

## 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 consider all comments submitted into the docket.

**Written Public Comment:** The docket will be opened to receive written comments September 2–13, 2025. Written comments must be received no later than September 13, 2025.

**Oral Public Comment:** This meeting will include time for members of the public to make an 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.

**Procedure for Oral Public Comment:** All persons interested in making an oral public comment at the September 18–19, 2025, ACIP meeting must submit a request at <https://www.cdc.gov/acip/meetings/index.html> between September 2–13, 2025, and no later than 11:59 p.m., EDT, September 13, 2025, 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 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 September 16, 2025. To accommodate the significant interest in participation in the oral public comment session of ACIP meetings, each speaker will be limited to three

minutes, and each speaker may speak only once per meeting.

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. 2025–16706 Filed 8–28–25; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10495 and CMS–855S]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, HHS.

**ACTION:** Notice; partial withdrawal.

**SUMMARY:** On Tuesday, August 5, 2025, the Centers for Medicare & Medicaid Services (CMS) published a notice document entitled, “Agency Information Collection Activities: Proposed Collection; Comment Request”. That notice invited public comments on five separate information collection requests, under Document Identifiers: CMS–10495, CMS 855S and CMS–R–131. Through the publication of this document, we are withdrawing the portion of the notice requesting public comment on the information collection request titled, “Data Collection and Submission for Open Payments” Form number: CMS–10495 (OMB control number: 0938–1237). We are also withdrawing the portion of the notice requesting public comment on the information collection titled, “Medicare Enrollment Application: Durable Medical Equipment, Prosthetics, Orthotics, and Supplies (DMEPOS) Suppliers.” Form number: CMS–855S (OMB control number: 0938–1056).

**DATES:** The original comment period for the document that published on August 5, 2025, remains in effect and ends October 6, 2025.

**SUPPLEMENTARY INFORMATION:** In FR document, 2025–14828, published on August 5, 2025 (90 FR 37515), we are withdrawing item 1 “Registration, Attestation, Dispute Resolution and Correction, Assumptions Document and Data Retention Requirements for Open Payments” which begins at the top of the left column on page 37516. We are also withdrawing item 2 “Medicare Enrollment Application—Durable Medical Equipment, Prosthetics, Orthotics and Supplies (DMEPOS) Suppliers,” which begins at the top of the middle column on page 37516.

**William N. Parham, III,**

*Director, Division of Information Collections and Regulatory Impacts, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2025–16572 Filed 8–28–25; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS–R–70, CMS–R–72 and CMS–10781]

#### 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 October 28, 2025.