

GENERAL SERVICES ADMINISTRATION

[Notice MG–2021–03; Docket No. 2021–0002; Sequence No. 25]

Office of Federal High-Performance Green Buildings; Green Building Advisory Committee; Notification of Upcoming Web-Based Meetings

AGENCY: Office of Government-wide Policy, General Services Administration (GSA).

ACTION: Notice of public meetings.

SUMMARY: Notice of these web-based public meetings/conference calls is being provided according to the requirements of the Federal Advisory Committee Act. This notice provides the schedule for one full Committee meeting of the Green Building Advisory Committee (Committee), which is open to the public. Interested individuals must register to attend and provide public comment as instructed below under **SUPPLEMENTARY INFORMATION**.

DATES: The Green Building Advisory Committee will hold a web-based meeting on Tuesday, November 16, 2021, from 11:00 a.m. to 4:00 p.m., Eastern Time (ET).

FOR FURTHER INFORMATION CONTACT: Michael Bloom, Designated Federal Officer, Office of Federal High-Performance Green Buildings, Office of Government-wide Policy, General Services Administration, at michael.bloom@gsa.gov or 312–805–6799. Additional information about the Committee, including meeting materials and agendas, will be available on-line at <http://www.gsa.gov/gbac>.

SUPPLEMENTARY INFORMATION:

Background

The Administrator of GSA established the Committee on June 20, 2011 (Federal Register/Vol. 76, No. 118) pursuant to Section 494 of the Energy Independence and Security Act of 2007 (EISA, 42 U.S.C. 17123). Under this authority, the Committee provides independent policy advice and recommendations to GSA to advance federal building innovations in planning, design, and operations to reduce costs, enable agency missions, enhance human health and performance, and minimize environmental impacts.

Procedures for Attendance and Public Comment

Contact Michael Bloom, at michael.bloom@gsa.gov, to register to attend this public web-based meetings. To register, submit your full name,

organization, email address, phone number. Requests to attend this web-based meeting must be received by 5:00 p.m. ET, on Wednesday, November 10, 2021. Meeting call-in information will be provided to interested parties who register by the deadline. (GSA will be unable to provide technical assistance to any listener experiencing technical difficulties. Testing access to the web-based meeting site before the meetings is recommended.)

Contact Mr. Bloom to register to provide public comment during the November 16th, 2021 meeting public comment period. Registered speakers/organizations will be allowed a maximum of five minutes each and will need to provide written copies of their presentations. Requests to provide public comment at the Committee meeting must be received by 5:00 p.m., ET, on Monday, November 8, 2021.

November 16, 2021 Meeting Agenda

- Updates and introductions
- Federal Building Decarbonization Task Group: Findings & Recommendations
- Environmental Justice and Equity Task Group: Findings & Recommendations
- Energy Storage Task Group: Final Vote to Accept Advice Letter
- New committee directions & topics to explore
- Public comment
- Next steps and closing comments

Kevin Kampschroer,

Federal Director, Office of Federal High-Performance Green Buildings, Office of Government-wide Policy, General Services Administration.

[FR Doc. 2021–23068 Filed 10–21–21; 8:45 am]

BILLING CODE 6820–14–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2021–0112]

Advisory Committee on Immunization Practices (ACIP)

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of meeting and request for comment.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC) announces the following meeting of the Advisory

Committee on Immunization Practices (ACIP). This meeting is open to the public. Time will be available for public comment. The meeting will be webcast live via the World Wide Web; for more information on ACIP please visit the ACIP website: <http://www.cdc.gov/vaccines/acip/index.html>.

DATES: The meeting will be held on November 2–3, 2021, from 10:00 a.m. to 5:00 p.m., EDT (times subject to change). The public may submit written comments from October 22, 2021 through November 3, 2021.

ADDRESSES: You may submit comments identified by Docket No. CDC–2021–0112 by any of the following methods:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments.

- *Mail:* Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H24–8, Atlanta, Georgia 30329–4027, Attn: ACIP Meeting.

Instructions: All submissions received must include the Agency name and Docket Number. All relevant comments received in conformance with the <https://www.regulations.gov> suitability policy will be posted without change to <https://www.regulations.gov>, including any personal information provided. For access to the docket to read background documents or comments received, go to <https://www.regulations.gov>.

Written public comments submitted up to 72 hours prior to the ACIP meeting will be provided to ACIP members before the meeting.

FOR FURTHER INFORMATION CONTACT: Stephanie Thomas, ACIP Committee Management Specialist, Centers for Disease Control and Prevention, National Center for Immunization and Respiratory Diseases, 1600 Clifton Road NE, MS–H24–8, Atlanta, Georgia 30329–4027; Telephone: (404) 639–8367; Email: ACIP@cdc.gov.

SUPPLEMENTARY INFORMATION: In accordance with 41 CFR 102–3.150(b), less than 15 calendar days' notice is being given for this meeting due to the exceptional circumstances of the COVID–19 pandemic and rapidly evolving COVID–19 vaccine development and regulatory processes. The Secretary of Health and Human Services has determined that COVID–19 is a Public Health Emergency. A notice of this ACIP meeting has also been posted on CDC's ACIP website at: <http://www.cdc.gov/vaccines/acip/index.html>. In addition, CDC has sent notice of this ACIP meeting by email to those who subscribe to receive email updates about ACIP.

Purpose: The committee is charged with advising the Director, CDC, on the

use of immunizing agents. In addition, under 42 U.S.C. 1396s, the committee is mandated to establish and periodically review and, as appropriate, revise the list of vaccines for administration to vaccine-eligible children through the Vaccines for Children program, along with schedules regarding dosing interval, dosage, and contraindications to administration of vaccines. Further, under provisions of the Affordable Care Act, section 2713 of the Public Health Service Act, immunization recommendations of the ACIP that have been approved by the CDC Director and appear on CDC immunization schedules must be covered by applicable health plans.

Matters To Be Considered: The agenda will include discussions on adult immunization schedule, child/adolescent immunization schedule, Ebola vaccine, hepatitis vaccines, Orthopoxviruses vaccine and COVID vaccines. Recommendation votes on adult immunization schedule, child/adolescent immunization schedule, hepatitis vaccine, Orthopoxviruses vaccine, Ebola vaccine and COVID vaccines are scheduled. No Vaccines for Children votes are scheduled. Agenda items are subject to change as priorities dictate. For more information on the meeting agenda visit <https://www.cdc.gov/vaccines/acip/meetings/meetings-info.html>.

Public Participation

Interested persons or organizations are invited to participate by submitting written views, recommendations, and data. Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on <https://www.regulations.gov>. Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted into the docket.

Written Public Comment: The docket will be opened to receive written comments on October 22, 2021. Written

comments must be received on or before November 3, 2021.

Oral Public Comment: This meeting will include time for members of the public to make an oral comment. Oral public comment will occur before any scheduled votes including all votes relevant to the ACIP's Affordable Care Act and Vaccines for Children Program roles. Priority will be given to individuals who submit a request to make an oral public comment before the meeting according to the procedures below.

Procedure for Oral Public Comment: All persons interested in making an oral public comment at the November 2–3, 2021 ACIP meeting must submit a request at <http://www.cdc.gov/vaccines/acip/meetings/> no later than 11:59 p.m., EDT, October 31, 2021, according to the instructions provided.

If the number of persons requesting to speak is greater than can be reasonably accommodated during the scheduled time, CDC will conduct a lottery to determine the speakers for the scheduled public comment session. CDC staff will notify individuals regarding their request to speak by email by November 1, 2021. To accommodate the significant interest in participation in the oral public comment session of ACIP meetings, each speaker will be limited to 3 minutes, and each speaker may only speak once per meeting.

The Director, Strategic Business Initiatives Unit, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, 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.

Kalwant Smagh,

*Director, Strategic Business Initiatives Unit,
Office of the Chief Operating Officer, Centers
for Disease Control and Prevention.*

[FR Doc. 2021–23222 Filed 10–20–21; 4:15 pm]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–2567, CMS–10790 and CMS–10463]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 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 must be received by December 21, 2021.

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.