your request to: *deshiree.belis@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: Assessment of NHLBI's Global Health Initiative Collaborating Centers of Excellence, 0925-New, National Heart, Lung, and Blood Institute (NHLBI), the National Institutes of Health (NIH).

Need and Use of Information Collection: This collection proposes to conduct a one-time outcome evaluation of the NHLBI Global Health Initiative Centers of Excellence (GHI COE) Program to examine the extent to which the program achieved its intended objectives in developing sustainable research and research training capacity, and advancing information about the prevention and treatment of chronic non-communicable chronic cardiovascular and pulmonary diseases (CVPD) in low- and middle-income country (LMIC) populations. The outcome evaluation will utilize a mixed-methods approach to comprehend each COE's processes, short term outcomes, and sustainability outcomes/efforts. Specifically, the evaluation will involve triangulating quantitative data sources (*e.g.,* archived systematic reporting data), and qualitative data sources (*e.g.,* archival data and key informant interview data). Data collected will be used to develop a Case Study report for each COE outlining their experience with implementing their program as well as a comprehensive cross-site Lessons Learned Report describing knowledge and experiences from the overall program, including similarities and differences across a variety of

project settings and conditions. Findings from interviews will be incorporated into the Case Studies report and Lessons Learned report, which will be used by CTRIS to inform NHLBI and NIH stakeholders about structural issues relevant to planning both global and domestic biomedical research and training programs with diverse operational conditions and challenges. Additionally, COEs may utilize the Case Studies report as a marketing tool to attract additional funding and media coverage.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 36.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Principal Investigators	9	1	1	9
Training Directors	9	1	1	9
Developed Country Partners	9	1	1	9
Trainees	9	1	1	9

Dated: February 23, 2015.

Lynn Susulske,

NHLBI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2015–05722 Filed 3–12–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

U.S. Customs and Border Protection

[1651–0100]

Agency Information Collection Activities: Petition for Remission or Mitigation of Forfeitures and Penalties Incurred

AGENCY: U.S. Customs and Border Protection, Department of Homeland Security.

ACTION: 30-Day notice and request for comments; extension of an existing collection of information.

SUMMARY: U.S. Customs and Border Protection (CBP) of the Department of Homeland Security will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act: Petition for Remission or Mitigation of Forfeitures and Penalties Incurred (CBP Form 4609). This is a proposed extension of an information collection that was previously approved. CBP is proposing that this information collection be extended with a change to the burden hours, but no changes to the information collected. This document is published to obtain comments from the public and affected agencies.

DATES: Written comments should be received on or before April 13, 2015 to be assured of consideration.

ADDRESSES: Interested persons are invited to submit written comments on this proposed information collection to the Office of Information and Regulatory Affairs, Office of Management and Budget. Comments should be addressed to the OMB Desk Officer for Customs and Border Protection, Department of Homeland Security, and sent via electronic mail to *oira_submission@omb.eop.gov* or faxed to (202) 395–5806.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Tracey Denning, U.S. Customs and Border Protection, Regulations and Rulings, Office of International Trade, 90 K Street NE., 10th Floor, Washington, DC 20229–1177, at 202–325–0265.

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** (79 FR 77019) on December 23, 2014, allowing for a 60-day comment period. This notice allows for an additional 30 days for public comments. This process is conducted in accordance with 5 CFR 1320.10. CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3507). The comments should address: (a) Whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden, including the use of automated collection techniques or the use of other forms of information technology; and (e) the annual costs to respondents or record keepers from the collection of information (total capital/startup costs and operations and maintenance costs). The comments that are submitted will be summarized and included in the CBP request for OMB approval. All comments will become a matter of public record. In this document, CBP is soliciting comments concerning the following information collection:

Title: Petition for Remission or Mitigation of Forfeitures and Penalties Incurred.

OMB Number: 1651–0100.

Form Number: Form 4609.

Abstract: CBP Form 4609, Petition for Remission or Mitigation of Forfeitures and Penalties Incurred, is completed and filed with the CBP Port Director by individuals who have been found to be in violation of one or more provisions of the Tariff Act of 1930, or other laws administered by CBP. Persons who violate the Tariff Act are entitled to file a petition seeking mitigation of any statutory penalty imposed or remission of a statutory forfeiture incurred. This petition is submitted on CBP Form 4609. The information provided on this form is used by CBP personnel as a basis for granting relief from forfeiture or penalty. CBP Form 4609 is authorized by 19 U.S.C. 1618 and provided for by 19 CFR 171.1. It is accessible at: *http://www.cbp.gov/sites/default/files/documents/CBP%20Form%204609.pdf*

Action: CBP proposes to extend the expiration date of this information collection with a change to the burden hours resulting from updated estimates of the number of responses. There are no changes to the information collected.

Type of Review: Extension (with change).

Affected Public: Businesses.

Estimated Number of Respondents: 1,610.