

your request. If you are requesting a change to your address, please be sure to include your name and the correct address. If you are requesting to delete your address from the mailing list, please include your name and address as it appeared on this notice. This email address is unable to accept comments.

OR
(2) Return the attached "Mailing List Update Form" (appendix 3).

Additional Information

Additional information about the project is available from the FERC website at www.ferc.gov using the eLibrary link. Click on the eLibrary link, click on "General Search" and enter the docket number in the "Docket Number" field. Be sure you have selected an appropriate date range. For assistance, please contact FERC Online Support at FercOnlineSupport@ferc.gov or (866) 208-3676, or for TTY, contact (202) 502-8659. The eLibrary link also provides access to the texts of all formal documents issued by the Commission, such as orders, notices, and rulemakings.

Public sessions or site visits will be posted on the Commission's calendar located at <https://www.ferc.gov/news-events/events/events> along with other related information.

(Authority: 18 CFR 2.1)

Dated: October 13, 2026

Carlos D. Clay,

Deputy Secretary.

[FR Doc. 2026-21191 Filed 10-15-26; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[FRL OPRM-FAD-244]

Environmental Impact Statements; Notice of Availability

Responsible Agency: Office of Federal Activities, General Information 202-993-3272 or <https://www.epa.gov/nepa>. Weekly receipt of Environmental Impact Statements (EIS)

Filed October 5, 2026 10 a.m. EST

Through October 13, 2026 10 a.m. EST

Pursuant to CEQ Guidance on 42 U.S.C. 4332.

Notice

Section 309(a) of the Clean Air Act requires that EPA make public its comments on EISs issued by other Federal agencies. EPA's comment letters on EISs are available at: <https://cdxapps.epa.gov/cdx-enepa-II/public/action/eis/search>.

EIS No. 20260132, Final, USFS, AK, Twin Mountain II Timber Sale, Review Period Ends: 11/30/2026, Contact: Christopher Dowling, 907-828-3304.

Dated: October 13, 2026

Nancy Abrams,

Deputy Director, Federal Activities Division.

[FR Doc. 2026-21183 Filed 10-15-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Meeting of the Advisory Council for the Elimination of Tuberculosis

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 Centers for Disease Control and Prevention (CDC) announces the following meeting of the Advisory Council for the Elimination of Tuberculosis (ACET). This meeting is open to the public, limited only by the number of audio and web conference lines (1,000 lines are available). Time will be available for public comment (registration is required to provide oral comment; see the Oral Public Comment section below).

DATES: The meeting will be held on December 2, 2026, from 12:00 p.m. to 4:30 p.m., EST, and on December 3, 2026, from 12:00 p.m. to 4 p.m., EST.

Written comments must be submitted by November 25, 2026. Registration to make oral comments must also be submitted by November 25, 2026.

ADDRESSES: The public meeting will be held virtually through Microsoft Teams. Advance registration is required to attend. Please register for each day of this meeting.

For registration on December 2, 2026: <https://events.gcc.teams.microsoft.com/event/ac21938b-f766-40e2-bf20-420cf442870f@9ce70869-60db-44fd-abe8-d2767077fc8f?source=copyLink>
<https://events.gcc.teams.microsoft.com/event/126b55ca-74b3-4907-857d-2709a2bb82b7@9ce70869-60db-44fd-abe8-d2767077fc8f?source=copyLink>
 LegacyShareLinkDialog. For registration on December 3, 2026: <https://events.gcc.teams.microsoft.com/event/126b55ca-74b3-4907-857d-2709a2bb82b7@9ce70869-60db-44fd-abe8-d2767077fc8f?source=copyLink>
 LegacyShareLinkDialog. Registration for virtual attendance will remain open through the meeting. Prior to the meeting, each individual

registrant will receive a registration confirmation along with an access link to the virtual meeting location. Written public comments and requests to make oral comments should be sent to nchhstppolicy@cdc.gov.

FOR FURTHER INFORMATION CONTACT:

ACET Committee Management, Office of Policy, Planning, and Partnerships, National Center for HIV, Viral Hepatitis, STD, and TB Prevention, Centers for Disease Control and Prevention. Email: nchhstppolicy@cdc.gov.

SUPPLEMENTARY INFORMATION:

Purpose: The Advisory Council for the Elimination of Tuberculosis is charged with providing advice and recommendations regarding the elimination of tuberculosis (TB) to the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, Centers for Disease Control and Prevention (CDC). Specifically, the Council makes recommendations regarding policies, strategies, objectives, and priorities; addresses the development and application of new technologies; provides guidance and review on CDC's Tuberculosis Prevention Research portfolio and program priorities; and reviews the extent to which progress has been made toward eliminating TB.

Matters to be Considered: The agenda will include discussions on: (1) CDC's National Center for HIV, Viral Hepatitis, STD, and TB Prevention Update; (2) CDC's Division of Tuberculosis Elimination Update; (3) Reported TB in the United States, 2025; (4) TB Diagnostics Rapid Acceleration of Diagnostics Tech initiative; (5) TB Workforce Update; (6) Tuberculosis Trials Consortium Update; (7) Study 38/CRUSH-TB Clinical Trial; and (8) Post-Tuberculosis Lung Disease and Mortality. Agenda items are subject to change as priorities dictate.

Public Participation

Written Public Comment: Members of the public are welcome to submit written comments in advance of the meeting. Written comments must be submitted by emailing nchhstppolicy@cdc.gov with subject line "ACET December 2026 Written Public Comment Registration" by November 25, 2026.

Oral Public Comment: Individuals who would like to make an oral comment during the public comment period must register by emailing nchhstppolicy@cdc.gov with subject line "ACET December 2026 Oral Public Comment Registration" by November 25, 2026. The public comment period is on December 3, 2026, at 2 p.m., EST.

Comments are limited to no more than 5 minutes each. If the number of persons requesting to speak is greater than can be reasonably accommodated during the scheduled time, CDC will conduct a random draw to determine the speakers for the scheduled public comment session. CDC staff will notify individuals regarding their request to speak by email by November 30, 2026 at 5 p.m. EST.

The Director, Office of Strategic Business Initiatives, Office of 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.

Kalwant Smagh,

Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2026-21125 Filed 10-15-26; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-10143]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and

clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by December 15, 2026.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier: CMS-10143/OMB Control Number: 0938-0958, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

FOR FURTHER INFORMATION CONTACT: William N. Parham at (410) 786-4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collection. More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

Under the PRA (44 U.S.C. 3501-3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term "collection of information" is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed

extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

Information Collections

1. Type of Information Collection

Request: Revision of a currently approved collection; *Title of Information Collection:* State Data for the Medicare Modernization Act (MMA); *Use:* The Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (Public Law 108-173) requires CMS to calculate the payment rates for the Phased-Down State Contribution (PDSC) for Part D each year using the latest available National Health Expenditure (NHE) estimates of per capita drug expenditure growth for the period 2003 to 2006, combined with the annual percentage increase (API) in average per capita aggregate Part D expenditures for 2007 and later years, as defined in section 1860D-2(b)(6) of the Social Security Act.

Section 103(a)(2) of the MMA addresses the PDSC process for the Medicare program. The PDSC is a partial recoupment from the states of ongoing Medicaid drug costs for dually eligible beneficiaries assumed by Medicare under the MMA, which (absent the MMA) would have been paid for by the states.

As CMS leverages MMA data on dual eligibility status into systems supporting all four parts of the Medicare program (which includes Original Medicare, Medicare Advantage, and Part D), it is important that dual eligibility status is accurate and up to date. Dual eligibility status can change at any time in a month. Daily status updates mitigate the risk of a lack of access to the correct level of benefit at the correct level of payment. Daily data exchange is critical to timely access to affordable benefits; without daily exchanges, CMS would experience a lag in its ability to automatically enroll these individuals in Medicare drug plans or deem them automatically eligible for the low-income subsidy for Part D premiums, deductibles, and copayments.

Since 2005, states have been submitting files at least monthly to CMS to identify all dually eligible beneficiaries. This includes full-benefit dually eligible beneficiaries and partial-benefit dually eligible beneficiaries (*i.e.*, those who get Medicaid help with Medicare premiums, and often for cost-sharing but are not otherwise eligible for Medicaid-covered services). The file is called the "MMA file" (after the Medicare Prescription Drug,