the Chief Data Officer, Board of Governors of the Federal Reserve System, *nuha.elmaghrabi@frb.gov,* (202) 452–3884.

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. In exercising this delegated authority, the Board is directed to take every reasonable step to solicit comment. In determining whether to approve a collection of information, the Board will consider all comments received from the public and other agencies.

During the comment period for this proposal, a copy of the proposed PRA OMB submission, including the draft reporting form and instructions, supporting statement (which contains more detail about the information collection and burden estimates than this notice), and other documentation, will be made available on the Board's public website at *https://www.federalreserve.gov/apps/reportingforms/home/review* or may be requested from the agency clearance officer, whose name appears above. Final versions of these documents will be made available at *https://www.reginfo.gov/public/do/PRAMain,* if approved.

Request for Comment on Information Collection Proposal

The Board invites public comment on the following information collection, which is being reviewed under authority delegated by the OMB under the PRA. Comments are invited on the following:

a. Whether the proposed collection of information is necessary for the proper performance of the Board's functions, including whether the information has practical utility;

b. The accuracy of the Board's estimate of the burden of the proposed 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;

d. Ways to minimize the burden of information collection on respondents, including through the use of automated collection techniques or other forms of information technology; and

e. Estimates of capital or startup costs and costs of operation, maintenance, and purchase of services to provide information.

At the end of the comment period, the comments and recommendations received will be analyzed to determine the extent to which the Board should modify the proposal.

Proposal Under OMB Delegated Authority To Extend for Three Years, Without Revision, the Following Information Collection

Collection title: Recordkeeping Requirements Associated with Regulation GG.

Collection identifier: FR GG.

OMB control number: 7100–0317.

General description of collection: Regulation GG—Prohibition on Funding of Unlawful internet Gambling (12 CFR part 233) is related to the Unlawful internet Gambling Enforcement Act of 2006. The FR GG is the Board's information collection associated with Regulation GG and requires participants in designated payment systems to establish written policies and procedures related to unlawful internet gambling. These recordkeeping requirements are contained in section 5 of Regulation GG. Policies and procedures reasonably designed to identify and block, or otherwise prevent or prohibit, restricted transactions are necessary because Congress found that internet gambling is primarily funded through personal use of payment system instruments, credit cards, and wire transfers.

Frequency: Event-generated.

Respondents: Depository institutions, card system operators, credit unions, and money transmitting business operators.

Total estimated number of respondents: 4,635.

Total estimated annual burden hours: 46,410.[1]

Board of Governors of the Federal Reserve System, March 4, 2024.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

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BILLING CODE 6210–01–P

[1] More detailed information regarding this collection, including more detailed burden estimates, can be found in the OMB Supporting Statement posted at *https://www.federalreserve.gov/apps/reportingforms/home/review.* On the page displayed at the link, you can find the OMB Supporting Statement by referencing the collection identifier, FR GG.

GENERAL SERVICES ADMINISTRATION

[Notice-Q–2024–02; Docket No. 2024–0002; Sequence No. 4]

Federal Secure Cloud Advisory Committee Notification of Upcoming Meeting

AGENCY: Federal Acquisition Service (Q), General Services Administration (GSA).

ACTION: Meeting notice.

SUMMARY: As stipulated by the Federal Advisory Committee Act (FACA), as amended, GSA is hereby giving notice of an open public meeting of the Federal Secure Cloud Advisory Committee (FSCAC). Information on attending and providing public comment is under the **SUPPLEMENTARY INFORMATION** section.

DATES: The open public meeting will be held on Thursday, March 28, 2024, from 10 a.m. to 3 p.m., eastern standard time (EST). The agenda for the meeting will be made available prior to the meeting online at *https://gsa.gov/fscac.*

ADDRESSES: The meeting will be held in person at 1800 F St NW, Room 1461, Washington, DC 20004. The meeting will also be accessible via live stream. Registrants will receive the live stream information before the meeting.

FOR FURTHER INFORMATION CONTACT: Michelle White, Designated Federal Officer (DFO), FSCAC, GSA, 703–489–4160, *fscac@gsa.gov.* Additional information about the Committee, including meeting materials and agendas, will be available online at *https://gsa.gov/fscac.*

SUPPLEMENTARY INFORMATION:

Background

GSA, in compliance with the FedRAMP Authorization Act of 2022, established the FSCAC, a statutory advisory committee in accordance with the provisions of FACA, as amended (5 U.S.C. 10). The Federal Risk and Authorization Management Program (FedRAMP) within GSA is responsible for providing a standardized, reusable approach to security assessment and authorization for cloud computing products and services that process unclassified information used by agencies.

The FSCAC will provide advice and recommendations to the Administrator of GSA, the FedRAMP Board, and agencies on technical, financial, programmatic, and operational matters regarding the secure adoption of cloud computing products and services. The FSCAC will ensure effective and ongoing coordination of agency

adoption, use, authorization, monitoring, acquisition, and security of cloud computing products and services to enable agency mission and administrative priorities. The purposes of the Committee are:

- To examine the operations of FedRAMP and determine ways that authorization processes can continuously be improved, including the following:
  - Measures to increase agency reuse of FedRAMP authorizations.
  - Proposed actions that can be adopted to reduce the burden, confusion, and cost associated with FedRAMP authorizations for cloud service providers.
  - Measures to increase the number of FedRAMP authorizations for cloud computing products and services offered by small businesses concerns (as defined by section 3(a) of the Small Business Act (15 U.S.C. 632(a)).
  - Proposed actions that can be adopted to reduce the burden and cost of FedRAMP authorizations for agencies.
- Collect information and feedback on agency compliance with, and implementation of, FedRAMP requirements.
- Serve as a forum that facilitates communication and collaboration among the FedRAMP stakeholder community.

The FSCAC will meet no fewer than three (3) times a calendar year. Meetings shall occur as frequently as needed, called, and approved by the DFO.

Purpose of the Meeting and Agenda

The March 28, 2024 public meeting will be dedicated to deliberations in order to determine what priority or priorities the Committee would like to work on next. Presentations may be held on updates to the Office of Management and Budget's (OMB) draft Memorandum titled ''Modernizing the Federal Risk Authorization Management Program (FedRAMP)'' (OMB Draft Memo), FedRAMP's updates in response to the OMB Draft Memo, and Third Party Assessment Organization (3PAO) user experiences with the FedRAMP process. A vote will be held to approve the priority or priorities the Committee chooses to work on next. The meeting agenda will be posted on *https://gsa.gov/fscac* prior to the March 28, 2024 meeting.

Meeting Attendance

This meeting is open to the public and can be attended in-person or virtually using the live stream link. Meeting registration and information is available at *https://gsa.gov/fscac.* Registration for attending the meeting in person is highly encouraged by 5 p.m. on Thursday, March 21, 2024 for easier building access. In-person public attendance is limited to the available space, and seating is available on a first come, first serve basis.

If you plan to attend virtually, you will need to register by 5 p.m. on Thursday, March 21, 2024 to obtain the virtual meeting information. After registration, individuals will receive meeting attendance information via email.

For information on services for individuals with disabilities, or to request accommodation for a disability, please email the FSCAC staff at *FSCAC@gsa.gov* at least 10 days prior to the meeting. Live captioning may be provided virtually, and ASL interpreters may be present onsite.

Public Comment

Members of the public will have the opportunity to provide oral public comment during the FSCAC meeting by indicating their preference when registering. Written public comments can be submitted at any time by completing the public comment form on our website, *https://gsa.gov/fscac.* All written public comments will be provided to FSCAC members in advance of the meeting if received by Wednesday, March 20, 2024.

Margaret Dugan,

Service-Level Liaison, Federal Acquisition Service, General Services Administration.

[FR Doc. 2024–04844 Filed 3–6–24; 8:45 am]

BILLING CODE 6820–34–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

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 updates to the currently approved information collection project: ''Implementation and Testing of Diagnostic Safety Resources.'' In accordance with the Paperwork Reduction Act of 1995, AHRQ invites the public to comment on this proposed information collection.

DATES: Comments on this notice must be received by May 6, 2024.

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

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

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

SUPPLEMENTARY INFORMATION:

Proposed Project

Implementation and Testing of Diagnostic Safety Resources

Patient safety is a pillar of the Agency for Healthcare Research and Quality's (AHRQ's) mission to support the highest quality healthcare. While progress has been made in many areas of patient safety, the field of diagnostic safety has emerged as a particular area of concern. It is estimated that every person in the United States will experience a diagnostic error in their lifetime (Institute of Medicine, 2015) which can lead to inappropriate, delayed, or withheld treatment and ultimately poor health outcomes, distress, and increased costs. Diagnostic errors can occur for many reasons: lack of meaningful engagement between clinicians, patients, and families; a fragmented healthcare system not designed to account for an increasingly complex diagnostic process; minimal (if any) feedback to clinicians about their diagnostic performance; and a culture that does not always support transparent disclosure of diagnostic errors (Institute of Medicine, 2015). Leaders in diagnostic excellence suggest that multi-pronged efforts are needed to address this complex problem and go beyond individual behaviors to system-level changes and empowering patients to engage in their care (Institute of Medicine, 2015; Henriksen, et al., 2017).

Improving diagnostic safety and quality is an AHRQ priority. In recognition of the multifaceted approach needed to effectively advance diagnostic safety, AHRQ recently supported the development of three tools to prevent diagnostic errors and have prioritized these tools for implementation and testing. These resources vary in the types of stakeholders they target, a critical