

approximately two to three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

#### SUPPLEMENTARY INFORMATION:

##### A. Purpose

The General Services Administration Acquisition Regulation (GSAR) clause 552.238–81 Modifications requires vendors to request a contract modification by submitting a request to the Contracting Officer for approval, except for electronic File updates. At a minimum, every request shall describe the proposed change(s) and provide the rationale for the requested change(s).

##### B. Annual Reporting Burden

*Respondents:* 14,376.

*Responses per Respondent:* 2.

*Total Responses:* 28,752.

*Hours per Response:* 3.5.

*Total Burden Hours:* 100,632.

##### C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary and whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected.

##### Obtaining Copies of Proposals

Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405; telephone 202–501–4755. Please cite OMB Control No. 3090–0302, "Modifications" in all correspondence.

**Jeffrey A. Koses,**

*Director, Office of Acquisition Policy, Office of Government-wide Policy.*

[FR Doc. 2019–07066 Filed 4–9–19; 8:45 am]

**BILLING CODE 6820–61–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—SIP19–004, Cancer Prevention and Control Research Network Coordinating Center and SIP19–005, Cancer Prevention and Control Research Network Collaborating Center; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—SIP19–004, Cancer Prevention and Control Research Network Coordinating Center and SIP19–005, Cancer Prevention and Control Research Network Collaborating Center; April 30, 2019 and May 1, 2019; 10:00 a.m.–6:30 p.m., EDT which was published in the **Federal Register** on Wednesday, February 13, 2019, Volume 84, Number 30, pages 3785.

The meeting is being amended to read as follows: Meeting date April 30, 2019. The meeting is closed to the public.

**FOR FURTHER INFORMATION CONTACT:** Jaya Raman, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway, Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488–6511, [kva5@cdc.gov](mailto:kva5@cdc.gov).

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.

**Sherri Berger,**

*Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2019–07062 Filed 4–9–19; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C.,

as amended, and the Determination of the Chief Operating Officer, CDC, pursuant to Public Law 92–463. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

*Name of Committee:* Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP)—RFA–OH–19–004, Commercial Fishing Occupational Safety Research Cooperative Agreement; RFA–OH–19–005, Commercial Fishing Occupational Training Project Grants.

*Date:* June 24–26, 2019.

*Time:* 8:00 a.m.–5:00 p.m. EDT.

*Place:* Virtual Meeting.

*Agenda:* To review and evaluate grant applications.

*For Further Information Contact:* Nina Turner, Ph.D., Scientific Review Officer, Office of Extramural Programs, 1095 Willowdale Road, Morgantown, West Virginia 26506, (304) 285–5976, [nxt2@cdc.gov](mailto:nxt2@cdc.gov).

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**Sherri Berger,**

*Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2019–07063 Filed 4–9–19; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[Docket No. CDC–2019–0028]

#### Advisory Committee on Immunization Practices (ACIP); Notice of Meeting and Request for Comment

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

**ACTION:** Notice of meeting.

**SUMMARY:** In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting of the Advisory Committee on Immunization Practices (ACIP). This meeting is open to the public, limited only by room seating. The meeting room accommodates 216 for public seating. Room 245, adjacent to the meeting

room, will be available once the meeting room reaches capacity, providing up to 18 additional seats. Time will be available for public comment. The meeting will be webcast live via the World Wide Web; for meeting registration and more information on ACIP please visit the ACIP website: <http://www.cdc.gov/vaccines/acip/index.html>.

**DATES:** The meeting will be held on June 26, 2019 8:00 a.m. to 5:30 p.m. and June 27, 2019 8:00 a.m. to 4:00 p.m. EDT. *Written comments must be received on or before June 29, 2019.*

**ADDRESSES:** You may submit comments, identified by Docket No. CDC-2019-0028 by any of the following methods:

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

- *Mail:* Centers for Disease Control and Prevention, 1600 Clifton Rd. NE, Mailstop A-27, Atlanta, GA 30329-4027

*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 <http://regulations.gov>, including any personal information provided. For access to the docket to read background documents or comments received, go to <http://www.regulations.gov>. Written public comments submitted by 72 hours prior to the ACIP meeting will be provided to the ACIP members before the meeting.

*Meeting location:* Centers for Disease Control and Prevention, 1600 Clifton Road NE, Tom Harkin Global Communications Center, Building 19, Kent 'Oz' Nelson Auditorium, Atlanta, Georgia 30329-4027.

**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, Atlanta, GA 30329-4027, telephone 404-639-8367, email [ACIP@cdc.gov](mailto:ACIP@cdc.gov).

**SUPPLEMENTARY INFORMATION:**

*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 (VFC) 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 Director of the Centers for Disease Control and Prevention and appear on CDC immunization schedules must be covered by applicable health plans.

**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 review and consider all comments submitted to the docket.

*Oral Public Comment:* This meeting will include time for members of the public to make an in-person 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. On-site, in-person registration for oral public comment at the meeting will only be available if there is time remaining in the oral public comment session after all individuals who submitted a request to make an oral comment before the meeting have had an opportunity to speak. There is no guarantee there will be an opportunity for on-site, in-person registration for oral public comment, and all individuals interested in requesting to make an oral public comment are strongly encouraged to submit a request according to the instructions below.

*Procedure for Oral Public Comment:* All persons interested in making an oral public comment at the June ACIP meeting must submit a request at <https://www.cdc.gov/vaccines/acip/meetings/> no later than 11:59 p.m., EDT,

June 12, 2019 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 each scheduled public comment session. CDC staff will notify individuals regarding their request to speak by email by June 14, 2019. 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.

*Written Public Comment:* Written comments must be received on or before June 29, 2019.

*Matters To Be Considered:* The agenda will include discussions on human papillomavirus vaccines, pneumococcal vaccines, influenza vaccines, meningococcal vaccines, hepatitis A vaccines, combination vaccines (diphtheria, tetanus, and pertussis; *Haemophilus influenzae* type b, hepatitis B, polio) dengue vaccine, pertussis vaccines, and herpes zoster vaccine. A recommendation vote is scheduled for human papillomavirus vaccines, pneumococcal vaccines, influenza vaccines, meningococcal vaccines, and hepatitis vaccines. A Vaccines for Children recommendation vote is scheduled for diphtheria, tetanus, and pertussis vaccines, *Haemophilus influenzae* type b vaccines, polio vaccines, hepatitis B vaccines, influenza vaccines, meningococcal vaccines, and hepatitis A vaccines. 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>.

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**Sherri A. Berger,**

*Chief Operating Officer, Centers for Disease Control and Prevention.*

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