

equivalent, passport number, financial account number, or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any “[t]rade secret or any commercial or financial information which is . . . privileged or confidential,” as discussed in Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you have to follow the procedure explained in FTC Rule 4.9(c), 16 CFR 4.9(c).³ Your comment will be kept confidential only if the FTC General Counsel grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at <https://ftcpubcommentworks.com/ftc/paestudypra>, by following the instructions on the web-based form. If this Notice appears at <http://www.regulations.gov/#!home>, you also may file a comment through that Web site.

If you file your comment on paper, write “PAE Reports: Paperwork Comment; Project No. P131203” on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H-113 (Annex J), 600 Pennsylvania Avenue NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at <http://www.ftc.gov> to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will

consider all timely and responsive public comments that it receives on or before December 16, 2013. You can find more information, including routine uses permitted by the Privacy Act, in the Commission’s privacy policy, at <http://www.ftc.gov/ftc/privacy.htm>.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 2013–28528 Filed 11–27–13; 8:45 am]

BILLING CODE 6750–01–P

GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090–00XX; Docket No. 2013–0001; Sequence No. 8]

Submission for OMB Review; MyUSA

AGENCY: Office of Citizen Services and Innovative Technologies (OCSIT), General Services Administration (GSA).

ACTION: Notice of request for comments regarding an extension to an existing OMB clearance.

SUMMARY: Under the provisions of the Paperwork Reduction Act (44 U.S.C. chapter 35), the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to approve a new information collection requirement concerning MyUSA.

DATES: Submit comments on or before December 30, 2013.

ADDRESSES: Submit comments identified by Information Collection 3090–00XX; MyUSA by any of the following methods:

- *Regulations.gov:* <http://www.regulations.gov>. Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link “Submit a Comment” that corresponds with “Information Collection 3090–00XX; MyUSA.” Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 3090–00XX; MyUSA” on your attached document.

- *Fax:* 202–501–4067.

- *Mail:* General Services Administration, Regulatory Secretariat (MVCB), 1800 F Street NW., 2nd Floor, Washington, DC 20405–0001. ATTN: IC 3090–00XX; MyUSA.

Instructions: Please submit comments only and cite Information Collection 3090–00XX; MyUSA, in all correspondence related to this collection. All comments received will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided.

www.regulations.gov, including any personal and/or business confidential information provided.

FOR FURTHER INFORMATION CONTACT:

Sarah Crane, Director, Office of Citizen Services and Innovative Technologies, General Services Administration, at telephone number 202–208–5855, or via email to Sarah.Crane@gsa.gov. For information pertaining to status or publication schedules, contact the Regulatory Secretariat at 202–501–4755.

SUPPLEMENTARY INFORMATION:

A. Purpose

MyUSA (<https://my.usa.gov>) provides an account to users that give them control over their interactions with government agencies and how government uses and accesses their personal information. Users have the option of creating a personal profile that can be reused across government to personalize interactions and streamline common tasks such as filling out forms. Government agencies can build applications that can request permission from the user to access their MyUSA Account and read their personal profile.

The information in the system is contributed voluntarily by the user and cannot be accessed by the government without explicit consent of the user; information is not shared between government agencies, except when the user gives explicit consent to share his or her information, and as detailed in the MyUSA System of Records Notice (SORN) (<http://www.gpo.gov/fdsys/pkg/FR-2013-07-05/pdf/2013-16124.pdf>).

The information collected is basic profile information, and may include: Name, home address, phone number, date of birth, gender, marital status and basic demographic information such as whether the individual is married, a veteran, a small business owner, a parent or a student.

Use of the system, and contribution of personal information, is completely voluntary.

A notice was published in the **Federal Register** at 78 FR 49270, on August 13, 2013. No comments were received.

B. Public Comments

Pursuant to section 3506(c)(2)(A) of the PRA, GSA specifically solicits comments and information to enable it to:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions 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

³ In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c), 16 CFR 4.9(c).

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]

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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]

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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).