about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT: Katherine Sleasman, Office of Program Support (Mail Code 7602M), Office of Chemical Safety and Pollution Prevention, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: (202) 566–1204; email address: *Sleasman.Katherine@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. What information is EPA particularly interested in?

Pursuant to PRA section 3506(c)(2)(A) (44 U.S.C. 3506(c)(2)(A)), EPA specifically solicits comments and information to enable it to:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility.

2. Evaluate the accuracy of the Agency's estimates of the burden of the proposed collection of information, including the validity of the methodology and assumptions used.

3. Enhance the quality, utility, and clarity of the information to be collected.

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses. In particular, EPA is requesting comments from very small businesses (those that employ less than 25) on examples of specific additional efforts that EPA could make to reduce the paperwork burden for very small businesses affected by this collection.

II. What information collection activity or ICR does this action apply to?

Title: Methylene Chloride; Regulation under TSCA § 6(a).

EPA ICR No.: 2556.04.

OMB Control No.: 2070–0204.

ICR Status: This ICR is currently approved through May 31, 2026. Under the PRA, an agency may not conduct or sponsor, and a person is not required to respond to, a collection of information, unless it displays a currently valid OMB control number. The OMB control numbers for EPA's regulations in title 40 of the Code of Federal Regulations (CFR), after appearing in the **Federal Register** when approved, are displayed either by publication in the **Federal Register** or by other appropriate means, such as on the related collection instrument or form, if applicable. The display of OMB control numbers for certain EPA regulations is consolidated in 40 CFR part 9.

Abstract: The Environmental Protection Agency (EPA) is consolidating the rule-related ICR titled, ''Methylene Chloride; Regulation under TSCA Section 6(a) (Final Rule; RIN 2070–AK70)(EPA ICR No. 2735.02; OMB Control No. 2070–0229)'' into ''Methylene Chloride; Regulation of Paint and Coating Removal for Consumer Use under TSCA Section 6(a) (EPA ICR No. 2556.03; OMB Control No. 2070–0204).'' This ICR describes the prohibition of the manufacture, process, and distribution of methylene chloride for all consumer use and most industrial and commercial uses and delay prohibition for two conditions: a requirement for a workplace chemical protection program (WCPP) and related workplace methylene chloride monitoring under 40 CFR 751. This ICR covers the information collection activities for downstream notification requirements through Safety Data Sheets; WCPP-related information such as recordkeeping and notification requirements associated with exposure monitoring; and recordkeeping for interim requirements for use of methylene chloride for refinishing wood pieces of artistic, cultural, or historic value and downstream notification for consumer use of paints and coatings.

The ICR, which is available in the docket along with other related materials, provides a detailed explanation of the collection activities and the burden estimate that is only briefly summarized here:

Form number(s): None.

Respondents/affected entities: Entities potentially affected by this ICR include Chemical and Allied Products Merchant Wholesaler firms and Basic Chemical Manufacturing firms North American Industrial Classification System (NAICS) codes identified in question 12 of the ICR.

Respondent's obligation to respond: Mandatory. Per 40 CFR 751 and 15 U.S.C. 2605(a)

Estimated number of potential respondents: 6,515.

Frequency of response: On occasion.

Total estimated average number of responses for each respondent: 7.41.

Total estimated burden: 72,699 hours (per year). Burden is defined at 5 CFR 1320.3(b).

Total estimated costs: $5,342,124 (per year), includes $4,583,912 annualized capital investment or maintenance and operational costs.

III. Are there changes in the estimates from the last approval?

There is an increase of 72,692 hours in the total estimated respondent burden compared with that identified in the ICR currently approved by OMB. This change, which is discussed in more detail in the ICR, reflects increase is a result of consolidating two existing ICRs into one. This change is an adjustment.

IV. What is the next step in the process for this ICR?

EPA will consider the comments received and amend the ICR as appropriate. The final ICR package will then be submitted to OMB for review and approval pursuant to 5 CFR 1320.12. EPA will issue another **Federal Register** document pursuant to 5 CFR 1320.5(a)(1)(iv) to announce the submission of the ICR to OMB and the opportunity to submit additional comments to OMB. If you have any questions about this ICR or the approval process, please contact the person listed under **FOR FURTHER INFORMATION CONTACT**.

Authority: 44 U.S.C. 3501 *et seq.*

Dated: August 15, 2025.

Nancy B. Beck,

Principal Deputy Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

[FR Doc. 2025–16216 Filed 8–22–25; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL DEPOSIT INSURANCE CORPORATION

[OMB No. 3064–0026;–0178]

Agency Information Collection Activities: Proposed Collection Renewal; Comment Request

AGENCY: Federal Deposit Insurance Corporation (FDIC).

ACTION: Notice and request for comment.

SUMMARY: The FDIC, as part of its obligations under the Paperwork Reduction Act of 1995, invites the general public and other Federal agencies to take this opportunity to comment on the request to renew the existing information collections described below (OMB Control No. 3064–0026 and –0178). The notices of proposed renewal for these information collections were previously published in the **Federal Register** on June 23, 2025, allowing for a 60-day comment period.

DATES: Comments must be submitted on or before September 24, 2025.

ADDRESSES: Interested parties are invited to submit written comments to the FDIC by any of the following methods:

- *Agency Website: https://www.fdic.gov/resources/regulations/federal-register-publications/.*
- *Email: comments@fdic.gov.* Include the name and number of the collection in the subject line of the message.
- *Mail:* Robert Meiers, Regulatory Attorney, MB–3013, Federal Deposit Insurance Corporation, 550 17th Street NW, Washington, DC 20429.
- *Hand Delivery:* Comments may be hand-delivered to the guard station at the rear of the 17th Street NW building (located on F Street NW), on business days between 7 a.m. and 5 p.m.

Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find these information collections by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: Robert Meiers, Regulatory Attorney, *Romeiers@fdic.gov,* MB–3013, Federal Deposit Insurance Corporation, 550 17th Street NW, Washington, DC 20429.

SUPPLEMENTARY INFORMATION: Proposal to renew the following currently approved collection of information:

1. *Title:* Reporting Requirements for Transfer Agents.

OMB Number: 3064–0026.

Form Number: TA–1.

Affected Public: Private sector, insured State nonmember banks and State savings associations.

Burden Estimate:

SUMMARY OF ESTIMATED ANNUAL BURDEN (OMB NO. 3064–0026)

Information collection (IC) (obligation to respond)	Type of burden (frequency of response)	Number of respondents	Number of responses per respondent	Average time per response (HH:MM)	Annual burden (Hours)
1. Transfer Agent Registration, 12 CFR 341.3 (Mandatory).	Reporting (Occasional)	1	1	01:15	1
2. Transfer Agent Amendment, 12 CFR 341.4 (Mandatory).	Reporting (Occasional)	1	1	00:10	0
3. Transfer Agent Deregistration, 12 CFR 341.5 (Mandatory).	Reporting (Occasional)	1	1	00:25	0
Total Annual Burden (Hours)					1

Source: FDIC.

General Description of Collection: Section 17A(c) of the Security Exchange Act of 1934 (the Act) requires all transfer agents for securities registered under section 12 of the Act or, if the security would be required to be registered except for the exemption from registration provided by section 12(g)(2)(B) or section 12(g)(2)(G), to ''fil[e] with the appropriate regulatory agency . . . an application for registration in such form and containing such information and documents . . . as such appropriate regulatory agency may prescribe as necessary or appropriate in furtherance of the purposes of this section.'' In general, an entity performing transfer agent functions for a security is required to register with its appropriate regulatory agency if the security is registered on a national securities exchange or if the issuer of the security has total assets exceeding $10 million and a class of equity security held of record by 2,000 persons or, for an issuer that is not a bank, bank holding company, or savings and loan holding company, by 500 persons who are not accredited investors. The Federal Reserve Board of Governors' Regulation H (12 CFR 208.31(a)) and Regulation Y (12 CFR 225.4(d)), the OCC's 12 CFR 9.20, and the FDIC's 12 CFR part 341 implement these provisions of the Act. To accomplish the registration of transfer agents, Form TA–1 was developed in 1975 as an interagency effort by the Securities and Exchange Commission and the agencies. The agencies primarily use the data collected on Form TA–1 to determine whether an application for registration should be approved, denied, accelerated or postponed, and they use the data in connection with their supervisory responsibilities. There is no change in the methodology or substance of this information collection. The estimated burden remains unchanged from the previous submission.

2. *Title:* Market Risk Capital Requirements.

OMB Number: 3064–0178.

Form Number: None.

Affected Public: Insured State nonmember banks and State savings associations.

Burden Estimate:

SUMMARY OF ESTIMATED ANNUAL BURDEN (OMB NO. 3064–0178)

Information collection (IC) (obligation to respond)	Type of burden (frequency of response)	Number of respondents	Number of responses per respondent	Average time per response (HH:MM)	Annual burden (Hours)
1. Prior Approval, 12 CFR 324.203(c)(1), 324.203(c)(2), 324.204(a)(2)(vi)(B), 324.206(b)(3), 324.208(a), 324.209(a) (Mandatory).	Reporting (Annual)	1	1	128:00	128
2. Policies and Procedures, 12 CFR 324.203(a)(1), 324.203(b)(1), 324.203(b)(2), 324.206(b)(3) (Mandatory).	Recordkeeping (Annual)	1	1	112:00	112

SUMMARY OF ESTIMATED ANNUAL BURDEN (OMB NO. 3064–0178)—Continued

Information collection (IC) (obligation to respond)	Type of burden (frequency of response)	Number of respondents	Number of responses per respondent	Average time per response (HH:MM)	Annual burden (Hours)
3. Trading and Hedging Strategy, 12 CFR 324.203(a)(2) (Mandatory).	Recordkeeping (Annual)	1	1	16:00	16
4. General Recordkeeping, 12 CFR 324.203(f) (Mandatory).	Recordkeeping (Annual)	1	1	24:00	24
5. Back testing, 12 CFR 324.205(c) (Mandatory).	Recordkeeping (Annual)	1	1	24:00	24
6. Stress testing, 12 CFR 324.209(c)(2) (Mandatory).	Recordkeeping (Annual)	1	4	08:00	32
7. Securitizations, 12 CFR 324.210(f)(1) (Mandatory).	Recordkeeping (Annual)	1	1	08:00	8
8. Disclosure Policy, 12 CFR 324.212(b) (Mandatory).	Recordkeeping (Annual)	1	1	40:00	40
9. Quantitative Disclosure, 12 CFR 324.212(c) (Mandatory).	Disclosure (Annual)	1	4	08:00	32
10. Qualitative Disclosure, 12 CFR 324.212(d) (Mandatory).	Disclosure (Annual)	1	1	12:00	12
Total Annual Burden (Hours):					428

Source: FDIC.

General Description of Collection: The FDIC's market risk capital rules (12 CFR part 324, subpart F) enhance risk sensitivity, increase transparency through enhanced disclosures and include requirements for the public disclosure of certain qualitative and quantitative information about the market risk of State nonmember banks and State savings associations (covered FDIC-supervised institutions). The market risk rule applies only if a bank holding company or bank has aggregated trading assets and trading liabilities equal to 10 percent or more of quarter-end total assets or $1 billion or more (covered FDIC-supervised institutions). Currently, only one FDIC-regulated entity meets the criteria of the information collection requirements that are located at 12 CFR 324.203 through 324.212. The collection of information is necessary to ensure capital adequacy appropriate for the level of market risk. Section 324.203(a)(1) requires covered FDIC-supervised institutions to have clearly defined policies and procedures for determining which trading assets and trading liabilities are trading positions and specifies the factors a covered FDIC-supervised institution must take into account in drafting those policies and procedures. Section 324.203(a)(2) requires covered FDIC-supervised institutions to have clearly defined trading and hedging strategies for trading positions that are approved by senior management and specifies what the strategies must articulate. Section 324.203(b)(1) requires covered FDIC-supervised institutions to have clearly defined policies and procedures for actively managing all covered positions and specifies the minimum requirements for those policies and procedures. Sections 324.203(c)(4) through (10) require the annual review of internal models and specify certain requirements for those models. Section 324.203(d) requires the internal audit group of a covered FDIC-supervised institution to prepare an annual report to the board of directors on the effectiveness of controls supporting the market risk measurement systems. Section 324.204(b) requires covered FDIC-supervised institutions to conduct quarterly back testing. Section 324.205(a)(5) requires institutions to demonstrate to the FDIC the appropriateness of proxies used to capture risks within value-at-risk models. Section 324.205(c) requires institutions to develop, retain, and make available to the FDIC value-at-risk and profit and loss information on sub portfolios for two years. Section 324.206(b)(3) requires covered FDIC-supervised institutions to have policies and procedures that describe how they determine the period of significant financial stress used to calculate the institution's stressed value-at-risk models and to obtain prior FDIC approval for any material changes to these policies and procedures. Section 324.207(b)(1) details requirements applicable to a covered FDIC-supervised institution when the covered FDIC-supervised institution uses internal models to measure the specific risk of certain covered positions. Section 324.208 requires covered FDIC-supervised institutions to obtain prior written FDIC approval for including equity positions in its incremental risk modeling. Section 324.209(a) requires prior FDIC approval for the use of a comprehensive risk measure. Section 324.209(c)(2) requires covered FDIC-supervised institutions to retain and report the results of supervisory stress testing. Section 324.210(f)(2)(i) requires covered FDIC-supervised institutions to document an internal analysis of the risk characteristics of each securitization position in order to demonstrate an understanding of the position. Section 324.212 applies to certain covered FDIC-supervised institutions that are not subsidiaries of bank holding companies, and requires quarterly quantitative disclosures, annual qualitative disclosures, and a formal disclosure policy approved by the board of directors that addresses the approach for determining the market risk disclosures it makes. The total estimated annual burden is 428 hours, which is a reduction of 4,032 hours from the 2022 submission. This reduction is due to a change in agency estimates. The FDIC's estimates significantly lowered because respondent institutions have generally already received prior approval for incremental risk modeling and the use of a comprehensive risk measure for one or more portfolios of correlation trading positions. Therefore, the agency predicts these respondents will not re-submit these models for approval, reducing the overall burden hours.

Request for Comment

Comments are invited on (a) whether the collection of information is necessary for the proper performance of the FDIC's functions, including whether

the information has practical utility; (b) the accuracy of the estimates of the burden of the information collection, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Federal Deposit Insurance Corporation.

Dated at Washington, DC, on August 21, 2025.

Jennifer M. Jones,

Deputy Executive Secretary.

[FR Doc. 2025–16200 Filed 8–22–25; 8:45 am]

BILLING CODE 6714–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifiers: CMS–10434 #26 and #47]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

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

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an overarching plan that defines the scope of the individual collections that would fall under its umbrella. This **Federal Register** notice seeks public comment on one or more of our collection of information requests that we believe are generic and fall within the scope of the umbrella. Interested persons are invited to submit 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 September 8, 2025.

ADDRESSES: When commenting, please reference the applicable form number (CMS–10434 #____) and the OMB control number (0938–1188). 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: CMS–10434____/OMB control number: 0938–1188, Room C4–26–05, 7500 Security Boulevard, Baltimore, MD 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/medicare/regulations-guidance/legislation/paperwork-reduction-act-1995/pra-listing*.

FOR FURTHER INFORMATION CONTACT: William N. Parham at 410–786–4669.

SUPPLEMENTARY INFORMATION: Following is a summary of the use and burden associated with the subject information collection(s). More detailed information can be found in the collection's supporting statement and associated materials (see **ADDRESSES**).

Generic Information Collection

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Child and Adult Core Set Measures; *Use:* CMS is required annually to revise and update the Child and Adult Core Sets. The review for the 2025 Core Set resulted in several substantive changes to the templates used for state Core Set reporting to CMS. State reporting of the Core Sets in the Quality Measurement Reporting system opens for 2025 Core Set reporting on September 3, 2025. *Form Number:* CMS–10434 #26 (OMB control number: 0938–1188); *Frequency:* Yearly, once, and occasionally; *Affected Public:* Individuals or households and State, Local, or Tribal Governments; *Number of Respondents:* 61,293; *Total Annual Responses:* 61,455; *Total Annual Hours:* 99,977. (For policy questions regarding this collection contact: Virginia (Gigi) Raney at 410–786–6117.)

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Health Home Core Sets Measures; *Use:* CMS is required annually to revise and update the Health Home Core Sets. The review for the 2025 Core Set resulted in several substantive changes to the templates used for state Core Set reporting to CMS. State reporting of the Core Sets in the Quality Measurement Reporting system opens for 2025 Core Set reporting on September 3, 2025. *Form Number:* CMS–10434 #47 (OMB control number: 0938–1188); *Frequency:* Yearly, once, and occasionally; *Affected Public:* State, Local, or Tribal Governments; *Number of Respondents:* 50; *Total Annual Responses:* 50; *Total Annual Hours:* 4,620. (For policy questions regarding this collection contact: Sara Rhoades at 410–786–4484.)

Evell Barco Holland,

Senior Technical Advisor, Regulations Development Group, Office of Strategic Operations and Regulatory Affairs.

[FR Doc. 2025–16171 Filed 8–22–25; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2025–N–2962]

Prescription Drug User Fee Act and Biosimilar User Fee Amendments Hiring and Retention Assessment; Public Meeting; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the following public meeting entitled ''Prescription Drug User Fee Act and Biosimilar User Fee Amendments Hiring and Retention Assessment.'' The topic to be discussed is a hiring and retention assessment