

The purpose of this analysis is to facilitate public comment on the Consent Agreement, and it is not intended to constitute an official interpretation of the proposed Decision and Order or the Order to Maintain Assets, or to modify their terms in any way.

By direction of the Commission.

**Donald S. Clark**

*Secretary.*

[FR Doc. 2011–23305 Filed 9–12–11; 8:45 am]

**BILLING CODE 6750–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Office of the Assistant Secretary for Planning and Evaluation; Meeting of the Advisory Council on Alzheimer's Research, Care, and Services

**AGENCY:** Assistant Secretary for Planning and Evaluation, HHS.

**ACTION:** Notice of meeting.

**SUMMARY:** This notice announces public meetings of the Advisory Council on Alzheimer's Research, Care, and Services (Advisory Council). Notice of these meetings is given under the Federal Advisory Committee Act (5 U.S.C. App. 2, section 10(a)(1) and (a)(2)). The Advisory Council on Alzheimer's Research, Care, and Services will provide advice on how to prevent or reduce the burden of Alzheimer's disease and related dementias on people with the disease and their caregivers. Representatives from the Department of Health and Human Services (HHS) will present inventories of Federal activities related to Alzheimer's disease and related dementias in three areas: research, clinical care, and long-term services and support. The representatives will also identify gaps and opportunities in these areas. The Advisory Council will discuss the inventories, gaps, and opportunities, and make recommendations to the Secretary for priority areas and actions for a national plan to address Alzheimer's disease and related dementias.

**Meeting Date:** September 27, 2011, 9:30 a.m. to 4 p.m.

**ADDRESSES:** The meeting will be held at Administration on Aging headquarters at 1 Massachusetts Ave., NW., Washington, DC, 20001, Room 5604/5403.

**Comments:** Time is allocated on the agenda to hear public comments at the end of the meeting. In lieu of oral comments, formal written comments may be submitted for the record to

Helen Lamont, OASPE, 200 Independence Ave., SW., Washington, DC 20201, Room 424E. Those submitting written comments should identify themselves and any relevant organizational affiliations.

#### FOR FURTHER INFORMATION CONTACT:

Helen Lamont (202) 690–7996, [helen.lamont@hhs.gov](mailto:helen.lamont@hhs.gov) **Note:** Although the meeting is open to the public, procedures governing security and the entrance to Federal buildings may change without notice. Those wishing to attend the meeting must call or e-mail Dr. Lamont by Thursday September 22, 2011, so that their name may be put on a list of expected attendees and forwarded to the security officers at the Administration on Aging. Space is limited to 40 participants.

**SUPPLEMENTARY INFORMATION:** Topics of the Meeting: The Advisory Council will hear presentations and provide feedback on inventories of Federal activities to address Alzheimer's disease and related dementias, gaps that can be addressed, and opportunities for collaboration. The Advisory Council is specifically charged with discussing and making recommendations to the Secretary on priorities for a national plan to address Alzheimer's disease and related dementias.

**Procedure and Agenda:** This meeting is open to the public. Representatives of HHS will present the inventories of Federal activities related to Alzheimer's disease and related dementias to the Advisory Council. The representatives will also identify gaps and opportunities in these areas. After each presentation, the Advisory Council will openly discuss the inventory and the findings. Interested persons may observe the discussion, but the Advisory Council will not hear public comments during this time. The Advisory Council will allow an open public session for any attendee to address issues specific to the inventories or topics that should be addressed by a national plan.

**Authority:** 42 U.S.C. 11225; Section 2(e)(3) of the National Alzheimer's Project Act. The panel is governed by provisions of Public Law 92–463, as amended (5 U.S.C. Appendix 2), which sets forth standards for the formation and use of advisory committees.

Dated: September 8, 2011.

**Sherry Glied,**

*Assistant Secretary for Planning and Evaluation.*

[FR Doc. 2011–23465 Filed 9–9–11; 11:15 am]

**BILLING CODE 4150–05–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[60Day–11–0666]

#### Proposed Data Collections Submitted for Public Comment and Recommendations

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 Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Daniel Holcomb, CDC Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to [omb@cdc.gov](mailto:omb@cdc.gov).

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; and (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. Written comments should be received within 60 days of this notice.

#### Proposed Project

National Healthcare Safety Network (NHSN) (OMB No. 0920–0666) exp. 05/31/2014—Revision—National Center for Emerging and Zoonotic Infectious Diseases (NCEZID), Centers for Disease Control and Prevention (CDC).

#### Background and Brief Description

The National Healthcare Safety Network (NHSN) is a system designed to accumulate, exchange, and integrate relevant information and resources among private and public stakeholders to support local and national efforts to protect patients and promote healthcare safety. Specifically, the data is used to determine the magnitude of various healthcare-associated adverse events and trends in the rates of these events among patients and healthcare workers with similar risks. The data will be used to detect changes in the epidemiology of