

Ends: 01/03/2017, Contact: Robert Johnson 305-244-4235.

The U.S. Department of the Interior's National Park Service (NPS) is adopting the U.S. Army Corps of Engineers Final EIS #20140215, filed with EPA 07/31/2014. The NPS was not a cooperating agency for this project. Therefore, recirculation of the document is necessary under Section 1506.3(b) of Council on Environmental Quality Regulations.

EIS No. 20160284, Draft, TVA, TN, Multiple Reservoir Land Management Plans, Comment Period Ends: 01/31/2017, Contact: Matthew Higdon 865-632-8051.

Dated: November 29, 2016.

Dawn Roberts,

Management Analyst, NEPA Compliance Division, Office of Federal Activities.

[FR Doc. 2016-29010 Filed 12-1-16; 8:45 am]

BILLING CODE 6560-50-P

GENERAL SERVICES ADMINISTRATION

[Notice-WWICC-2016-05; Docket No. 2016-0006; Sequence No. 5]

World War One Centennial Commission; Notification of Upcoming Public Advisory Meeting

AGENCY: World War One Centennial Commission, GSA.

ACTION: Meeting notice.

SUMMARY: Notice of this meeting is being provided according to the requirements of the Federal Advisory Committee Act, 5 U.S.C. App. 10(a)(2). This notice provides the schedule and agenda for the December 15, 2016 meeting of the World War One Centennial Commission (the Commission). The meeting is open to the public.

DATES: *Meeting date:* The meeting will be held on Thursday, December 15, 2016 starting at 9:00 a.m. Eastern Standard Time (EST), and ending no later than 10:00 a.m., EST.

The meeting will be held at the Offices of the World War 1 Centennial Commission at 1800 G Street NW., Washington, DC 20006, Street Level. This location is handicapped accessible. The meeting will be open to the public. Persons attending in person are requested to refrain from using perfume, cologne, and other fragrances (see <http://www.access-board.gov/about/policies/fragrance.htm> for more information).

Written Comments may be submitted to the Commission and will be made part of the permanent record of the

Commission. Comments must be received by 5:00 p.m., EST, December 9, 2016 and may be provided by email to daniel.dayton@worldwar1centennial.gov.

Contact Daniel S. Dayton at daniel.dayton@worldwar1centennial.org to register to comment during the meeting's 30-minute public comment period.

Registered speakers/organizations will be allowed 5 minutes and will need to provide written copies of their presentations. Requests to comment, together with presentations for the meeting must be received by 5:00 p.m., EST, Friday, December 9, 2016. Please contact Mr. Dayton at the email address above to obtain meeting materials.

FOR FURTHER INFORMATION CONTACT:

Daniel S. Dayton, Designated Federal Officer, World War 1 Centennial Commission, 701 Pennsylvania Avenue NW., 123, Washington, DC 20004-2608; or telephone 202-380-0725 (note: this is not a toll-free number).

SUPPLEMENTARY INFORMATION:

Background

The World War One Centennial Commission was established by Public Law 112-272 (as amended), as a commission to ensure a suitable observance of the centennial of World War I, to provide for the designation of memorials to the service of members of the United States Armed Forces in World War I, and for other purposes. Under this authority, the Committee will plan, develop, and execute programs, projects, and activities to commemorate the centennial of World War I, encourage private organizations and State and local governments to organize and participate in activities commemorating the centennial of World War I, facilitate and coordinate activities throughout the United States relating to the centennial of World War I, serve as a clearinghouse for the collection and dissemination of information about events and plans for the centennial of World War I, and develop recommendations for Congress and the President for commemorating the centennial of World War I. The Commission does not have an appropriation and operated solely on donated funds.

Agenda: Thursday, December 15, 2016

Old Business

- Acceptance of minutes of last meeting.
- Public Comment Period.

New Business

- Executive Director's Report—Mr. Dayton.

- Fundraising Report—Ambassador Sedgwick.

- Memorial Report—Mr. Fountain.
- Education Report—Dr. O'Connell.
- Endorsements—(RFS)—Dr.

Seefried.

- International Report—Dr. Seefried.
- Report on April 6 Event—Dr.

Seefried.

Other Business

- Chairman's Report.
- Set Next Meeting.
- Motion to Adjourn.

Dated: November 28, 2016.

Daniel S. Dayton,

Designated Federal Official, World War I Centennial Commission.

[FR Doc. 2016-28920 Filed 12-1-16; 8:45 am]

BILLING CODE 6820-95-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-17-17FB; Docket No. CDC-2016-0113]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection entitled "Understanding Relationship Dynamics and Conflict Survey." CDC will use the information collected to ascertain which factors or groups of factors may influence violence perpetration that occurs within adult intimate partner relationships.

DATES: Written comments must be received on or before January 31, 2017.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2016-0113 by any of the following methods:

- *Federal eRulemaking Portal:* [Regulations.gov](http://www.regulations.gov). Follow the instructions for submitting comments.
- *Mail:* Leroy A. Richardson, Information Collection Review Office,

Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. All relevant comments received will be posted without change to *Regulations.gov*, including any personal information provided. For access to the docket to read background documents or comments received, go to *Regulations.gov*.

Please note: All public comment should be submitted through the Federal eRulemaking portal (*Regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact the Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS-D74, Atlanta, Georgia 30329; phone: 404-639-7570; Email: *omb@cdc.gov*.

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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed 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 estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. 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.

Proposed Project

Understanding Relationship Dynamics and Conflict Survey—New—National Center for Injury Prevention and Control (NCIPC), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Intimate partner violence (IPV) is a substantial public health problem in the United States. Over a third of women and over a quarter of men have experienced rape, physical violence, and/or stalking by an intimate partner. Recognition of the importance and prevalence of this issue has fueled research to examine the causes, correlates, and outcomes of IPV over the past several decades. However, studies

across various IPV research domains (*e.g.*, etiology, prevention efficacy and intervention effectiveness) tend to view IPV as an isolated occurrence and rarely consider the contextual situation in which IPV occurs. For example, existing models may not distinguish between an act of physical violence perpetrated during an argument from an act of physical violence perpetrated as a constellation of physical, sexual, and psychological violence by one partner toward another for the purpose of dominating and controlling that partner.

To that end, we need more information about the factors or groups of factors that influence violence perpetration within adult intimate partner relationships. This project will take a critical first step by collecting information from adults in the United States about their attitudes, perceptions, beliefs and experiences with violence in intimate relationships. In the future, this information can help develop a standardized measurement scheme that will distinguish among different contextual forms of IPV perpetration so that effective violence prevention strategies can be targeted and implemented.

The respondent universe consists of 2,210 adults (18 years or older) from two populations: The general population who live in the United States and incarcerated individuals who live in Indiana. Half of the incarcerated group will have an IPV-related offense record and half will not. Data will be collected through an online survey of Mechanical Turk (MT) workers and an in-person survey of incarcerated individuals. Data analysis will include a combination of Factor Analysis and Latent Profile Analysis.

CDC will seek a two-year approval from the Office of for this new collection. There are no cost to respondents other than their time spent responding to the survey/screener.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Response burden (hours)	Total burden hours
Mechanical Turk Survey Respondents.	Screener	4,300	1	5/60	358
Mechanical Turk Survey Respondents.	Understanding Relationship Dynamics and Conflict	1,000	1	1	1,000
Incarcerated Survey Respondents ...	Understanding Relationship Dynamics and Conflict.	105	1	75/60	131
Total	1,489

Leroy A. 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. 2016-28899 Filed 12-1-16; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-R-26]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services.

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: The necessity and utility of the proposed information collection for the proper performance of the agency's functions; the accuracy of the estimated burden; ways to enhance the quality, utility, and clarity of the information to be collected; and 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 31, 2017.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number _____, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

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:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS-R-26 Clinical Laboratory Improvement Amendments (CLIA) Regulations

Under the 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 requires federal agencies to publish a 60-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.

Information Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Clinical

Laboratory Improvement Amendments (CLIA) Regulations; *Use:* The information is necessary to determine an entity's compliance with the Congressionally-mandated program with respect to the regulation of laboratory testing (CLIA). In addition, laboratories participating in the Medicare program must comply with CLIA requirements as required by section 6141 of OBRA 89. Medicaid, under the authority of section 1902(a)(9)(C) of the Social Security Act, pays for services furnished only by laboratories that meet Medicare (CLIA) requirements. *Form Number:* CMS-R-26 (OMB Control Number: 0938-0612); *Frequency:* Monthly, occasionally; *Affected Public:* Private sector—Business or other for-profits and Not-for-profit institutions, State, Local or Tribal Governments, and the Federal government; *Number of Respondents:* 70,861; *Total Annual Responses:* 1,979,300; *Total Annual Hours:* 14,975,785. (For policy questions regarding this collection contact Raelene Perfetto at 410-786-6876).

Dated: November 29, 2016.

William N. Parham, III,
Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2016-29011 Filed 12-1-16; 8:45 am]

BILLING CODE 4120-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10340, CMS-10476, CMS-10525, and CMS-10630]

Agency Information Collection Activities: Submission for OMB Review; 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 (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