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. Telecommunications Device for the Deaf (TDD) users may contact (202) 263–4869, Board of Governors of the Federal Reserve System, Washington, DC, 20551.

SUPPLEMENTARY INFORMATION: On June 15, 1984, the Office of Management and Budget (OMB) delegated to the Board authority under the Paperwork Reduction Act (PRA) to approve of 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 Federal Reserve's functions; including whether the information has practical utility;

b. The accuracy of the Federal Reserve'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 Federal Reserve should modify the proposal prior to giving final approval.

Proposal To Approve Under OMB Delegated Authority the Extension for Three Years, With Revision, of the Following Report

Report title: Recordkeeping and Disclosure Requirements Associated with Consumer Financial Protection Bureau's (CFPB) Regulation M (Consumer Leasing).

Agency form number: FR M.

OMB control number: 7100–0202.

Frequency: Disclosures, 461 times per year; and Advertising, quarterly.

Respondents: State member banks with assets of $10 billion or less that are not affiliated with an insured depository institution with assets over $10 billion (irrespective of the consolidated assets of any holding company); non-depository affiliates of such state member banks; and non-depository affiliates of bank holding companies that are not affiliated with an insured depository institution with assets over $10 billion.

Estimated number of respondents: 5.

Estimated average hours per response: Disclosures, 6.5 minutes; and advertising, 25 minutes.

Estimated annual burden hours: Disclosures, 250 hours; and advertising, 8 hours.

General description of report: The CLA and Regulation M require lessors uniformly to disclose to consumers the costs, liabilities, and terms of consumer lease transactions. Disclosures are provided to consumers before they enter into lease transactions and in advertisements that state the availability of consumer leases on particular terms. The regulation generally applies to consumer leases of personal property in which the contractual obligation does not exceed $50,000, adjusted annually for inflation, and has a term of more than four months.[1] The CLA does not provide exemptions for small entities.

Proposed revisions: The Board proposes to revise the methodology for estimating burden for disclosures to provide additional clarity and transparency into the calculation. Specifically, the Board proposes to estimate disclosure burden using the estimated average number of lease contracts each Board-supervised institution initiates annually, assuming it takes approximately 6.5 minutes to populate and provide each disclosure.

Legal authorization and confidentiality: The Board's Legal Division has determined that sections 105(a) and 187 of TILA (15 U.S.C. 1604(a) and 1667f respectively, authorize the CFPB to issue regulations to carry out the provisions of the CLA. The CFPB's Regulation M, 12 CFR part 1013, implements these statutory provisions. An institution's recordkeeping and disclosure obligations under Regulation M are mandatory. Because the Board does not collect any information pursuant to the CFPB's Regulation M, no issue of confidentiality normally arises. In the event the Board were to retain information regarding consumer leases during the course of an examination, the information regarding the consumer and the lease would be kept confidential pursuant to section (b)(8) of the Freedom of Information Act (5 U.S.C. 522 (b)(8)).

[1] For 2018, the Regulation M threshold is $55,800.

Board of Governors of the Federal Reserve System, April 2, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–06947 Filed 4–4–18; 8:45 am]

BILLING CODE 6210–01–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, without revision, the voluntary Federal Reserve Clearance for Board Public website Usability Surveys (FR 3076, OMB No. 7100–0366). On June 15, 1984, the Office of Management and Budget (OMB) delegated to the Board authority under the Paperwork Reduction Act (PRA) to approve of 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.

DATES: Comments must be submitted on or before June 4, 2018.

ADDRESSES: You may submit comments, identified by FR 3076, 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 from the Board's website at *http://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 3515, 1801 K Street (between 18th and 19th Streets 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 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 PRA OMB submission, including the proposed reporting form and instructions, supporting statement, and other documentation will be placed into OMB's public docket files, once approved. These documents will also be made available on the Federal Reserve 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. Telecommunications Device for the Deaf (TDD) users may contact (202) 263–4869, Board of Governors of the Federal Reserve System, Washington, DC 20551.

SUPPLEMENTARY INFORMATION:

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 Federal Reserve's functions; including whether the information has practical utility;

b. The accuracy of the Federal Reserve'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 Federal Reserve should modify the proposal prior to giving final approval.

Proposal To Approve Under OMB Delegated Authority the Extension for Three Years, Without Revision, of the Following Report

Report title: Federal Reserve Clearance for Board Public website Usability Surveys.

Agency form number: FR 3076.

OMB control number: 7100–0366.

Frequency: As needed.

Respondents: Individuals.

Estimated number of respondents: Surveys: 100, Focus Groups: 20.

Estimated average hours per response: Surveys: 0.25, Focus Groups: 1.5.

Estimated annual burden hours: 420.

General description of report: The FR 3076 is used to gather qualitative and quantitative information directly from users or potential users of the Board's website such as Congress, other government agencies, the public, economic educators, economists, financial institutions, financial literacy groups, and community development groups and more. Participation is voluntary.

The FR 3076 may seek information from users or potential users of various Board web pages, including press releases, data releases and downloads, reports, supervision manuals, brochures, new web pages, audio, video, and use of social media. Information gathered may also include general input on users' interests and needs, feedback on website navigation and layout, distribution channels, or other factors which may affect the ability of users to locate and access content online.

Qualitative surveys conducted using the FR 3076 would include data gathering methods such as focus groups and individual interviews. Quantitative surveys conducted using the FR 3076 would include surveys conducted online or via mobile device, by phone or by mail, emails, or a combination of these methods. The Board may contract with an outside vendor to conduct focus groups, interviews, or surveys, or the Board may collect the data directly. As the Board's public website continues to evolve, the Board may seek input from users or potential users of Board's public website on questions such as the following:

• Did you find the content and layout relevant and of value?

• How did you find the content you were looking for?

• Was the navigation useful?

• How did you learn about the content?

• How did you access the content? (*e.g.:* paper copy distributed at an event, online, or mobile device). If online or through a mobile device, was the document printed, viewed on a tablet, or on a computer screen?

• What suggestions do you have for improving the format and appearance of online presentation? (*e.g.:* readability—font size, charts, and graphs; organization of information; and navigating—indexing, search tools, and links)

• What other information would be of value to enhance the online tool or information?

Legal authorization and confidentiality: The Board uses its website and social media to communicate important information to the public about a variety of different issues. The Board is required to provide certain information on its website. For example, under section 2B of the Federal Reserve Act the Board is required to provide certain reports, audits, and other information that ''the Board reasonably believes is necessary or helpful to the public in understanding the accounting, financial reporting, and internal controls of the Board and the Federal reserve banks.'' 12 U.S.C. 225b(c). In addition, the Board uses its website to provide the public with information about a variety of other matters, including information

about the Board, its actions, and the economy. The responses to the FR 3076 help the Board determine how to most effectively communicate this information to the public in order to fulfill its statutory responsibilities. The FR 3076 is voluntary. The information collected by the FR 3076 is not considered to be confidential.

Board of Governors of the Federal Reserve System, April 2, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–06948 Filed 4–4–18; 8:45 am]

BILLING CODE 6210–01–P

GOVERNMENT ACCOUNTABILITY OFFICE

Request for Nominations for Board of Governors of the Patient-Centered Outcomes Research Institute (PCORI)

AGENCY: Government Accountability Office (GAO).

ACTION: Request for letters of nomination and resumes.

SUMMARY: The Patient Protection and Affordable Care Act gave the Comptroller General of the United States responsibility for appointing 19 members to the Board of Governors of the Patient-Centered Outcomes Research Institute. In addition, the Directors of the Agency for Healthcare Research and Quality and the National Institutes of Health, or their designees, are members of the Board. As the result of terms ending in September 2018, GAO is accepting nominations in the following categories required in statute: A physician, a nurse, a representative of patients and health care consumers, a representative of private payers, a representative of a state or a federal health program or agency, and a representative of pharmaceutical, device, or diagnostic manufacturers or developers. Nominations should be sent to the email or mailing address listed below. Acknowledgement of submissions will be provided within a week of submission.

DATES: Letters of nomination and resumes should be submitted no later than May 4, 2018, to ensure adequate opportunity for review and consideration of nominees prior to appointment.

ADDRESSES: Submit letters of nomination and resumes by either of the following methods: Email: *PCORI@gao.gov.* Include PCORI Nominations in the subject line of the message, or Mail: U.S. GAO, Attn: PCORI Board Nominations, 441 G Street NW, Washington, DC 20548.

FOR FURTHER INFORMATION CONTACT: Rashmi Agarwal at (202) 512–4077 or *agarwalr@gao.gov* if you do not receive an acknowledgement or need additional information. For general information, contact GAO's Office of Public Affairs, (202) 512–4800.

Authority: [Sec. 6301 and Sec. 10602, Pub. L. 111–148].

Gene L. Dodaro,

Comptroller General of the United States.

[FR Doc. 2018–06999 Filed 4–4–18; 8:45 am]

BILLING CODE 1610–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2002–D–0093]

Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation; Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a guidance for industry entitled ''Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation.'' This guidance document finalizes the revised draft of the same name that published on October 30, 2015. This guidance provides recommendations to applicants on the chemistry, manufacturing, and controls (CMC); pharmacokinetics and bioavailability; and labeling documentation for liposome drug products submitted in new drug applications (NDAs) and abbreviated new drug applications (ANDAs), reviewed by the Center for Drug Evaluation and Research (CDER).

DATES: The announcement of the guidance is published in the **Federal Register** on April 5, 2018.

ADDRESSES: You may submit either electronic or written comments on Agency guidances 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–2002–D–0093 (formerly 2002D–0337) for ''Liposome Drug Products: Chemistry, Manufacturing, and Controls; Human Pharmacokinetics and Bioavailability; and Labeling Documentation.'' 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.

- 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