

Date first accepted by the commission	Docket No.	Matter name
7. 11/24/09 .....	C 4275	SCI/Palm.
8. 3/25/10 .....	C 4284	SCI/Keystone.
Hospitals and other clinics		
9. 03/30/06 .....	C 4159	Fresenius AG.
10. 10/07/09 .....	D 9338	Carilion Clinic.
11. 11/25/10 .....	C 4309	Universal/PSI.
12. 07/21/11 .....	C 4339	Cardinal/Biotech.
13. 09/02/11 .....	C 4334	Davita/DSI.
14. 02/28/12 .....	C 4348	Fresenius AG.
15. 10/5/12 .....	C 4372	Universal/Ascend.

By direction of the Commission.

**Donald S. Clark,**

*Secretary.*

[FR Doc. 2015-00666 Filed 1-15-15; 8:45 am]

**BILLING CODE 6750-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Advisory Council on Alzheimer's Research, Care, and Services; Meeting

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

**ACTION:** Notice of meeting.

**SUMMARY:** This notice announces the public meeting of the Advisory Council on Alzheimer's Research, Care, and Services (Advisory Council). The Advisory Council on Alzheimer's Research, Care, and Services provides advice on how to prevent or reduce the burden of Alzheimer's disease and related dementias on people with the disease and their caregivers. During the January meeting, the Advisory Council will hear a presentation on IOM's final expert panel on Advanced Dementia, which will provide additional recommendations for the Council to consider. The Advisory Council will spend the majority of the meeting considering recommendations made by each of the three subcommittees for updates to the 2015 National Plan.

**DATES:** The meeting will be held on January 26th, 2014 from 9 a.m. to 5 p.m. EDT.

**ADDRESSES:** The meeting will be held in the Great Hall in the Hubert H. Humphrey Building, 200 Independence Avenue SW., Washington, DC 20201.

**Comments:** Time is allocated mid-morning on the agenda to hear public comments. The time for oral comments will be limited to two (2) minutes per individual. In lieu of oral comments, formal written comments may be submitted for the record to Rohini Khillan, OASPE, 200 Independence Avenue SW., Room 424E, Washington,

DC 20201. Comments may also be sent to [napa@hhs.gov](mailto:napa@hhs.gov). Those submitting written comments should identify themselves and any relevant organizational affiliations.

#### FOR FURTHER INFORMATION CONTACT:

Rohini Khillan (202) 690-5932, [rohini.khillan@hhs.gov](mailto:rohini.khillan@hhs.gov). Note: Seating may be limited. Those wishing to attend the meeting must send an email to [napa@hhs.gov](mailto:napa@hhs.gov) and put "January 26 Meeting Attendance" in the Subject line by Friday, January 16, so that their names may be put on a list of expected attendees and forwarded to the security officers at the Department of Health and Human Services. Any interested member of the public who is a non-U.S. citizen should include this information at the time of registration to ensure that the appropriate security procedure to gain entry to the building is carried out. Although the meeting is open to the public, procedures governing security and the entrance to Federal buildings may change without notice. If you wish to make a public comment, you must note that within your email.

**SUPPLEMENTARY INFORMATION:** 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)). Topics of the Meeting: The Advisory Council will hear presentations on the basics of long-term care, including presentations on programs, settings, and payers. The Council will use a portion of the meeting to review the work it has accomplished thus far towards the 2025 goals, and then discuss the process for developing recommendations for the 2015 update to the National Plan. The Council will also hear presentations from the three subcommittees (Research, Clinical Care, Long-Term Services and Supports, and Ethics).

**Procedure and Agenda:** This meeting is open to the public. Please allow 30 minutes to go through security and walk to the meeting room. The meeting will also be webcast at [www.hhs.gov/live](http://www.hhs.gov/live).

**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: January 5, 2015.

**Richard G. Frank,**

*Assistant Secretary for Planning and Evaluation.*

[FR Doc. 2015-00517 Filed 1-15-15; 8:45 am]

**BILLING CODE 4150-28-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[30Day-15-14AYC]

### Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) 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; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) 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; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639-7570 or send an email to [omb@cdc.gov](mailto:omb@cdc.gov). Written comments and/or suggestions regarding the items contained in this notice should be directed to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395-5806. Written comments should be received within 30 days of this notice.

### Proposed Project

Behavioral Risk Factor Surveillance System (BRFSS)—Existing Collection Without an OMB Control Number—National Center for Chronic Disease Prevention and Health Promotion

(NCCDPHP)—Centers for Disease Control and Prevention (CDC).

### Background and Brief Description

CDC is requesting OMB approval to conduct information collection for the Behavioral Risk Factor Surveillance System (BRFSS) for three years beginning with the 2015 data collection cycle. The BRFSS is a nationwide system of customized, cross-sectional telephone health surveys sponsored by CDC. Information collection is conducted in a continuous, three-part telephone interview process: screening, participation in a common BRFSS core survey, and participation in optional question modules that states use to customize survey content. BRFSS coordinators in health departments in U.S. states, territories, and the District of Columbia (collectively referred to as states) are responsible for questionnaire content and survey administration. CDC provides the states with technical and methodological assistance.

The BRFSS produces state-level information on adults 18 years and

older primarily on the health risk behaviors, health conditions, and preventive health practices that are associated with chronic diseases, infectious diseases, and injury. This information is used by state and local health departments to plan and evaluate public health programs at the state or sub-state level. For most states and territories, the BRFSS provides the only source of data amenable to state and local level health and health risk indicators.

Information collected through the BRFSS is also used by the federal government and other entities. CDC makes annual BRFSS data sets available for public use and provides guidance on statistically appropriate uses of the data. CDC's authority to collect this information is provided by the Public Health Service Act. Participation in the BRFSS is voluntary and there are no costs to respondents other than their time. The total estimated annualized burden hours are 255,915.

### ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hr)
U.S. General Population .....	Landline Screener .....	440,486	1	1/60
	Cell Phone Screener .....	223,334	1	1/60
Adults ≥ 18 Years .....	BRFSS Core Survey .....	494,650	1	15/60
	BRFSS Optional Modules .....	484,757	1	15/60

**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. 2015-00562 Filed 1-15-15; 8:45 am]

BILLING CODE 4163-18-P

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier CMS-10114]

### 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 *February 17, 2015*.

**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) 395-5806 *OR*, Email: [OIRA\\_submission@omb.eop.gov](mailto: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](mailto:Paperwork@cms.hhs.gov).

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