subcontractors for that first-tier subcontractor's preceding completed fiscal year at *http://www.fsrs.gov,* if—

(i) In the subcontractor's preceding fiscal year, the subcontractor received—

(A) 80 percent or more of its annual gross revenues from Federal contracts (and subcontracts), loans, grants (and subgrants), cooperative agreements, and other forms of Federal financial assistance; and

(B) $25,000,000 or more in annual gross revenues from Federal contracts (and subcontracts), loans, grants (and subgrants), cooperative agreements, and other forms of Federal financial assistance; and

(ii) The public does not have access to information about the compensation of the executives through periodic reports filed under section 13(a) or 15(d) of the Securities Exchange Act of 1934 (15 U.S.C. 78m(a), 78o(d)) or section 6104 of the Internal Revenue Code of 1986. (To determine if the public has access to the compensation information, see the U.S. Security and Exchange Commission total compensation filings at *http://www.sec.gov/answers/execomp.htm.*)

FAR 52.204–10(d)(1) requires contractors to report the names and total compensation of each of the five most highly compensated executives for its preceding completed fiscal year as part of the contractor's annual registration requirement in the System for Award Management (SAM) (FAR provision 52.204–7). The burden for the SAM information collection is covered under OMB Control No. 9000–0189, Certain Federal Acquisition Regulation Part 4 Requirements.

This collection of information is required to comply with section 2 of the Federal Funding Accountability and Transparency Act of 2006 (Pub. L. 109–282) (FFATA), as amended by section 6202 of the Government Funding Transparency Act of 2008 (Pub. L. 110–252). The statute required the Office of Management and Budget to establish a free, public, online database containing full disclosure of all Federal contract award information. The public may view first-tier subcontract award data at *usaspending.gov.*

C. Annual Burden

Respondents: 42,231.

Total Annual Responses: 266,169.

Total Burden Hours: 506,617.

Obtaining Copies: Requesters may obtain a copy of the information collection documents from the GSA Regulatory Secretariat Division by calling 202–501–4755 or emailing *GSARegSec@gsa.gov.* Please cite OMB Control No. 9000–0177, Reporting Executive Compensation and First-tier Subcontract Awards.

Janet Fry,

Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.

[FR Doc. 2024–11523 Filed 5–23–24; 8:45 am]

BILLING CODE 6820–EP–P

GENERAL SERVICES ADMINISTRATION

[Notice-ME–2024–02; Docket No. 2024–0002; Sequence No. 26]

Notice of General Services Administration's Office of Government-Wide Policy Annual IT Modernization Summit AGENCY: Office of Government-Wide Policy (OGP), General Services Administration (GSA).

ACTION: Notice of hybrid summit.

SUMMARY: GSA, in partnership with the Information Technology Industry Council (ITI), is hosting the Annual IT Modernization Summit to bring the federal and industry communities together for a series of panel discussions covering a broad spectrum of topics such as IPv6, infrastructure, artificial intelligence, and advanced computing.

DATES: Tuesday, June 11, from 9:00 a.m. to 4:00 p.m., Eastern Time (ET).

ADDRESSES: All attendees, including industry partners, must register for the event here: *https://gsa.zoomgov.com/webinar/register/2817141347446/WN_wqOmHBZcQ0-DiER5s7ZG2Q.*

Members of the press are invited to attend but are required to register with the GSA Press Office via email to *press@gsa.gov* by Friday, June 7, 2024.

FOR FURTHER INFORMATION CONTACT: GSA's IT Modernization Division(*dccoi@gsa.gov*) and/or Tom Santucci *thomas.santucci@gsa.gov 202–230–4822.*

SUPPLEMENTARY INFORMATION:

Background

The current policy climate indicates the federal government is taking concrete steps to solidify the United States as a leader in IT modernization. This is evident with policy releases such as M–21–07 and M–24–10.

It is widely recognized that full transition to IPv6 is the only viable option to ensure future growth and innovation in internet technology and services. The federal government must expand and enhance its strategic commitment to the transition to IPv6 in order to keep pace with and capitalize on industry trends. The Office of Management and Budget's (OMB) M–21–07 ''Completing the Transition to IPv6'' provides key milestone dates for agencies to transition into IPv6-only technology stacks. In FY24, agencies are expected to reach 50% IPv6-only network environments. The Federal IPv6 Task Force has been helping agencies overcome challenges and identify best practices for achieving a full transition to IPv6.

Artificial Intelligence (AI) is making waves in both the private and public sectors as one of the most powerful technologies available. With the release of M–24–10 ''Advancing Governance, Innovation, and Risk Management for Agency Use of Artificial Intelligence,'' OMB outlines the expectations for agencies to strengthen AI governance, advance responsible AI innovation, and manage the risks from the use of AI.

Format

The annual IT Modernization Summit convenes leaders from the federal government and industry to discuss their experiences with federal IT modernization. The summit will include keynote speakers, panel discussions, and Fireside Chats with featured speakers.

If you have questions for the panelists, you can email them to *dccoi@gsa.gov.*

Special Accommodations

ASL Interpreter will be in attendance.

Zoom will have the option to enable closed captioning. If additional accommodations are needed, please note them on the Zoom Webinar registration form.

Live In-Person and Webinar Speakers (Subject to Change Without Notice)

Hosted by:

- Tom Santucci, *Director, IT Modernization; Office of Government-wide Policy*
- Robert Sears, *Director, N-Wave, National Oceanic and Atmospheric Administration; Office of Chief Information Officer (CIO)*
- Gordon Bitko, Executive Vice President of Policy, Public Sector, Information Technology Industry Council *(ITIC)*

Agenda Topic Areas

- Past to Present to Future of Information Technology
- Power of AI and Data Centers
- IPv6 and the Cloud

- High Performance Computing (HPC)

Thomas Santucci,

Director, Division of IT Modernization, Office of Technology Policy, Office of Government-wide Policy, General Services Administration.

[FR Doc. 2024–11436 Filed 5–23–24; 8:45 am]

BILLING CODE 6820–68–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[Docket No. ATSDR–2024–0001]

Availability of Three Draft Toxicological Profiles

Correction

In notice document 2024–09662, appearing on pages 36820 through 36821 in the issue of Friday, May 3, 2024, make the following correction:

On page 36820, in the first column, in the **DATES** section, on the second line, ''May 3, 2024'' should read ''August 1, 2024''.

[FR Doc. C1–2024–09662 Filed 5–23–24; 8:45 am]

BILLING CODE 0099–10–D

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2024–0043]

Meeting of the Advisory Committee on Immunization Practices

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

ACTION: Notice 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.

DATES: The meeting will be held on June 26, 2024, from 8 a.m. to 5:10 p.m., EDT, June 27, 2024, from 8 a.m. to 5:15 p.m., EDT, and June 28, 2024, from 8:30 a.m. to 12 p.m., EDT (times subject to change); see the ACIP website for updates: *https://www.cdc.gov/vaccines/acip/index.html*).

Written comments must be received between June 3–17, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0043, by either of the methods listed below. CDC does not accept comments by email.

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Ms. Stephanie Thomas, ACIP Meeting, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24–8, Atlanta, Georgia 30329–4027. Attn: Docket No. CDC–2024–0043.

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.*

The meeting will be webcast live via the World Wide Web. The webcast link can be found on the ACIP website at *https://www.cdc.gov/vaccines/acip/index.html.*

FOR FURTHER INFORMATION CONTACT: Stephanie Thomas, Committee Management Specialist, Advisory Committee on Immunization Practices, National Center for Immunization and Respiratory Diseases, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H24–8, Atlanta, Georgia 30329–4027. Telephone: (404) 639–8836; Email: *ACIP@cdc.gov.*

SUPPLEMENTARY INFORMATION:

Purpose: The Advisory Committee on Immunization Practices (ACIP) is charged with advising the Director, Centers for Disease Control and Prevention (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 applicable provisions of the Affordable Care Act and Section 2713 of the Public Health Service Act, immunization recommendations of ACIP that have been approved by the Director, CDC, and appear on CDC immunization schedules generally must be covered by applicable health plans.

Matters to be Considered: The agenda will include discussions on influenza vaccines, chikungunya vaccine, COVID–19 vaccines, pneumococcal vaccines, meningococcal vaccines, Human papillomavirus (HPV) vaccines, Respiratory Syncytial Virus (RSV) vaccines for adults, RSV vaccines for maternal and pediatric populations, a combined diphtheria and tetanus toxoids and acellular pertussis, inactivated poliovirus, Haemophilus influenzae type B conjugate, and hepatitis B vaccine (Vaxelis®), and dengue vaccine. Recommendation votes are scheduled for influenza vaccines, COVID–19 vaccines, pneumococcal vaccine, Vaxelis®, and RSV vaccines for adults. A Vaccines for Children (VFC) vote is scheduled for Vaxelis®. Agenda items are subject to change as priorities dictate. For more information on the meeting agenda, visit *https://www.cdc.gov/vaccines/acip/meetings/index.html.*

Meeting Information: The meeting will be webcast live via the World Wide Web. For more information on ACIP, please visit the ACIP website: *https://www.cdc.gov/vaccines/acip/index.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 June 3–17, 2024. Written comments must be received by June 17, 2024.

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