FEDERAL RESERVE SYSTEM

Agency Information Collection Activities: Announcement of Board Approval Under Delegated Authority and Submission to OMB

AGENCY: Board of Governors of the Federal Reserve System.

SUMMARY: The Board of Governors of the Federal Reserve System (Board) is adopting a proposal to extend for three years, with revision, the New Hire Information Collection (FR 27; OMB No. 7100–0375).

DATES: The information collection revisions are effective June 9, 2025.

FOR FURTHER INFORMATION CONTACT: Federal Reserve Board Clearance Officer—Nuha Elmaghrabi—Office of the Chief Data Officer, Board of Governors of the Federal Reserve System, *nuha.elmaghrabi@frb.gov,* (202) 452–3884.

Office of Management and Budget (OMB) Desk Officer for the Federal Reserve Board, Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Room 10235, 725 17th Street NW, Washington, DC 20503, or by fax to (202) 395–6974.

SUPPLEMENTARY INFORMATION: On June 15, 1984, OMB delegated to the Board authority under the Paperwork Reduction Act (PRA) to approve and assign OMB control numbers to collections of information conducted or sponsored by the Board. Board-approved collections of information are incorporated into the official OMB inventory of currently approved collections of information. The OMB inventory, as well as copies of the PRA Submission, supporting statements (which contain more detailed information about the information collections and burden estimates than this notice), and approved collection of information instrument(s) are available at *https://www.reginfo.gov/public/do/PRAMain.* These documents are also available on the Federal Reserve Board's public website at *https://www.federalreserve.gov/apps/reportingforms/review* or may be requested from the agency clearance officer, whose name appears above. On the page displayed at the link above, you can find the supporting information by referencing the collection identifier, FR 27.

Final Approval Under OMB Delegated Authority of the Extension for Three Years, With Revision, of the Following Information Collection

Collection title: New Hire Information Collection.

Collection identifier: FR 27.

OMB control number: 7100–0375.

General description of collection: This information collection provides for the electronic collection of certain personnel information from new hires using a secure web-based portal, the New Hire Portal, before the first day of employment of a new hire. As part of the onboarding process for new hires, a Human Resources (HR) professional at the Board identifies the necessary information that must be collected from the new hire, which is dependent upon the type of hire that the person is. The types of hires include Regular Hires, which are hires who are being hired into a non-intern position and not transferring from a non-federal agency, including a Governor or Board officer; Intern Hires, which are hires being hired into an intern position; and Federal Transfers, which are hires who are transferring from another federal agency. Once the HR professional has identified the types of information that will be necessary, the new hire is sent an email asking him or her to provide certain information through the New Hire Portal prior to their official start date.

The New Hire Portal is broken out into different sections and each section corresponds to the hardcopy forms that new employees previously filled out and provided to the Board during or after the first day of New Employee Orientation. The information collection now involves a new hire electronically providing this personnel information and filling out the applicable sections of the New Hire Portal before their first day of orientation. The sections of the portal that each new hire is asked to complete electronically depend upon the type of position that the new hire has been offered at the Board.

Frequency: Event-generated.

Respondents: The FR 27 panel comprises individuals who are new hires to the Board but have not yet become employees.

Total estimated number of respondents: Regular Hire, 235; Intern Hire, 131; Federal Transfer, 39; Governor/Officer, 9; Contingent Worker, 637.

Estimated average hours per response: Regular Hire, 1.78; Intern Hire, 1.71; Federal Transfer, 1.95; Governor/Officer, 1.86; Contingent Worker, 1.5.

Total estimated change in burden: 1,298.

Total estimated annual burden hours: 1,691.

Current actions: On March 11, 2025, the Board published a notice in the **Federal Register** (90 FR 11738) requesting public comment for 60 days on the extension, with revision, of the FR 27. The Board proposed to revise the FR 27 by changing the collection platform from the New Hire Portal to Workday Onboarding in June 2025, adding two new categories of hires categorized as Officers/Governors and Contingent Workers, restructuring sections into individual tasks, adding new data fields, removing data fields, and relabeling data fields. There will also be a change in the login process, as all respondents, except Contingent Workers, will be required to complete identity proofing and set up a secure account through a separate system, *Login.gov,* before accessing Workday Onboarding. The Designation of Beneficiary Unpaid Compensation of Deceased Employee form and Executive Death and Dismemberment Benefit for Officers and Governors questionnaire, previously not subject to PRA, will now be collected prior to a hire's first day of employment. The Board will no longer collect information for use on the FEGLI Program Designation of Beneficiary form through the New Hire Information Collection. The Board will also no longer collect information to populate the state tax form prior to the hire's start date, so it will no longer be subject to PRA. The Board will begin collecting Form I–9 (Employment Eligibility Verification) electronically through Workday Onboarding, rather than via a PDF. The comment period for this notice expired on May 12, 2025. The Board received one comment from one individual. The comment expressed support for the extension, with revision, of the FR 27. The Board will adopt the extension, with revision, of the FR 27 as originally proposed.

Board of Governors of the Federal Reserve System, June 2, 2025.

Benjamin W. McDonough,

Deputy Secretary and Ombuds of the Board.

[FR Doc. 2025–10239 Filed 6–4–25; 8:45 am]

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OFFICE OF GOVERNMENT ETHICS

Agency Information Collection Activities; Information Collection Renewal; Comment Request for Qualified Trust Model Certificates and Model Trust Documents

AGENCY: Office of Government Ethics (OGE).

ACTION: Notice of request for agency and public comments.

SUMMARY: After this first round notice and public comment period, the U.S. Office of Government Ethics (OGE) plans to request that the Office of

Management and Budget (PMB) renew its approval under the Paperwork Reduction Act for an existing information collection consisting of twelve model certificates and model documents for qualified trusts.

DATES: Written comments by the public and the agencies on this proposed extension are invited and must be received on or before August 4, 2025.

ADDRESSES: Comments may be submitted to OGE by any of the following methods:

Email: usoge@oge.gov (Include reference to ''OGE qualified trust model certificates and model trust documents paperwork comment'' in the subject line of the message.)

Mail: Office of Government Ethics, 250 E Street SW, Suite 750, Attention: Jennifer Matis, Associate Counsel, Washington, DC 20005–3917.

Instructions: Comments may be posted on OGE's website, *www.oge.gov.* Sensitive personal information, such as account numbers or Social Security numbers, should not be included. Comments generally will not be edited to remove any identifying or contact information.

FOR FURTHER INFORMATION CONTACT: Jennifer Matis at the U.S. Office of Government Ethics; telephone: 202–482–9216; TTY: 800–877–8339; Email: *jmatis@oge.gov.* Copies of the model documents as currently approved are available on OGE's website, *www.oge.gov.*

SUPPLEMENTARY INFORMATION:

Title: Executive Branch Qualified Trust Documents.

Type of Information Collection: Extension of a currently approved collection.

Type of Review Request: Regular.

Respondents: Any current or prospective executive branch officials who seek to establish or have established a qualified blind or diversified trust under the Ethics in Government Act of 1978 to avoid conflicts of interest while in office.

Estimated Average Annual Number of Respondents: 2.

Total Estimated Time per Response: 20 minutes to 100 hours (see table below for detailed explanation).

Estimated Average Total Annual Burden: 120 hours.

OMB Control Number: 3209–0007.

Abstract: OGE is the supervising ethics office for the executive branch of the Federal Government under the Ethics in Government Act of 1978 (EIGA). Accordingly, OGE administers the qualified trust program for the executive branch. Presidential nominees to executive branch positions subject to Senate confirmation and any other executive branch officials may seek OGE approval for EIGA-qualified blind or diversified trusts as one means to avoid conflicts of interest. The requirements for EIGA-qualified blind and diversified trusts are set forth in section 13104(f) of the Ethics in Government Act, 5 U.S.C. 13104(f), and OGE's implementing financial disclosure regulations at subpart D of 5 CFR part 2634.

In order to ensure that all applicable requirements are met, OGE is the sponsoring agency for twelve model certificates and model trust documents for qualified blind and diversified trusts. See 5 CFR 2634.402(e)(3), 2634.402(f)(3), 2634.404(e) through (g), 2634.405(d)(2), 2634.407(a); 2634.408(b)(1) through (3), 2634.408(d)(4), 2634.409, and 2634.414. The various model certificates and model trust documents are used by settlors, trustees, and other fiduciaries in establishing and administering these qualified trusts. OGE plans to submit these model certificates and model trust documents (described in detail in the table below) to OMB for renewed approval pursuant to the Paperwork Reduction Act of 1995, 44 U.S.C. chapter 35.

The twelve model documents, along with their burden estimates, are as follows:

Model qualified trust documents	Estimated burden
(A) Blind Trust Communications (Expedited Procedure for Securing Approval of Proposed Communications)	20 minutes per communication.
(B) Model Qualified Blind Trust Provisions	100 hours per model.
(C) Model Qualified Diversified Trust Provisions	100 hours per model.
(D) Model Qualified Diversified Trust Provisions (For Use in the Case of Multiple Fiduciaries)	100 hours per model.
(E) Model Qualified Blind Trust Provisions (For Use in the Case of an Irrevocable Pre-Existing Trust)	100 hours per model.
(F) Hybrid Version of the Model Qualified Diversified Trust Provisions	100 hours per model.
(G) Model Qualified Blind Trust Provisions (For Use in the Case of Multiple Fiduciaries)	100 hours per model.
(H) Model Qualified Diversified Trust Provisions (For Use in the Case of an Irrevocable Pre-Existing Trust)	100 hours per model.
(I) Model Confidentiality Agreement Provisions (For Use in the Case of a Privately Owned Business)	2 hours per agreement.
(J) Model Confidentiality Agreement Provisions (For Use in the Case of Investment Management Activities)	2 hours per agreement.
Model Trust Certificates	
(K) Certificate of Independence	20 minutes per certificate.
(L) Certificate of Compliance	20 minutes per certificate.

These estimates are based on the amount of time imposed on professional trust administrators or private representatives. OGE notes that only one set of the various model trust provisions (items (B) through (H)) will be prepared for a single qualified trust, and only prior to the establishment of that qualified trust. Likewise, other model documents listed above are used in connection with establishing the qualified trust (items (I), (J), and (K)). The remaining model documents are used after the trust's creation (items (A) and (L)). Accordingly, OGE notes that the majority of the time burden for any given trust is imposed during the creation of the trust.

At the present time, there are no active qualified trusts in the executive branch. However, OGE anticipates possible limited use of these model documents during the forthcoming three-year period. OGE estimates that there may be an average of one individual per year who initiates a qualified trust using these model documents during calendar years 2026 through 2028. OGE has accordingly estimated the average annual number of respondents to be two, which represents one respondent establishing a qualified trust and one respondent maintaining a previously established qualified trust. Based on the above, OGE estimates an average annual time burden during the next three years of 120 hours. Using an estimated rate of $300 per hour for the services of a professional trust administrator or private representative, the estimated annual cost burden is $36,000.

Under OMB's implementing regulations for the Paperwork Reduction Act, any recordkeeping, reporting, or disclosure requirement contained in a rule of general applicability is deemed to involve ten or more persons. See 5 CFR 1320.3(c)(4)(i). OGE intends to submit all twelve qualified trust model certificates and model documents described above (all of which are included under OMB paperwork control number 3209–0007) for a three-year extension of approval without modification.

Request for Comments: Agency and public comments are invited specifically on the need for and practical utility of this information collection, on the accuracy of OGE's burden estimate, on the enhancement of quality, utility, and clarity of the information collected, and on minimizing the burden to the public. Comments received in response to this notice will be summarized for, and may be included with, the OGE request for extension of OMB approval. The comments will also become a matter of public record.

Specifically, OGE seeks public comment on the following:

- Do the model qualified blind trusts provide sufficient direction to establish a trust under the Qualified Trust Program? If not, what provisions could be clearer or what language should be changed?
- Do the model qualified diversified trusts provide sufficient direction to establish a trust under the Qualified Trust Program? If not, what provisions could be clearer or what language should be changed?
- Do the Additional Trust Documents provide sufficient information for individuals to comply with the logistical requirements (*e.g.,* procedure for securing approval of proposed communications) of the Qualified Trust Program? If not, what provisions could be clearer or what language should be changed?

Approved: May 28, 2025.

Shelley K. Finlayson,

Chief of Staff and Program Counsel, Office of Government Ethics.

[FR Doc. 2025–10211 Filed 6–4–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2024–D–1528]

Transfer of a Premarket Notification (510(k)) Clearance—Questions and Answers; Draft Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, Department of Health and Human Services (HHS).

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of the draft guidance entitled ''Transfer of a Premarket Notification (510(k)) Clearance—Questions and Answers.'' This guidance provides information on the most frequently asked questions regarding transfer of ownership from one 510(k) holder to another. This draft guidance is not final nor is it for implementation at this time.

DATES: Submit either electronic or written comments on the draft guidance by August 4, 2025 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

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–2024–D–1528 for ''Transfer of a Premarket Notification (510(k)) Clearance—Questions and Answers.'' Received comments 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 available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://*