

The USPSTF was first established in 1984 under the auspices of the U.S. Public Health Service. AHRQ provides ongoing scientific, administrative, and dissemination support for the USPSTF's operation. See 42 U.S.C. 299b-4(a)(3). Members are appointed by the Secretary of the U.S. Department of Health and Human Services to serve four-year terms. New members are selected each year to replace those members who are completing their appointments.

The USPSTF rigorously evaluates the effectiveness of clinical preventive services and formulating or updating recommendations regarding the appropriate provision of preventive services. Current USPSTF recommendations and associated evidence reviews are available on the internet (www.uspreventiveservicestaskforce.org).

USPSTF members meet three times a year for two days in the Washington, DC area or virtually if necessary. A significant portion of the USPSTF's work occurs between meetings during video conference calls and via email discussions. Member duties include prioritizing topics, designing research plans, reviewing and commenting on systematic evidence review reports, discussing evidence and making recommendations on preventive services, reviewing stakeholder comments, drafting final recommendation documents, and participating in workgroups on specific topics and methods. Members can expect to receive frequent emails, participate in multiple video conference calls each month, and have periodic interaction with stakeholders. AHRQ estimates that members devote approximately 250 hours a year outside of in-person meetings to their USPSTF duties. The members are all volunteers and do not receive any compensation beyond support for travel to attend the thrice yearly meetings and trainings.

Dated: April 21, 2026.

Roger D. Klein,

Director.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2026-N-2366]

Commissioner's National Priority Voucher (CNPV) Pilot Program; Public Hearing; Request for Comments; Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an amendment to the notice of the public hearing related to the Commissioner's National Priority Voucher (CNPV) Pilot Program. This meeting was announced in the **Federal Register** of March 23, 2026. The amendment is being made to reflect a change in the **DATES, ADDRESSES, FOR FURTHER INFORMATION CONTACT,** and *IV. Participating in Public Hearing* portions of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Mallika Mundkur, Deputy Chief Medical Officer, Office of the Commissioner, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301-796-8800, CNPVPublicMeeting@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of March 23, 2026, 91 FR 13849, FDA announced that a public hearing related to the CNPV Pilot Program would be held on June 12, 2026. On page 13849, in the third column, the **DATES** portion of the document is changed to read as follows:

- The public hearing will be held with an in-person and virtual option (*i.e.*, hybrid) on June 4, 2026, from 1:00 p.m. to 4:00 p.m. Eastern Time. Meeting registration, including requests for participation in the public hearing, can be found at the following website: <https://www.fda.gov/news-events/fda-meetings-conferences-and-workshops/commissioners-national-priority-voucher-cnpv-pilot-program-public-hearing-06042026>. All requests for participation, including those who wish to present during the public hearing, must be received by April 24, 2026, through the meeting registration page. Questions about meeting registration and participation should be sent to CNPVPublicMeeting@fda.hhs.gov and include the title of this notice: "Commissioner's National Priority Voucher (CNPV) Pilot Program; Public Hearing; Request for Comments").

On page 13850, in the first column, the second and third paragraphs of the **ADDRESSES** portion of the document are changed to read as follows:

- Additional details, such as any changes to the time of the public hearing and registration information, will be posted at <https://www.fda.gov/news-events/fda-meetings-conferences-and-workshops/commissioners-national-priority-voucher-cnpv-pilot-program-public-hearing-06042026>. The online web conference meeting link can be accessed at <https://www.fda.gov/news-events/fda-meetings-conferences-and-workshops/commissioners-national-priority-voucher-cnpv-pilot-program-public-hearing-06042026> on the day of the meeting.

- All written requests for participation in the pilot program must be received by April 24, 2026 (email to: CNPVPublicMeeting@fda.hhs.gov).

On page 13850, in the third column, the **FOR FURTHER INFORMATION CONTACT** portion of the document is changed to read as follows:

- Mallika Mundkur, Deputy Chief Medical Officer, Office of the Commissioner, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301-796-8800, CNPVPublicMeeting@fda.hhs.gov.

On page 13852, in the second column, the *IV. Participating in Public Hearing* portion of the **SUPPLEMENTARY INFORMATION** section is changed to read as follows:

- **Registration:** To register to attend or participate in the free public hearing, please visit the following website: <https://www.fda.gov/news-events/fda-meetings-conferences-and-workshops/commissioners-national-priority-voucher-cnpv-pilot-program-public-hearing-06042026>.

- **Written Notice of Participation:** All written requests for participation must be received by April 24, 2026, 11:59 p.m. Eastern Time (email to: CNPVPublicMeeting@fda.hhs.gov).

- **Transcripts:** A link to the transcript will also be available on the internet at <https://www.fda.gov/news-events/fda-meetings-conferences-and-workshops/commissioners-national-priority-voucher-cnpv-pilot-program-public-hearing-06042026>.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

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