of chemical exposure to the skin. The final publication, which will address public comments, will be available on the NIOSH website and in the NIOSH docket (153–F) and in *Regulations.gov* (CDC–2024–0085).

Background: In 2009, NIOSH published Current Intelligence Bulletin (CIB) 61: A Strategy for Assigning New NIOSH Skin Notations [NIOSH 2009]. The CIB presents a strategic framework that is a form of hazard identification designed to do the following:

- Ensure that the assigned skin notations reflect the contemporary state of scientific knowledge
- Provide transparency behind the assignment process
- Communicate the hazards of chemical exposures of the skin
- Meet the needs of health professionals, employers, and other interested parties in protecting workers from chemical contact with the skin.

This strategy involves the assignment of multiple skin notations for distinguishing systemic (SYS), direct (DIR), and sensitizing (SEN) effects caused by exposure of skin (SK) to chemicals. Chemicals that are highly or extremely toxic and may be potentially lethal or life-threatening following exposures of the skin are designated with the systemic subnotation (FATAL). Potential irritants and corrosive chemicals are indicated by the direct effects subnotations (IRR) and (COR), respectively. The five draft Skin Notation Profiles available for review were developed following the framework in NIOSH CIB 61.

Reference

NIOSH [2009]. Current Intelligence Bulletin 61: A strategy for assigning new NIOSH skin notations. Cincinnati, OH: U.S. Department of Health and Human Services, Centers for Disease Control and Prevention, National Institute for Occupational Safety and Health, DHHS (NIOSH) Publication 2009–147, *https://www.cdc.gov/niosh/docs/2009-147/.*

Dated: October 23, 2024.

John J. Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Department of Health and Human Services.

[FR Doc. 2024–24983 Filed 10–25–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10856]

Agency Information Collection Activities: Submission for OMB Review; Comment Request; Withdrawal

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

ACTION: Notice; withdrawal.

SUMMARY: On Thursday, September 26, 2024, the Centers for Medicare & Medicaid Services (CMS) published a 30-day Paperwork Reduction Act of 1995 notice entitled, ''Agency Information Collection Activities: Submission for OMB Review; Comment Request.'' The notice invited public comment on Document Identifier: CMS–10856; Title of Information Collection: Medicaid Managed Care and Supporting Regulations; and Form Number: CMS–10856 (OMB control number 0938–1453). Through the publication of this document we are withdrawing the September 26, 2024, notice in its entirety.

DATES: This withdrawal is applicable on October 28, 2024.

SUPPLEMENTARY INFORMATION: Through the publication of this notice we are withdrawing FR document 2024–21982 which published in the **Federal Register** on September 26, 2024 (89 FR 78875). Upon further review the associated collection of information request was not ready for public review and comment. The 30-day notice will republish at a date to be determined.

William N. Parham, III,

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

[FR Doc. 2024–25019 Filed 10–25–24; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10912]

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 27, 2024.

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**).

CMS–10912 Medicare Transaction Facilitator for 2026 and 2027 under Sections 11001 and 11002 of the Inflation Reduction Act (IRA) Information Collection Request

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:* New Collection; *Title of Information Collection:* Medicare Transaction Facilitator for 2026 and 2027 under Sections 11001 and 11002 of the Inflation Reduction Act (IRA) Information Collection Request; *Use:* Under the authority in sections 11001 and 11002 of the Inflation Reduction Act of 2022 (Pub. L. 117–169), the Centers for Medicare & Medicaid Services (CMS) is implementing the Medicare Drug Price Negotiation Program, codified in sections 1191 through 1198 of the Social Security Act (''the Act''). The Act establishes the Negotiation Program to negotiate maximum fair prices (''MFPs''), defined at 1191(c)(3) of the Act, for certain high expenditure, single source selected drugs covered under Medicare Part B and Part D (''selected drugs''). In accordance with section 1193(a) of the Act, any Primary Manufacturer of a selected drug that continues to participate in the Negotiation Program and reaches agreement upon an MFP must provide access to the MFP to MFP-eligible individuals, defined in section 1191(c)(2)(A) of the Act, and to pharmacies, mail order services, other dispensing entities, providers and suppliers with respect to such MFP-eligible individuals who are dispensed that selected drug during a price applicability period. The purpose of this information collection request (ICR) is for CMS to collect information from manufacturers of drugs covered under Part D selected for negotiation under the Inflation Reduction Act for the initial price applicability years 2026 and 2027 and the dispensing entities that dispense the selected drugs to MFP-eligible individuals. To facilitate the effectuation of the MFP, CMS will engage a Medicare Transaction Facilitator (''MTF''). The MTF system will be composed of two modules: the MTF Data Module (MTF DM), and the MTF Payment Module (MTF PM).

Medicare Transaction Facilitator Data Elements: The MTF system will be composed of two modules: the MTF Data Module (MTF DM), and the MTF Payment Module (MTF PM). Primary Manufacturers participating in the Negotiation Program are required to participate in the MTF DM. Further, CMS intends to propose in future rulemaking to require Part D plan sponsors to include in their pharmacy agreements provisions requiring dispensing entities to participate in the MTF DM for purposes of data exchange. As such, for the purposes of this ICR, CMS assumes full participation in the MTF DM by affected Primary Manufacturers and dispensing entities. Meanwhile, participation in the MTF PM, for use in passing through payment from the Primary Manufacturer to dispensing entities, will be optional for Primary Manufacturers; as a result, dispensing entities may receive fund transfers from the MTF PM, or via an alternative process established by a Primary Manufacturer. As discussed in section 40.4 of the Medicare Drug Price Negotiation Program: Final Guidance, Implementation of Sections 1191–1198 of the Social Security Act for Initial Price Applicability Year 2027 and Manufacturer Effectuation of the Maximum Fair Price (MFP) in 2026 and 2027 (''final guidance''), CMS will engage the MTF DM to facilitate the exchange of certain claim-level data elements and payment elements for selected drugs. The data exchange component of the MTF will involve both the transmission of certain claim-level data elements to the Primary Manufacturer and receipt of claim-level payment elements from the Primary Manufacturer. Both Primary Manufacturers and dispensing entities will need to provide certain information at the onset of their enrollment in the MTF DM system to facilitate effectuation of the MFP via refunds from Primary Manufacturers. Both Primary Manufacturers and dispensing entities will be able to submit complaints and disputes through their participation in the MTF DM. Primary Manufacturers will also submit information to fulfill their requirement to provide an MFP Effectuation Plan and transmit recurring data submissions reflecting their payment elements, as described in the final guidance. Given these information collection requirements, this ICR includes the following forms: (A) Drug Price Negotiation Program MTF DM Dispensing Entity and Third-Party Support Enrollment Form; (B) Drug Price Negotiation Program MTF DM Primary Manufacturer Maximum Fair Price (MFP) Effectuation Plan Form; (C) Drug Price Negotiation Program MTF DM Primary Manufacturer Payment Elements Form; and (D) Drug Price Negotiation Program Complaint and Dispute Intake Form. *Form Number:* CMS–10912 (OMB control number: 0938-New); *Frequency:* Once and Daily; *Affected Public:* Private sector, Business or other for-profit, and individuals; *Number of Respondents:* 85,853; *Total Annual Responses:* 93,120; *Total Annual Hours:* 821,560. (For policy questions regarding this collection contact Brennan Folsom at 667–414–0014.)

William N. Parham, III,

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

[FR Doc. 2024–25009 Filed 10–25–24; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10261]

Agency Information Collection Activities: Submission for OMB Review; 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