ESTIMATED REALLOTMENT AMOUNTS OF FY 2018 LIHEAP FUNDS

| Grantee name | Reallotment amount |
|---|---|
| Alaska | $1,579,924 |
| Five Sandoval Indian Pueblos, INC | 16,089 |
| Hoh Indian Tribe | 4,378 |
| Little River Band of Ottawa Indians | 47,440 |
| Northern Cheyenne Tribe | 45,607 |
| Three Affiliated Tribes | 140,582 |
| Turtle Mountain Band of Chippewa Indians | 5,108 |
| Total | $1,839,128 |

**Statutory Authority:** 42 U.S.C. 8626.

**Elizabeth Leo,**

*Senior Grants Policy Specialist, Division of Grants Policy, Office of Administration.*

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**BILLING CODE 4184–80–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2019–N–3612]**

### Vaccines and Related Biological Products Advisory Committee; Notice of Meeting

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or the Agency) announces a forthcoming public advisory committee meeting of the Vaccines and Related Biological Products Advisory Committee (VRBPAC). The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. At least one portion of the meeting will be closed to the public. Members will participate via teleconference.

**DATES:** The meeting will be held on October 9, 2019, from 8:30 a.m. to 3:30 p.m.

**ADDRESSES:** FDA White Oak Campus, 10903 New Hampshire Avenue, Bldg. 31 Conference Center, the Great Room (Rm. 1503), Silver Spring, MD 20993–0002. For those unable to attend in person, the meeting will also be webcast and will be available at the following link: *https://collaboration.fda.gov/vrbpac100919/.* Answers to commonly asked questions including information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

**FOR FURTHER INFORMATION CONTACT:** Capt. Serina Hunter-Thomas, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 6338, Silver Spring, MD 20993–0002, 240–402–5771, *serina.hunter-thomas@fda.hhs.gov,* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the Agency's website at *https://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

**SUPPLEMENTARY INFORMATION:**

*Agenda:* On October 9, 2019, under topic I, the Center for Biologics Evaluation and Research's (CBER) VRBPAC will meet in open session to hear an overview of the research programs in the Laboratory of Hepatitis Viruses (LIR) and the Laboratory of Vector-Borne Viral Diseases (LVVD), Division of Viral Products, Office of Vaccines Research and Review, CBER, FDA. Also, on October 9, 2019, under topic II, the committee will meet in open session to discuss and make recommendations on the selection of strains to be included in an influenza virus vaccine for the 2020 southern hemisphere influenza season.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's website after the meeting. Background material is available at *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link.

*Procedure:* On October 9, 2019, from 8:30 a.m. to approximately 10 a.m. and from 11 a.m. to 3:30 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person on or before October 2, 2019. Oral presentations from the public will be scheduled between approximately 9:45 a.m. and 10 a.m. for the overview portion of the LHV/LVVD Site Visit (topic I), and from 1:30 p.m. to 2:15 p.m. for the influenza strain selection portion of the meeting (topic II). Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation on or before September 24, 2019. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by September 25, 2019.

Persons attending FDA's advisory committee meetings are advised that the Agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Capt. Serina Hunter-Thomas at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our website at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: August 16, 2019.

**Lowell J. Schiller,**

*Principal Associate Commissioner for Policy.*

[FR Doc. 2019–18410 Filed 8–26–19; 8:45 am]

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