Participation.'' A copy of the comments may also be submitted to the OMB desk officer for the FDIC: Office of Information and Regulatory Affairs, Office of Management and Budget, New Executive Office Building, Washington, DC 20503.

FOR FURTHER INFORMATION CONTACT: Jennifer Jones, at the FDIC mailing address above or by phone at 202–898–6768.

SUPPLEMENTARY INFORMATION:

Proposal for the Following New Generic Collection of Information

1. *Title:* Generic Clearance for Prize Competition Participation.

OMB Number: 3064–NEW.

Form Number: None.

Affected Public: Innovators; technologists, coders, engineers and developers; consumers of financial services; consumer advocates; academics; members of trade groups and other associations; individuals connected to financial institutions, community banks, and financial and bank service and technology providers; software, data, and technology firms; and other members of the public.

Estimated Burden per Prize Competition:

Estimated Annual Number of Respondents: 300.

Estimated Average Time per Response: 20 hours.

Total Estimated Annual Burden per Prize Competition: 6,000 hours.

General Description of Collection: The FDIC seeks generic clearance for the collection of information requested from potential participants (including innovators; technologists, coders, engineers and developers; consumers of financial services; consumer advocates; academics; members of trade groups and other associations; individuals connected to financial institutions, community banks, and financial and bank service and technology providers; software, data, and technology firms; and other members of the public) with respect to solicitations for expressions of interest to participate in FDIC-sponsored or co-sponsored prize competitions of various types, including point solution competitions (designed to spur the development of solutions for a particular problem) and exposition (designed competitions to identify and promote a broad range of ideas and practices to facilitate further development by third parties). Prize competitions and the opportunity to submit applications to participate will be announced on the agency's publicly accessible government website, as well as possibly through other forms of public communication, such as publication in the **Federal Register**, issuance of Financial Institution Letters, use of *challenge.gov* website maintained by the U.S. General Services Administration, or social media advertisement.

In order for the FDIC to determine which applicants will be eligible and selected to participate in FDIC prize competitions, the FDIC will request that potential participants provide their name, contact information, address, and such other information that may be necessary to evaluate applicants' qualifications and ability to participate in the event as well as to match the applicants' anticipated role to the needs of the competition. Applicants will also be asked to acknowledge the terms and conditions of participating in the prize competition. Information will be collected during prize competitions through the solutions to the challenges or problems presented.

This information collection will be voluntary. Collection in the form of application will be conducted primarily online with alternative methods made available. Collection during the events will be in-person or electronic. The FDIC will consult with OMB regarding each specific information collection during the approval period.

The FDIC estimates that over the three-year clearance period of this request, up to five (5) competitions will be conducted across various divisions of the agency, involving a variety of topics and challenges associated with underserved communities and financial inclusion; consumer protection; the FDIC's use of information technology and data (including artificial intelligence and machine learning); and financial and technologically-driven innovation in banking. The total hourly burden attributed to this generic clearance will be 30,000 hours (6,000 hours per prize competition × 5 competitions per year).

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 estimate 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.

Dated at Washington, DC, on June 20, 2019.

Federal Deposit Insurance Corporation

Valerie Best,

Assistant Executive Secretary.

[FR Doc. 2019–13477 Filed 6–24–19; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL RESERVE SYSTEM

Proposed Agency Information Collection Activities; Comment Request

AGENCY: Board of Governors of the Federal Reserve System.

ACTION: Notice; request for comment.

SUMMARY: The Board of Governors of the Federal Reserve System (Board) invites comment on a proposal to extend for three years, with revision, the Interagency Policy Statement on Funding and Liquidity Risk Management (FR 4198; OMB No. 7100–0326).

DATES: Comments must be submitted on or before August 26, 2019.

ADDRESSES: You may submit comments, identified by *FR 4198,* by any of the following methods:

- *Agency Website: http://www.federalreserve.gov.* Follow the instructions for submitting comments at *http://www.federalreserve.gov/apps/foia/proposedregs.aspx.*
- *Email: regs.comments@federalreserve.gov.* Include OMB number in the subject line of the message.
- *FAX:* (202) 452–3819 or (202) 452–3102.
- *Mail:* Ann E. Misback, Secretary, Board of Governors of the Federal Reserve System, 20th Street and Constitution Avenue NW, Washington, DC 20551.

All public comments are available on the Board's website at *https://www.federalreserve.gov/apps/foia/proposedregs.aspx* as submitted, unless modified for technical reasons. Accordingly, your comments will not be edited to remove any identifying or contact information. Public comments may also be viewed electronically or in paper form in Room 146, 1709 New York Avenue NW, Washington, DC 20006, between 9:00 a.m. and 5:00 p.m. on weekdays. For security reasons, the Board requires that visitors make an appointment to inspect comments. You may do so by calling (202) 452–3684. Upon arrival, visitors will be required to present valid government-issued photo identification and to submit to security

screening in order to inspect and photocopy comments.

Additionally, commenters may send a copy of their comments to the Office of Management and Budget (OMB) Desk Officer—Shagufta Ahmed—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.

FOR FURTHER INFORMATION CONTACT: A copy of the Paperwork Reduction Act (PRA) OMB submission, including the proposed reporting form and instructions, supporting statement, and other documentation will be placed into OMB's public docket files, if approved. These documents will also be made available on the Board's public website at *http://www.federalreserve.gov/apps/reportforms/review.aspx* or may be requested from the agency clearance officer, whose name appears below.

Federal Reserve Board Clearance Officer—Nuha Elmaghrabi—Office of the Chief Data Officer, Board of Governors of the Federal Reserve System, Washington, DC 20551, (202) 452–3829.

SUPPLEMENTARY INFORMATION: On June 15, 1984, OMB delegated to the Board authority under the PRA to approve and assign OMB control numbers to collection of information requests and requirements 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.

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, With Revision, the Following Information Collection

Report title: Interagency Policy Statement on Funding and Liquidity Risk Management.

Agency form number: FR 4198.

OMB control number: 7100–0326.

Frequency: Annually.

Respondents: Bank holding companies, savings and loan holding companies, state-licensed branches and agencies of foreign banks (other than insured branches), corporations organized or operating under sections 25 or 25A of the Federal Reserve Act (agreement corporations and Edge corporations), and state member banks (collectively, financial institutions).

Estimated number of respondents: Implementing recordkeeping, 30; ongoing recordkeeping, 4,789.

Estimated average hours per response: Implementing recordkeeping, 160 hours; ongoing recordkeeping, 32 hours.

Estimated annual burden hours: 158,048 hours.

General description of report: The Interagency Policy Statement on Funding and Liquidity Risk Management (Guidance) [1] states that financial institutions should develop and document liquidity risk management policies and procedures commensurate with the institution's complexity, risk profile, and scope of operations. Sections 3 and 6 of the Guidance provide that financial institutions should maintain such policies and procedures. Section 6 of the Guidance states that financial institutions should have a contingency funding plan (CFP) that sufficiently addresses potential adverse liquid events and emergency cash flow requirements, and section 34 of the Guidance states that the CFP should be documented.

Proposed revisions: The Board is proposing to revise the FR 4198 to account for all of the recordkeeping provisions set forth in the Guidance related to liquidity risk management policies, procedures, and assumptions, and CFPs. The FR 4198 currently does not account for the recordkeeping provisions related to CFPs and does not fully account for the recordkeeping provisions related to liquidity risk management policies, procedures, and assumptions.

Legal authorization and confidentiality: The recordkeeping provisions of the Guidance are authorized pursuant to sections 9(6), 25, and 25A of the Federal Reserve Act [2] (for state member banks, agreement corporations, and Edge corporations, respectively); section 5(c) of the Bank Holding Company Act [3] (for bank holding companies); section 10(b)(3) of the Home Owners' Loan Act [4] (savings and loan holding companies); and section 7(c)(2) of the International Banking Act [5] (state-licensed branches and agencies of foreign banks, other than insured branches). Because the recordkeeping provisions are contained within guidance, which is nonbinding, they are voluntary.[6] There are no reporting forms associated with this information collection.

Because these records would be maintained at each banking organization, the Freedom of Information Act (FOIA) would only be implicated if the Board obtained such records as part of the examination or supervision of a banking organization. In the event the records are obtained by the Board as part of an examination or supervision of a financial institution, this information may be considered confidential pursuant to exemption 8 of the FOIA, which protects information contained in ''examination, operating, or condition reports'' obtained in the bank supervisory process (5 U.S.C. 552(b)(8)). In addition, the information may also be kept confidential under exemption 4 for the FOIA, which protects ''commercial or financial information obtained from a person [that is] privileged or confidential'' (5 U.S.C. 552(b)(4)).

Consultation outside the agency: The Guidance was published jointly by the Board, the Office of the Comptroller of

[1] ''Interagency Policy Statement on Funding and Liquidity Risk Management,'' 75 FR 13656 (March 22, 2010). The Guidance was published jointly by the Board, the Office of the Comptroller of the Currency, the Office of Thrift Supervision, the Federal Deposit Insurance Corporation, and the National Credit Union Administration.

[2] 12 U.S.C. 324, 602, and 625, respectively.

[3] 12 U.S.C. 1844(c).

[4] 12 U.S.C. 1467a(b)(3).

[5] 12 U.S.C. 3105(c)(2).

[6] *See* SR 18–5/CA 18–7: Interagency Statement Clarifying the Role of Supervisory Guidance (Sept. 11, 2018).

the Currency, the Office of Thrift Supervision, the Federal Deposit Insurance Corporation, and the National Credit Union Administration. There has been no consultation outside of the Federal Reserve System with regard to the current proposal to extend the FR 4198 for three years, with revision.

Board of Governors of the Federal Reserve System, June 20, 2019.

Michele Taylor Fennell,

Assistant Secretary of the Board.

[FR Doc. 2019–13490 Filed 6–24–19; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1147]

Agency Information Collection Activities; Proposed Collection; Comment Request; Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. 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 of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions of our guidance document entitled ''Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition.''

DATES: Submit either electronic or written comments on the collection of information by August 26, 2019.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before August 26, 2019. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of August 26, 2019. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

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–2013–N–1147 for ''Preparing a Claim of Categorical Exclusion or an Environmental Assessment for Submission to the Center for Food Safety and Applied Nutrition.'' Received comments, those filed in a timely manner (see **ADDRESSES**), 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.

- 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.gpo.gov/fdsys/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://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: 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. ''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 (44 U.S.C. 3506(c)(2)(A)) requires Federal