publishing this notice that summarizes the following proposed collection(s) of information for public comment.

1. *Type of Information Collection Request:* Reinstatement with change of a previously approved information collection; *Title of Information Collection:* State-based Exchange Annual Reporting Tool (SMART); *Use:* The ACA § 1313(a)(1) and its implementing regulations require State Exchanges to keep an accurate accounting of all activities, receipts, and expenditures, and to submit a report annually to CMS concerning such accounting. Instructions governing specific facets of the activities covered by the report are contained both in the ACA and 45 CFR 155.1200, 155.1210. CMS uses the SMART as the reporting tool to ensure compliance with regulatory requirements.

CMS uses the information collected from the SMART to determine if a state is maintaining a compliant, operational Exchange. It also provides a mechanism to collect innovative approaches to meeting challenges encountered by states during the preceding year, as well as to provide information to CMS regarding potential changes in priorities and approaches for the upcoming year. If CMS determines a state to be non-compliant through the review of required documentation, it will issue a formal letter asking the state to develop and submit a Corrective Action Plan (CAP). CMS may also provide technical assistance to help State Exchanges address potential areas of non-compliance, as needed. *Form Number:* CMS–10507 (OMB control number: 0938–1244); *Frequency:* Annually; *Affected Public:* State, Local or Tribal Government; *Number of Respondents:* 23; *Number of Responses:* 23; *Total Annual Hours:* 4,792. (For questions regarding this collection, contact Tiffany Y. Animashaun at *Tiffany.Animashaun@cms.hhs.gov.*)

William N. Parham, III

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10394 and CMS–10544]

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 May 18, 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: __/OMB Control Number: __, 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 collections. More detailed information can be found in each 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 Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Application To Be a Qualified Entity To Receive Medicare Data for Performance Measurement/Reapplication/Annual Report Worksheet; *Use:* Section 10332 of the Patient Protection and Affordable Care Act (ACA) requires the Secretary to make standardized extracts of Medicare claims data under Parts A, B, and D available to ''qualified entities'' (QEs) for evaluating performance of providers of services and suppliers, implement the requirements outlined in the legislation, CMS established the Qualified Entity Certification Program (QECP) to evaluate an organization's eligibility across three areas: (1) organizational and governance capabilities, (2) addition of claims data from other sources (as

required in the statute), and (3) data privacy and security.

This collection covers the application through which organizations provide information to CMS to determine whether they will be approved as a qualified entity. This collection also covers the triennial re-application through which organizations provide information to CMS to determine whether they are approved to continue as a qualified entity and the QECP Annual Report Worksheet (ARW). *Form Number:* CMS–10394 (OMB control number: 0938–1144); *Frequency:* Yearly; *Affected Public:* Business or other for-profits; *Number of Respondents:* 40; *Total Annual Responses:* 70; *Total Annual Hours:* 5,800. (For policy questions regarding this collection contact Kari Gaare at *kari.gaare@cms.hhs.gov.*)

2. *Type of Information Collection Request:* Re-instatement without change of a previously approved collection; *Title of Information Collection:* Good Cause Processes; *Use:* Section 1851(g)(3)(B)(i) of the Act provides that MA organizations may terminate the enrollment of individuals who fail to pay basic and supplemental premiums after a grace period established by the plan. Section 1860D–1(b)(1)(B)(v) of the Act generally directs us to establish rules related to enrollment, disenrollment, and termination for Part D plan sponsors that are similar to those established for MA organizations under section 1851 of the Act. Consistent with these sections of the Act, subpart B in each of the Parts C and D regulations sets forth requirements with respect to involuntary disenrollment procedures at 42 CFR 422.74 and 423.44, respectively. In addition, section 1876(c)(3)(B) establishes that individuals may be disenrolled from coverage as specified in regulations.

These good cause provisions authorize CMS to reinstate a disenrolled individual's enrollment without interruption in coverage if the non-payment is due to circumstances that the individual could not reasonably foresee or could not control, such as an unexpected hospitalization. At its inception, the process of accepting, reviewing, and processing beneficiary requests for reinstatement for good cause was carried out exclusively by CMS. *Form Number:* CMS–10544 (OMB control number: 0938–1271); *Frequency:* Occasionally; *Affected Public:* Individuals and Households, Private Sector and Business or other for-profits; *Number of Respondents:* 54,789; *Total Annual Responses:* 54,789; *Total Annual Hours:* 36,490. (For policy questions regarding this collection contact AnhViet Nguyen at 667–290–9745 or *anhviet.nguyen@cms.hhs.gov*).

William N. Parham, III,

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

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2026–N–2279]

Preparation for International Cooperation on Cosmetics Regulation Twentieth Annual Meeting (ICCR–20); Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is soliciting comments from the public on various topics pertaining to the regulation of cosmetics in preparation for the twentieth International Cooperation on Cosmetics Regulation annual meeting (ICCR–20). We may use this input to help us prepare for the ICCR–20 meeting that will be held July 7 to 9, 2026, in Tokyo, Japan.

DATES: Either electronic or written comments on this notice must be submitted by May 18, 2026.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of May 18, 2026. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2026–N–2279 for ''Preparation for International Cooperation on Cosmetics Regulation Twentieth Annual Meeting (ICCR–20); Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly