

(9) Are there biological indices or metrics that should be used to aid in the interpretation of biomonitoring data for 1-BP? What is the most appropriate biomarker that can confirm and quantify occupational exposures to 1-BP?

(10) Should acute exposure recommendations, such as a short term exposure limit (STEL), be derived for 1-BP? If so, what data support the development of the STEL?

(11) NIOSH provided Globally Harmonized System (GHS) of Classification and Labelling of Chemicals designations for health endpoints evaluated in the criteria document. Please comment on the utility of these classifications for hazard communication. Are these classifications helpful for employers?

II. Public Meeting

NIOSH will hold a public meeting on the draft document *Criteria for a Recommended Standard: Occupational Exposure to 1-Bromopropane (1-BP)* to allow commenters to provide oral comments on the draft document, to inform NIOSH about additional relevant data or information, and to ask questions on the draft document and NIOSH recommendations.

This meeting is open to the public. Attendance is limited only by the space available. The meeting room accommodates 100 people. The meeting will be open to a limited number of participants through a conference call phone number and Webcast live on the Internet.

Notification of intent to attend the meeting, for in-person and remote participation, must be made to the NIOSH Docket Office, at nioshdocket@cdc.gov, (513) 533-8611, no later than March 16, 2016. Priority for attendance will be given to those providing oral comments. Other requests to attend the meeting will then be accommodated on a first-come, first-served basis.

Registration is required for in-person attendance and remote participation. Because this meeting is being held at a federal site, pre-registration is required on or before March 16, 2016 and a government-issued photo ID (driver's license, military ID or passport) will be required to obtain entrance to the facility. There will be an airport-type security check. Non-US citizens need to register by February 24, 2016 to allow sufficient time for mandatory facility security clearance procedures to be completed. This information will be transmitted to the CDC Security Office for approval. An email confirming registration will be sent from NIOSH for both in-person participation and audio conferencing participation.

Oral comments will be permitted for 15 minutes. If additional time becomes available, presenters will be notified. All requests to present should contain the name, address, telephone number, and relevant business affiliations of the presenter, topic of the presentation, whether you will be presenting in person or by phone, and the approximate time requested for the presentation. An email confirming registration will be sent from the NIOSH Docket Office and will include details needed to participate. Oral comments given at the meeting will be recorded and included in the docket.

After reviewing the requests for presentations, NIOSH will notify the presenter when his/her presentation is scheduled. If a participant is not in attendance when his/her presentation is scheduled to begin, the remaining participants will be heard in order. After the last scheduled speaker is heard, participants who missed their assigned times may be allowed to speak, limited by time available.

Attendees who wish to speak but did not submit a request for the opportunity to make a presentation may be given this opportunity after the scheduled speakers are heard, at the discretion of the presiding officer and limited by time available.

You may submit comments, identified by CDC 2016-0003 and NIOSH 057-A, by either of the following methods:

- *Federal eRulemaking Portal:* 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.

Instructions: All information received in response to this notice must include the agency name and docket number [CDC-2016-0003; NIOSH 057-A]. All relevant comments received will be posted without change to www.regulations.gov including any personal information provided. All information will be available for public examination and copying at the NIOSH Docket Office, 1150 Tusculum Avenue, Room 155, Cincinnati, Ohio 45226.

Non-U.S. Citizens: Because of CDC Security Regulations, any non-U.S. citizen wishing to attend this meeting must provide the following information in writing to the NIOSH Docket Officer at the address below no later than February 24, 2016.

Name:
Gender:
Date of Birth:
Place of Birth (city, province, state, country):
Citizenship:
Passport Number:

Date of Passport Issue:
Date of Passport Expiration:
Type of Visa:
U.S. Naturalization Number (if a naturalized citizen):
U.S. Naturalization Date (if a naturalized citizen):
Visitor's Organization:
Organization Address:
Organization Telephone Number:
Visitor's Position/Title within the Organization:

This information will be transmitted to the CDC Security Office for approval. Visitors will be notified as soon as approval has been obtained.

Public Review

The external review of the draft document has been (1) developed in accordance with Office of Management and Budget (OMB) guidelines, (2) is consistent with NIOSH peer review practice, and (3) is meant to ensure that credible and appropriate science is reflected within the draft document.

Dated: February 4, 2016.

John Howard,

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

[FR Doc. 2016-02650 Filed 2-9-16; 8:45 am]

BILLING CODE 4163-19-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

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) announces a meeting for the initial review of applications in response to Funding Opportunity Announcement (FOA) DP16-004, Childhood Obesity Research Demonstration 2.0.

Time and Date: 10:00 a.m.–6:00 p.m., EST, March 15–16, 2016 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92-463.

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to FOA DP16-004, Childhood Obesity Research Demonstration 2.0.

Contact Person for More Information: Jaya Raman, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway, Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488-6511, kva5@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.

Catherine Ramadei,

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

[FR Doc. 2016-02582 Filed 2-9-16; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number CDC-2016-0001; NIOSH-260-A]

Draft Current Intelligence Bulletin: Health Effects of Occupational Exposure to Silver Nanomaterials; Notice of Public Meeting; Availability of Document for Comment; 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 January 21, 2016, the Director of 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 3425] announcing the availability of the following draft document for public comment entitled *Draft Current Intelligence Bulletin: Health Effects of Occupational Exposure to Silver Nanomaterials*. Written comments were to be received by March 21, 2016. NIOSH is extending the public comment period until April 22, 2016.

DATES: NIOSH is extending the comment period on the document published January 21, 2016 (81 FR 3425). Electronic or written comments must be received by April 22, 2016.

ADDRESSES: You may submit comments, identified by CDC-2016-0001 and

docket number NIOSH-260-A, by any of the following methods:

- **Federal eRulemaking Portal:** 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.

FOR FURTHER INFORMATION CONTACT:

Charles Geraci, NIOSH, Education and Information Division, Nanotechnology Research Center, 1090 Tusculum Avenue, Cincinnati, Ohio 45226, telephone (513) 533-8339 (not a toll free number).

Dated: February 3, 2016.

John Howard,

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

[FR Doc. 2016-02647 Filed 2-9-16; 8:45 am]

BILLING CODE 4163-19-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-276, CMS-1957, CMS-10599 and CMS-10600]

Agency Information Collection Activities: Proposed Collection; 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 (the 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 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates 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 must be received by April 11, 2016:

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number __, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

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.

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

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786-1326.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS-276 Prepaid Health Plan Cost Report
 CMS-1957 Social Security Office (SSO) Report of State Buy-in Problem
 CMS-10599 Medicare Prior Authorization of Home Health Services Demonstration
 CMS-10600 Evaluation of the Medicare Patient Intravenous Immunoglobulin Demonstration

Under the PRA (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management