

members of the Committee (when additional voting members are added to the Committee to provide needed expertise, a quorum will be based on the combined total of regular and added members) or (2) to comprise a quorum when, because of unforeseen circumstances, a quorum is or will be lacking. Because of the size of the Committee and the variety in the types of issues that it will consider, FDA may, in connection with a particular committee meeting, specify a quorum that is less than a majority of the current voting members. The Agency's regulations (21 CFR 14.22(d)) authorize a committee charter to specify quorum requirements.

If functioning as a medical device panel, a non-voting representative of consumer interests and a non-voting representative of industry interests will be included in addition to the voting members.

Further information regarding the most recent charter and other information can be found at <https://www.fda.gov/advisory-committees/vaccines-and-related-biological-products-advisory-committee/charter-vaccines-and-related-biological-products-advisory-committee> or by contacting the Designated Federal Officer (see **FOR FURTHER INFORMATION CONTACT**). In light of the fact that no change has been made to the committee name or description of duties, no amendment will be made to 21 CFR 14.100.

This document is issued under the Federal Advisory Committee Act (5 U.S.C. app.). For general information related to FDA advisory committees, please visit us at <https://www.fda.gov/AdvisoryCommittees/default.htm>.

Dated: January 24, 2022.

**Lauren K. Roth,**

*Associate Commissioner for Policy.*

[FR Doc. 2022-01858 Filed 1-28-22; 8:45 am]

**BILLING CODE 4164-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

#### Meeting of the Advisory Committee on Childhood Vaccines

**AGENCY:** Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** In accordance with the Federal Advisory Committee Act, this notice announces that the Advisory

Commission on Childhood Vaccines (ACCV) will hold public meetings for the 2022 calendar year (CY).

Information about the ACCV, agendas, and materials for these meetings can be found on the ACCV website at <https://www.hrsa.gov/advisory-committees/vaccines/index.html>.

**DATES:** ACCV meetings will be held on:

- March 3, 2022, 10:00 a.m. Eastern Time (ET)–4:00 p.m. ET;
- June 2, 2022, 10:00 a.m. ET–4:00 p.m. ET;
- September 1, 2022, 10:00 a.m. ET–4:00 p.m. ET; and
- December 1, 2022, 10:00 a.m. ET–4:00 p.m. ET.

**ADDRESSES:** Meetings may be held in-person or virtually. For updates on how the meeting will be held, visit the ACCV website 30 business days before the meeting date, where instructions for joining meetings either in-person or remotely will be posted. In-person ACCV meetings will be held at 5600 Fishers Lane, Rockville, Maryland 20857. For meeting information updates, go to the ACCV website meeting page at <https://www.hrsa.gov/advisory-committees/vaccines/meetings.html>.

#### FOR FURTHER INFORMATION CONTACT:

Annie Herzog, Division of Injury Compensation Programs, HRSA, 5600 Fishers Lane, 08N186B, Rockville, Maryland 20857; 301-443-6634; or [ACCV@HRSA.gov](mailto:ACCV@HRSA.gov).

**SUPPLEMENTARY INFORMATION:** The ACCV provides advice and recommendations to the Secretary of HHS on policy, program development, and other issues related to the implementation of the National Vaccine Injury Compensation Program and concerning other matters as described under section 2119 of the Public Health Service Act (42 U.S.C. 300aa-19).

Since priorities dictate meeting times, be advised that times and agenda items are subject to change. Refer to the ACCV website listed above for any meeting updates that may occur. For CY 2022 meetings, agenda items may include, but are not limited to: Updates from the Division of Injury Compensation Programs, Department of Justice, Office of Infectious Disease and HIV/AIDS Policy (HHS), Immunization Safety Office (Centers for Disease Control and Prevention), National Institute of Allergy and Infectious Diseases (National Institutes of Health) and Center for Biologics, Evaluation and Research (Food and Drug Administration). Refer to the ACCV website listed above for all current and updated information concerning the CY 2022 ACCV meetings, including draft

agendas and meeting materials posted 5 calendar days before the meeting(s).

Members of the public will have the opportunity to provide comments. Public participants may submit written statements in advance of the scheduled meeting(s). Oral comments will be honored in the requested order and may be limited as time allows. Requests to submit a written statement or make oral comments to ACCV should be sent to Annie Herzog using the contact information above at least 5 business days before the meeting date(s).

Individuals who need special assistance or another reasonable accommodation should notify Annie Herzog using the contact information listed above at least 10 business days before the meeting(s) they wish to attend. If in-person meetings occur, they will be held in a federal government building and attendees must go through a security check to enter the building. Non-U.S. Citizen attendees must notify HRSA of their planned attendance at least 20 business days before the meeting to facilitate their entry into the building. All attendees are required to present government-issued identification before entry.

**Maria G. Button,**

*Director, Executive Secretariat.*

[FR Doc. 2022-01848 Filed 1-28-22; 8:45 am]

**BILLING CODE 4165-15-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Center for Scientific Review; Notice of Closed Meetings

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

The meetings 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. 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:* Biobehavioral and Behavioral Processes Integrated Review Group; Biobehavioral Mechanisms of Emotion, Stress and Health Study Section.

*Date:* February 24–25, 2022.

*Time:* 10:00 a.m. to 8:00 p.m.