DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2016–0090; Docket Number NIOSH 288–A]

A Performance Test Protocol for Closed System Transfer Devices Used During Pharmacy Compounding and Administration of Hazardous Drugs; Extension of Comment Period

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice and extension of comment period.

SUMMARY: On September 15, 2016 the National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), published a notice in the **Federal Register** [81 FR 63482] announcing a public meeting and request for public comment on a draft testing protocol.

Written comments were to be received by December 7, 2016. NIOSH initially extended the public comment period to June 7, 2017 [81 FR 88687]. NIOSH extended the comment period again to August 30, 2017 [82 FR 25290]. NIOSH is extending the public comment period to close on February 28, 2018. The longer timeframe will allow companies to test the protocol with the proposed challenge agents and permit full participation in the protocol design process.

FOR FURTHER INFORMATION CONTACT: Deborah V. Hirst, NIOSH, Alice Hamilton Laboratories, 1090 Tusculum Avenue, MS R–5, Cincinnati, Ohio 45226, telephone (513) 841–4141 (not a toll free number), Email: *DHirst@cdc.gov.*

ADDRESSES: You may submit comments, identified by CDC–2016–0090 and Docket Number NIOSH 288–A, by either of the following two methods:

- *Federal eRulemaking Portal: http://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, Ohio 45226–1998.

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2017–15727 Filed 7–25–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Sunshine Act Meeting: Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ATSDR)

TIMES AND DATES:

8:30 a.m.–4:30 p.m., EDT, September 13, 2017

8:30 a.m.–11:30 a.m., EDT, September 14, 2017

PLACE: CDC, 4770 Buford Highway, Building 102, Conference Room 2202, Atlanta, Georgia 30341.

STATUS: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention announces the meeting of the BSC, NCEH/ATSDR. This meeting is open to the public. The meeting room accommodates approximately 60 people. The public is also welcome to listen to the meeting by joining the teleconference at the USA toll-free, dial-in number, 1–888–790–2009 Passcode: 7865774. The deadline for notification of attendance is August 30, 2017. The public comment period is scheduled on Wednesday, September 13, 2017 from 2:00 p.m. until 2:15 p.m.; from 2:40 p.m. until 2:55 p.m.; and from 3:25 p.m. until 3:40 p.m., and on Thursday, September 14, 2017 from 10:10 a.m. until 10:25 a.m. EDT (15 minutes). Individuals wishing to make a comment during Public Comment period, please email your name, organization, and phone number by Monday, September 4, 2017 to Dr. William Cibulas at *wic1@cdc.gov.*

MATTERS TO BE CONSIDERED: The Secretary, Department of Health and Human Services (HHS) and by delegation, the Director, CDC and Administrator, NCEH/ATSDR, are authorized under Section 301 (42 U.S.C. 241) and Section 311 (42 U.S.C. 243) of the Public Health Service Act, as amended, to: (1) Conduct, encourage, cooperate with, and assist other appropriate public authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes, diagnosis, treatment, control, and prevention of physical and mental diseases and other impairments; (2) assist states and their political subdivisions in the prevention of infectious diseases and other preventable conditions and in the promotion of health and wellbeing; and (3) train state and local personnel in health work. The BSC, NCEH/ATSDR provides advice and guidance to the Secretary, HHS; the Director, CDC and Administrator, ATSDR; and the Director, NCEH/ATSDR, regarding program goals, objectives, strategies, and priorities in fulfillment of the agency's mission to protect and promote people's health. The Board provides advice and guidance that will assist NCEH/ATSDR in ensuring scientific quality, timeliness, utility, and dissemination of results. The Board also provides guidance to help NCEH/ATSDR work more efficiently and effectively with its various constituents and to fulfill its mission in protecting America's health. The agenda items for the BSC Meeting will include NCEH/ATSDR Director Updates; Noise-Induced Hearing Loss; NCEH/ATSDR Program Responses to BSC Guidance and Action Items; Lead Poisoning Prevention Program Updates; Flint Registry; Revision of blood lead level reference value (status); Discussion of Legislative Requirements of new Lead Exposure Poisoning Federal Advisory Committee; Amyotrophic Lateral Sclerosis (ALS) Program Update; Environmental Health Tracking Program update; updates from the National Institute of Environmental Health Sciences, the National Institute for Occupational Safety and Health, the US Department of Energy and the US Environmental Protection Agency.

Agenda items are subject to change as priorities dictate.

CONTACT PERSON FOR MORE INFORMATION: Shirley Little, NCEH/ATSDR, CDC, 4770 Buford Highway, Mail Stop F–45, Atlanta, Georgia 30341; Telephone 770/488–0577, Email: *snl7@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and

Prevention and the Agency for Toxic Substances and Disease Registry.

Claudette Grant,

Acting Director, Management Analysis and Services Office Centers for Disease Control and Prevention.

[FR Doc. 2017–15782 Filed 7–24–17; 11:15 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Sunshine Act Meeting: Board of Scientific Counselors, National Center for Health Statistics (NCHS)

TIMES AND DATES:

11:00 a.m.–5:30 p.m., EDT, September 6, 2017

8:30 a.m.–1:00 p.m., EDT, September 7, 2017

PLACE: NCHS Headquarters, 3311 Toledo Road, Hyattsville, Maryland 20782.

STATUS: In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC), National Center for Health Statistics (NCHS) announces the following meeting of the aforementioned committee. This meeting is open to the public; however, visitors must be processed in accordance with established federal policies and procedures. For foreign nationals or non-U.S. citizens, pre-approval is required (please contact Gwen Mustaf, 301–458–4500, *glm4@cdc.gov,* or Virginia Cain, *vcain@cdc.gov* at least 10 days in advance for requirements). All visitors are required to present a valid form of picture identification issued by a state, federal or international government. As required by the Federal Property Management Regulations, Title 41, Code of Federal Regulation, Subpart 101–20.301, all persons entering in or on Federal controlled property and their packages, briefcases, and other containers in their immediate possession are subject to being x-rayed and inspected. Federal law prohibits the knowing possession or the causing to be present of firearms, explosives and other dangerous weapons and illegal substances. The meeting room accommodates approximately 78 people.

MATTERS TO BE CONSIDERED: This committee is charged with providing advice and making recommendations to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NCHS, regarding the scientific and technical program goals and objectives, strategies, and priorities of NCHS. The agenda includes welcome remarks by NCHS leadership; update from the Division of Health Care Statistics; update on National Committee on Vital and Health Statistics (NCVHS) activities; update on improving data collection.

Requests to make oral presentations should be submitted in writing to the contact person listed below. All requests must contain the name, address, telephone number, and organizational affiliation of the presenter. Written comments should not exceed five single-spaced typed pages in length and must be received by August 22, 2017. Agenda items are subject to change as priorities dictate.

CONTACT PERSON FOR MORE INFORMATION: Virginia S. Cain, Ph.D., Director of Extramural Research, NCHS/CDC, 3311 Toledo Road, Room 2627, Hyattsville, Maryland 20782, telephone (301) 458–4500, email *vcain@cdc.gov.*

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and the Agency for Toxic Substances and Disease Registry.

Claudette Grant,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2017–15783 Filed 7–24–17; 11:15 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–1984–14, CMS–10326, CMS–2088–17, CMS–10452, CMS–10320 and CMS–10418]

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 other aspect of this collection of information, including 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 on the collection(s) of information must be received by the OMB desk officer by August 25, 2017.

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.*

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 Web site address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

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: William Parham at (410) 786–4669.

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. 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 (44 U.S.C.