2. Activity-Based Stock Purchase Requirement Submissions

FHFA estimates that the average number of daily transactions between Banks and members that will require the exchange of information to confirm the member's activity-based stock purchase requirement will be 300, and that there will be an average of 261 working days per year, resulting in an estimated 78,300 submissions annually. The estimate for the average preparation time per submission is 0.2 hours. Accordingly, the estimate for the annual hour burden associated with activity-based stock purchase requirement submissions is (78,300 submissions × 0.2 hours per submission) = 15,660 hours.

E. Comment Request

In accordance with the requirements of 5 CFR 1320.8(d), FHFA published an initial notice and request for public comments regarding this information collection in the **Federal Register** on August 8, 2023.[4] The 60-day comment period closed on October 10, 2023. FHFA received no substantive comments.

FHFA requests written comments on the following: (1) whether the collection of information is necessary for the proper performance of FHFA functions, including whether the information has practical utility; (2) the accuracy of FHFA's estimates of the burdens of the collection of information; (3) ways to enhance the quality, utility, and clarity of the information collected; and (4) 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.

Shawn Bucholtz,

Chief Data Officer, Federal Housing Finance Agency.

[FR Doc. 2023–23067 Filed 10–18–23; 8:45 am]

BILLING CODE 8070–01–P

FEDERAL RESERVE SYSTEM

Sunshine Meeting Notice

AGENCY HOLDING THE MEETING: Board of Governors of the Federal Reserve System.

TIME AND DATE: 1:00 p.m. on Wednesday, October 25, 2023.

PLACE: Martin Federal Reserve Board Building, C Street entrance between 20th and 21st Streets NW, Washington, DC 20551.

STATUS: Open.

On the day of the meeting, you will be able to view the meeting via webcast from a link available on the Board's website. You do not need to register to view the webcast of the meeting. A link to the meeting documentation will also be available approximately 20 minutes before the start of the meeting. Both links may be accessed from the Board's website at *www.federalreserve.gov*.

If you plan to attend the open meeting in person, we ask that you notify us in advance and provide your name, date of birth, and social security number (SSN) or passport number. You may provide this information by calling 202–452–2474 or you may register online *www.federalreserve.gov*. You may pre-register until close of business on October 24, 2023. You also will be asked to provide identifying information, including a photo ID, before being admitted to the Board meeting. The Public Affairs Office must approve the use of cameras; please email *media@frb.gov* for further information. If you need an accommodation for a disability, please contact Penelope Beattie on 202–452–3982. For users of telephone systems via text telephone (TTY) or any TTY-based Telecommunications Relay Services (TRS), please call 202–263–4869 or dial 7–1–1 from any telephone, anywhere in the United States.

Privacy Act Notice: The information you provide will be used to assist us in prescreening you to ensure the security of the Board's premises and personnel. In order to do this, we may disclose your information consistent with the routine uses listed in the Privacy Act Notice for BGFRS–32, including to appropriate federal, state, local, or foreign agencies where disclosure is reasonably necessary to determine whether you pose a security risk or where the security or confidentiality of your information has been compromised. We are authorized to collect your information by 12 U.S.C. 243 and 248, and Executive Order 9397. In accordance with Executive Order 9397, we collect your SSN so that we can keep accurate records, because other people may have the same name and birth date. In addition, we use your SSN when we make requests for information about you from law enforcement and other regulatory agency databases. Furnishing the information requested is voluntary; however, your failure to provide any of the information requested may result in disapproval of your request for access to the Board's premises. You may be subject to a fine or imprisonment under 18 U.S.C. 1001 for any false statements you make in your request to enter the Board's premises.

MATTERS TO BE CONSIDERED:

Discussion Agenda

1. Proposed revisions to the Board's debit interchange fee cap.

Notes: 1. For those attending in person, the staff memo will be available to attendees on the day of the meeting in paper. Meeting documentation will be available on the Board's website about 20 minutes before the start of the meeting.

2. This meeting will be recorded for the benefit of those unable to attend. The webcast recording and a transcript of the meeting will be available after the meeting on the Board's website *http://www.federalreserve.gov/aboutthefed/boardmeetings/*.

For questions please contact: Public Affairs Office at *media@frb.gov*.

SUPPLEMENTARY INFORMATION: You may access the Board's website at *www.federalreserve.gov* for an electronic announcement. (The website also includes procedural and other information about the open meeting.)

Dated: October 16, 2023.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2023–23116 Filed 10–18–23; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2023–D–4356]

Enforcement Policy for Non-Invasive Remote Monitoring Devices Used To Support Patient Monitoring; Guidance for Industry and Food and Drug Administration Staff; 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 final guidance entitled ''Enforcement Policy for Non-Invasive Remote Monitoring Devices Used to Support Patient Monitoring.'' The enforcement policy described in this guidance applies to modified devices where the original device was a legally marketed, non-invasive remote monitoring device listed in the guidance that measures or detects common physiological parameters and that is used to support patient monitoring. The guidance is intended to describe the enforcement policy for limited modifications to the indications, functionality, or hardware

[4] *See* 88 FR 53484 (Aug. 8, 2023).