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 through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses.

C. Annual Reporting Burden

Respondents: 10,000.
Responses per Respondent: 1.
Total annual responses: 10,000.
Hours per Response: .25.
Total Burden Hours: 2,500.
Obtaining Copies of Proposals: Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (MVCB), 1800 F Street NW., 2nd Floor, Washington, DC 20405–0001, telephone 202–501–4755. Please cite OMB Control No. 3090–00XX, MyUSA, in all correspondence.

Dated: November 25, 2013.

Casey Coleman,

Chief Information Officer.

[FR Doc. 2013–28715 Filed 11–27–13; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Public Meeting of the Presidential Commission for the Study of Bioethical Issues

AGENCY: Presidential Commission for the Study of Bioethical Issues, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

ACTION: Notice of meeting.

SUMMARY: The Presidential Commission for the Study of Bioethical Issues (the Commission) will conduct its fifteenth meeting on December 18, 2013. At this meeting, the Commission will discuss the BRAIN Initiative and ongoing work in neuroscience.

DATES: The meeting will take place Wednesday, December 18, 2013, from 9:00 a.m. to approximately 5:15 p.m.

ADDRESSES: The Hamilton Crowne Plaza Hotel, 1001 14th Street NW., Washington, DC 20005. Telephone (202) 682–0111.

FOR FURTHER INFORMATION CONTACT: Hillary Wicai Viers, Communications Director, Presidential Commission for the Study of Bioethical Issues, 1425 New York Avenue NW, Suite C–100, Washington, DC 20005. Telephone: 202–233–3960. Email: *Hillary.Viers@bioethics.gov.* Additional information may be obtained at *www.bioethics.gov.*

SUPPLEMENTARY INFORMATION: Pursuant to the Federal Advisory Committee Act of 1972, Public Law 92–463, 5 U.S.C. app. 2, notice is hereby given of the fifteenth meeting of the Commission. The meeting will be open to the public with attendance limited to space available. The meeting will also be webcast at *www.bioethics.gov.*

Under authority of Executive Order 13521, dated November 24, 2009, the President established the Commission. The Commission is an expert panel of not more than 13 members who are drawn from the fields of bioethics, science, medicine, technology, engineering, law, philosophy, theology, or other areas of the humanities or social sciences. The Commission advises the President on bioethical issues arising from advances in biomedicine and related areas of science and technology. The Commission seeks to identify and promote policies and practices that ensure scientific research, health care delivery, and technological innovation are conducted in a socially and ethically responsible manner.

The main agenda item for the Commission's fifteenth meeting is to discuss the BRAIN Initiative and ongoing work in neuroscience.

The draft meeting agenda and other information about the Commission, including information about access to the webcast, will be available at *www.bioethics.gov.*

The Commission welcomes input from anyone wishing to provide public comment on any issue before it. Respectful debate of opposing views and active participation by citizens in public exchange of ideas enhances overall public understanding of the issues at hand and conclusions reached by the Commission. The Commission is particularly interested in receiving comments and questions during the meeting that are responsive to specific sessions. Written comments will be accepted at the registration desk and comment forms will be provided to members of the public in order to write down questions and comments for the Commission as they arise. To accommodate as many individuals as possible, the time for each question or comment may be limited. If the number of individuals wishing to pose a question or make a comment is greater than can reasonably be accommodated during the scheduled meeting, the Commission may make a random selection.

Written comments will also be accepted in advance of the meeting and are especially welcome. Please address written comments by email to *info@bioethics.gov,* or by mail to the following address: Public Commentary, Presidential Commission for the Study of Bioethical Issues, 1425 New York Avenue NW., Suite C–100, Washington, DC 20005. Comments will be publicly available, including any personally identifiable or confidential business information that they contain. Trade secrets should not be submitted.

Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, should notify Esther Yoo by telephone at (202) 233–3960, or email at *Esther.Yoo@bioethics.gov* in advance of the meeting. The Commission will make every effort to accommodate persons who need special assistance.

Dated: November 12, 2013.

Lisa M. Lee,

Executive Director, Presidential Commission for the Study of Bioethical Issues.

[FR Doc. 2013–28621 Filed 11–27–13; 8:45 am]

BILLING CODE 4154–06–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30 Day–14–13AAH]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call (404) 639–7570 or send an email to *omb@cdc.gov.* Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

CDC Work@Health Program: Phase 2 Training and Technical Assistance Evaluation—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

The Centers for Disease Control and Prevention (CDC) is establishing the Work@Health Program, a comprehensive worksite health promotion training program, to support employers' efforts to create healthy work environments and provide employees with opportunities to make healthy lifestyle choices. The Work@Health Program will train and support small, mid-size, and large employers with three primary goals: (1) Increase understanding of the training needs of employers and the best way to deliver skill-based training to them; (2) Increase employers' level of knowledge and awareness of worksite health program concepts and principles as well as tools and resources to support the design, implementation, and evaluation of effective worksite health strategies and interventions; and (3) Increase the number of science-based worksite health programs, policies, and practices in place at participating employers' worksites and increase the access and opportunities for employees to participate in them.

CDC is requesting OMB approval to initiate Phase 2 information collection. Phase 2 procedures were informed by a needs assessment and pilot test that were conducted in fall 2013 (''CDC Work@Health Program: Phase 1,'' OMB No. 0920–0989, exp. 9/30/2014). In Phase 2, CDC will offer training in four models (formats): (1) A ''Hands-on'' instructor-led workshop model (T1), (2) a self-paced ''Online'' model (T2), (3) a combination or ''Blended'' model (T3), and (4) a ''Train-the-Trainer'' model (T4) designed to prepare qualified individuals to train employers through the Hands-on, Online, or Blended models.

To evaluate the training, information will be collected from approximately 540 employers and approximately 60 individuals who are interested in becoming trained/certified instructors for the Work@Health Program. Respondents will include employers/employees, trainees who participate in the four training models, and training and technical assistance instructors, coaches and subject matter experts.

CDC will use the information collected to evaluate the effectiveness of the Work@Health Program in terms of (1) increasing employer's knowledge and awareness of worksite health concepts, principles, and resources and (2) increasing the number of science-based worksite health programs, policies and practices in place at participating employers' worksites. The information will also be used to identify the best way(s) to deliver skill-based training and technical support to employers in the area of worksite health.

OMB approval is requested for two years. Participation in Work@Health is voluntary and there are no costs to participants other than their time and cost of travel to the training. The total estimated annualized burden hours are 1,601.

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
Interested Employer	Employer Application Form	600	1	20/60
Employers Participating in Work@Health™	CDC Worksite Health Scorecard.	540	1	30/60
	Organizational Assessment	540	1	15/60
	Employer Follow-up Survey	270	1	15/60
	Case Study Interviews with Senior Leadership.	3	1	1
	Case Study Interviews with Employees.	6	1	1
Trainees Participating in the Work@Health™ Program (Hands-on, Online, Blended models).	Trainee KAB Survey	1,080	1	20/60
	Trainee Reaction Survey—Hands-On Model.	180	1	15/60
	Trainee Reaction Survey—Online Model.	180	1	15/60
	Trainee Reaction Survey—Blended Model.	180	1	15/60
	Trainee Technical Assistance Survey.	1,080	1	15/60
	Case Study Interviews with Selected Trainees.	15	1	1
	Trainee Focus Group Discussion Guide.	11	1	1.5
Interested Train-the-Trainer Participants	Train-the-Trainer Application Form.	60	1	30/60
Trainees Participating in the Work@Health™ Program (Train-the-Trainer model).	Train-the-Trainer Participant Survey.	60	1	20/60
	Trainee Reaction Survey—Train-the-Trainer Model.	30	1	15/60
Train-the-Trainer Trainee Technical Assistance Survey		60	1	15/60
Trainees participating in the Work@Health™ Program Wave 2.	Wave 2 Trainee Reaction Survey.	150	1	15/60
Work@Health™ Instructors/Coaches	Instructor/Coach Group Discussion Guide.	21	1	30/60

Leroy Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2013–28592 Filed 11–27–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–194]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (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.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by December 30, 2013:

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be received by the OMB desk officer via one of the following transmissions:

OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–6974 *OR,*

Email: *OIRA_submission@omb.eop.gov.*

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at *http://www.cms.hhs.gov/PaperworkReductionActof1995.*

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@cms.hhs.gov.*

3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term ''collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Reinstatement without change of a previously approved collection; *Title of Information Collection:* Medicare Disproportionate Share Adjustment (DSH) Procedures and Criteria and Supporting Regulations; *Use:* Section 1886(d)(5)(F) of the Social Security Act provides for additional payment to hospitals that serve a disproportionate share of the indigent patient population. This payment is an add-on to the set amount per case that we pay to hospitals under the Medicare Inpatient Prospective Payment System. To meet the qualifying criteria for this additional DSH payment, a hospital must prove that a disproportionate percentage of its patients are low income using Supplemental Security Income and Medicaid as proxies for this determination. Once a hospital qualifies for the DSH payment, we also determine the hospital's payment adjustment. *Form Number:* CMS–R–194 (OCN: 0938–0691); *Frequency:* Occasionally; *Affected Public:* Private sector (business or other for-profits and not-for-profit institutions); *Number of Respondents:* 800; *Total Annual Responses:* 800; *Total Annual Hours:* 400. (For policy questions regarding this collection contact JoAnne Cerne at 410–786–4530.)

Dated: November 22, 2013.

Martique Jones,

Deputy Director, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2013–28524 Filed 11–27–13; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10512, CMS–R–153 and CMS–10277]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including any of the following subjects: (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.

DATES: Comments must be received by January 28, 2014.

ADDRESSES: When commenting, please reference the document identifier or OMB control number (OCN). To be assured consideration, comments and