SUMMARY OF ESTIMATED ANNUAL BURDEN—Continued

[OMB No. 3064–0087]

Information collection (obligation to respond)	Type of burden (frequency of response)	Number of respondents	Number of responses per respondent	Time per response (HH:MM)	Annual burden (hours)
Total Annual Burden (Hours)					338,905

Source: FDIC.

General Description of Collection: Respondents must establish and maintain procedures designed to monitor and ensure their compliance with the requirements of the Bank Secrecy Act and the implementing regulations promulgated by the Department of Treasury at 31 CFR chapter X. Respondents must also keep records evidencing that they have provided training for appropriate personnel. There is no change in the method or substance of the collection. The overall increase in burden hours is a result of economic fluctuation. In particular, the total number of respondents has increased while the hours per response remain the same.

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 July 7, 2023.

James P. Sheesley,

Assistant Executive Secretary.

[FR Doc. 2023–14823 Filed 7–12–23; 8:45 am]

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GENERAL SERVICES ADMINISTRATION

[Notice–MRB–2023–03; Docket No. 2023–0001; Sequence No. 22]

Regulatory Information Systems Center; Announcement of Public Listening Sessions

AGENCY: Office of Government-wide Policy, General Services Administration (GSA).

ACTION: Notice.

SUMMARY: To assist with the Regulatory Information website (*Reginfo.gov*) user experience research, the Regulatory Information Systems Center (RISC) will be hosting public listening sessions. The purpose of these listening sessions is to collect public input on the usability of *Reginfo.gov.* In turn, RISC will use the input to inform future enhancements to *Reginfo.gov.*

DATES: RISC will hold web-based public listening sessions on Tuesday, August 8, 2023, from 1:00 p.m. to 3:00 p.m. Eastern Standard Time (EST) and on Thursday, August 10, 2023, from 10 a.m. to 12:00 p.m. EST.

ADDRESSES: The virtual listening sessions will be open to the public and held via the Zoom Webinar Platform. Virtual attendance information will be provided upon registration. Registration information is located on Eventbrite: *https://www.eventbrite.com/e/risc-presents-reginfo-public-comment-session-tickets-668851050497.*

In addition to the listening sessions, written public comments are being accepted via email. To submit a written public comment, send an email to *risc@gsa.gov.* Please include ''Reginfo.gov Public Comment'' in the subject line. In the body of the email, please include your name, company name (if applicable), and years of *Reginfo.gov* usage.

FOR FURTHER INFORMATION CONTACT: Please contact Mr. Wesley Weston, Senior Program Analyst, RISC, 202–251–7769 or by email at *wesley.weston@gsa.gov.*

SUPPLEMENTARY INFORMATION:

Background

Reginfo.gov assists users who want to find federal regulatory information and provides a variety of graphical displays constituting a ''Regulatory Dashboard.'' Users can select and identify rules under review by agency, economic significance, stage of rulemaking, or other characteristics, and compare the results for different agencies.

Reginfo.gov provides information on the following areas:

- Federal regulatory agendas and regulatory plans to include brief synopsis and timetables for action on rules that Federal departments and agencies are considering.
- Rules under review by the Office of Information and Regulatory Affairs (OIRA) prior to initial publication or final adoption are listed.
- OIRA reviews of information collections, such as forms and surveys, under the Paperwork Reduction Act (PRA), are listed together with a complete inventory of currently approved information collections.

Reginfo.gov gives the public searchable access to this information to make more transparent the activities of OIRA and Federal agencies in rulemaking and information collection.

Specifically, RISC invites public comment on the following questions:

1. On the homepage, do you find the visual graphs helpful? If not, what other tool would you recommend to present the information?

2. How do you feel about the overall navigation of the site? Do you feel the main navigation covers what you are looking for when visiting the site?

3. Do you find the current search options useful? Have you had difficulty using the search option based on its current location?

4. Do you feel the ''Contact us'' information or ''Getting help'' is easily found when visiting the site?

5. Have you been able to find answers to questions you were looking for? Did you have to use another site? Please explain.

6. How would you like real time information presented?

7. Have you used the mobile app which is available to both android and

IOS users? If so, what has been your experience with the app?

8. Overall, what information is missing when visiting this site?

9. Overall, what improvements would you suggest to make the site better?

Meeting Registration

This meeting is open to the public and will be accessible by webcast. All public attendees will need to register to obtain the meeting webcast information. All registrants will be asked to provide their name, affiliation, and email address. After registration, individuals will receive webcast access information via email.

Public Participation

The public listening sessions will start at 1:00 p.m. EST, on August 8, 2023, and 10:00 a.m. EST on August 10, 2023. The RISC team first will provide opening remarks. The meetings will then transition to public comments. Any oral comments presented should be brief and limited to the subjects described in this Notice so all participants will have an opportunity to speak.

Members of the public who wish to present oral comments must notify RISC no later than Monday, August 7, 2023, via email at *risc@gsa.gov.* The email should (1) identify specific subject(s) on which you wish to provide comments; and (2) state the organization or entity you are representing or that you are speaking as a member of the public.

Boris Arratia,

Regulatory Information Service Center Director, Office of Government-wide Policy.

[FR Doc. 2023–14842 Filed 7–12–23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request; Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project ''*Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery.*''

DATES: Comments on this notice must be received by September 11, 2023.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at *doris.lefkowitz@AHRQ.hhs.gov.*

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

Generic Clearance for the Collection of Qualitative Feedback on Agency Service Delivery

The information collection activity will garner qualitative customer and stakeholder feedback in an efficient, timely manner, in accordance with the Administration's commitment to improving service delivery. By qualitative feedback we mean information that provides useful insights on perceptions and opinions, but are not statistical surveys that yield quantitative results that can be generalized to the population of study. This feedback will provide insights into customer or stakeholder perceptions, experiences, and expectations, provide an early warning of issues with service, or focus attention on areas where communication, training or changes in operations might improve delivery of products or services. These collections will allow for ongoing, collaborative and actionable communications between the Agency and its customers and stakeholders. It will also allow feedback to contribute directly to the improvement of program management. The current clearance was approved on November 2, 2020 (OMB Control Number 0935–0179) and will expire on November 30, 2023. Feedback collected under this generic clearance will provide useful information, but it will not yield data that can be generalized to the overall population. This type of generic clearance for qualitative information will not be used for quantitative information collections that are designed to yield reliably actionable results, such as monitoring trends over time or documenting program performance. Such data uses require more rigorous designs that address: (1) the target population to which generalizations will be made; (2) the sampling frame; (3) the sample design (including stratification and clustering); (4) the precision requirements or power calculations that justify the proposed sample size; (5) the expected response rate; (6) methods for assessing potential nonresponse bias; (7) the protocols for data collection; (8) and any testing procedures that were or will be undertaken prior to fielding the study. Depending on the degree of influence the results are likely to have, such collections may still be eligible for submission for other generic mechanisms that are designed to yield quantitative results.

Below we provide AHRQ's projected average annual estimates for the next three years:

Current Actions: New collection of information.

Type of Review: New Collection.

Affected Public: Individuals and Households, Businesses and Organizations, State, Local or Tribal Government.

Average Expected Annual Number of Activities: 10.

Respondents: 10,900.

Annual Responses: 10,900.

Frequency of Response: Once per request.

The total number of respondents across all 10 activities each year is 10,900.

Average Minutes per Response: 19.

Burden Hours: 3,383.

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 Office of Management and Budget control number.

Request for Comments

In accordance with the Paperwork Reduction Act, comments on AHRQ's information collection are requested with regard to any of the following: (a) Whether the proposed collection of information is necessary for the proper performance of AHRQ healthcare research and healthcare information dissemination functions, including whether the information will have practical utility; (b) the accuracy of AHRQ's estimate of burden (including hours and costs) of the proposed collection(s) of information; (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 upon the respondents, including the use of automated collection techniques or other forms of information technology. Comments submitted in response to this notice will be summarized and included in the Agency's subsequent request for OMB approval of the proposed information collection. All comments will become a matter of public record.