

states, the Pacific Islands, and U.S. territories (for a total of 62 awardees) to improve surge capacity and enhance community and hospital preparedness for public health emergencies. These 62 awardees are responsible for enhancing the preparedness of the nation's nearly 6000 hospitals. These awards are authorized under section 391C–2 of the Public Health Service (PHS) Act.

For this data collection the situation will dictate how often the data will be collected using the web-based interface known as HAvBED. For a large scale emergency data will be collected nationally from all 62 HPP awardees to include all 6000 hospitals in HAvBED system. For smaller scale events data collection will be targeted to individual states or regions. Data may also be

gathered during exercises. Notifications for data collection are sent to the affected states through the HPP program staff. The data gathered from the hospitals are reported to the HHS Secretary's Operations Center to inform situational awareness and national preparedness.

ANNUAL ESTIMATED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses/ respondent	Average burden hours per response	Total burden hours
Hospital staff (training)	6,000	1	1	6,000
Hospital staff (data collection)	6,000	102	1	612,000
State/Territory Preparedness staff (training)	62	1	1	62
State/Territory Preparedness staff (data collection)	62	102	3	18,972
Total	31,154

Seleda Perryman,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Notice of Meeting: Secretary's Advisory Committee on Genetics, Health, and Society

Pursuant to Public Law 92–463, notice is hereby given of the twenty-second meeting of the Secretary's Advisory Committee on Genetics, Health, and Society (SACGHS), U.S. Public Health Service. The meeting will be held from 8:30 a.m. to approximately 5:30 p.m. on Tuesday, June 15, 2010, and from 8 a.m. to approximately 2:45 p.m. on Wednesday, June 16, 2010, at the Washington Plaza Hotel, 10 Thomas Circle, NW., Washington, DC 20005. The meeting will be open to the public with attendance limited to space available. The meeting will also be Web cast.

The main agenda item will be an exploratory session on the implications of affordable whole-genome sequencing. The meeting will also include updates and discussions on other issues SACGHS has been addressing, including the work of the Secretary's Advisory Committee on Heritable Disorders in Newborns and Children related to the retention and use of dried blood spot specimens from newborn screening.

As always, the Committee welcomes hearing from anyone wishing to provide

public comment on any issue related to genetics, health and society. Please note that because SACGHS operates under the provisions of the Federal Advisory Committee Act, all public comments will be made available to the public. Individuals who would like to provide public comment should notify the SACGHS Executive Secretary, Ms. Sarah Carr, by telephone at 301–496–9838 or e-mail at carrs@od.nih.gov. The SACGHS office is located at 6705 Rockledge Drive, Suite 750, Bethesda, MD 20892. Anyone planning to attend the meeting who needs special assistance, such as sign language interpretation or other reasonable accommodations, is also asked to contact the Executive Secretary.

Under authority of 42 U.S.C. 217a, Section 222 of the Public Health Service Act, as amended, the Department of Health and Human Services established SACGHS to serve as a public forum for deliberations on the broad range of human health and societal issues raised by the development and use of genetic and genomic technologies and, as warranted, to provide advice on these issues. The draft meeting agenda and other information about SACGHS, including information about access to the Web cast, will be available at the following Web site: http://oba.od.nih.gov/SACGHS/sacghs_meetings.html.

Dated: April 16, 2010.

Jennifer Spaeth,

Director, NIH Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier CMS–10316 and CMS–10209]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare & Medicaid Services (CMS) is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this 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.

1. *Type of Information Collection Request:* New collection; *Title of Information Collection:* Medicare Prescription Drug Plan (PDP) and Medicare Advantage Prescription Drug Plan (MA–PD) Disenrollment Reasons Survey; *Use:* The Medicare Prescription Drug, Improvement, and Modernization