

Washington, DC 20551-0001, not later than August 4, 2022.

*A. Federal Reserve Bank of Kansas City (Jeffrey Imgarten, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198-0001:* 1. J. Scott Schrempp, Hartington, Nebraska, Christine Rossiter, Elkhorn, Nebraska, and Mary Rossiter, Macon, Georgia; to become members of Rossiter Family Control Group, a group acting in concert, to retain voting shares of Cedar Bancorp, and thereby indirectly retain voting shares of Bank of Hartington, both of Hartington, Nebraska.

*B. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414:* 1. The ATPB Exempt Bank Trust, and the ATPB Non-Exempt Bank Trust, both of La Crosse, Wisconsin, Andrew R. Bosshard, La Crosse, Wisconsin, and Ashley B. Sawyer, Washington, DC, as co-trustees to both trusts, Alexandra Tana Pizitz Bosshard, Washington, DC, as investment advisor and with power to appoint or remove trustees, and Elizabeth Bosshard-Blackey, Edina, Minnesota, as trust protector and with power to appoint or remove trustees; to become members of the Bosshard Family Control Group, a group acting in concert, to acquire voting shares of Bosshard Financial Group, Inc., La Crosse, Wisconsin, and thereby indirectly acquire voting shares of One Community Bank, Oregon, Wisconsin, and Farmers State Bank-Hillsboro, Hillsboro, Wisconsin.

Board of Governors of the Federal Reserve System.

**Margaret McCloskey Shanks,**  
*Deputy Secretary of the Board.*

[FR Doc. 2022-15508 Filed 7-19-22; 8:45 am]

**BILLING CODE P**

## FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

### Notice of Board Meeting

**DATES:** July 26, 2022 at 10 a.m.

**ADDRESSES:** Telephonic. Dial-in (listen only) information: Number: 1-202-599-1426, Code: 303 807 400#; or via web: <https://teams.microsoft.com/join/19%3ameeting-NjVmZmM5NzktM2Q5NS00MmYzLWFlNmYtMjBhNTEExNTdkYTM3%40thread.v2/0?context=%7b%22Tid%22%3a%223f6323b7-e3fd-4f35-b43d-1a7afae5910d%22%2c%22Oid%22%3a%227c8d802c-5559-41ed-9868-8bfad5d44af9%22%7d>

**FOR FURTHER INFORMATION CONTACT:** Kimberly Weaver, Director, Office of External Affairs, (202) 942-1640.

### SUPPLEMENTARY INFORMATION:

#### Board Meeting Agenda

##### Open Session

1. Approval of the June 28, 2022 Board Meeting Minutes
2. Monthly Reports
  - (a) Participant Activity Report
  - (b) Legislative Report
3. Quarterly Report
  - (c) Investment Policy
  - (d) Budget Review
  - (e) Audit Status
4. Internal Audit Update

##### Closed Session

5. Information covered under 5 U.S.C. 552b (c)(9)(B).

*Authority:* 5 U.S.C. 552b (e)(1).

Dated: July 14, 2022.

**Dharmesh Vashee,**

*General Counsel, Federal Retirement Thrift Investment Board.*

[FR Doc. 2022-15438 Filed 7-19-22; 8:45 am]

**BILLING CODE 6760-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifiers: CMS-10507]

#### Agency Information Collection Activities: Submission for OMB Review; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. 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 or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of

the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments on the collection(s) of information must be received by the OMB desk officer by August 19, 2022.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' website address at website address at: <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing>.

**FOR FURTHER INFORMATION CONTACT:** William Parham at (410) 786-4669.

**SUPPLEMENTARY INFORMATION:** Under the Paperwork Reduction Act of 1995 (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. The term "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 agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* State-based Exchange Annual Report Tool (SMART); *Use:* The annual report is the primary vehicle to insure comprehensive compliance with all reporting requirements contained in the Affordable Care Act (ACA). It is

specifically called for in Section 1313(a)(1) of the Act which requires a State Based Exchange (including an Exchange using the Federal Platform) to keep an accurate accounting of all activities, receipts, and expenditures, and to submit a report annually to the Secretary concerning such accounting. CMS will use the information collected from States to assist in determining if a State is maintaining a compliant operational Exchange. *Form Number:* CMS-10507 (OMB control number: 0938-1244); *Frequency:* Annually; *Affected Public:* State, Local, or Tribal governments; *Number of Respondents:* 21; *Total Annual Responses:* 21; *Total Annual Hours:* 4,281. (For policy questions regarding this collection contact Shilpa Gogna at 301-492-4257.)

Dated: July 14, 2022.

**William N. Parham, III,**

*Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2022-15404 Filed 7-19-22; 8:45 am]

**BILLING CODE 4120-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA-2011-N-0921]

#### **Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Standards for the Growing, Harvesting, Packaging, and Holding of Produce for Human Consumption**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or we) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Submit written comments (including recommendations) on the collection of information by August 19, 2022.

**ADDRESSES:** To ensure that comments on the information collection are received, OMB recommends that written comments be submitted to [https://](https://www.reginfo.gov/public/do/PRAMain)

[www.reginfo.gov/public/do/PRAMain](https://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting “Currently under Review—Open for Public Comments” or by using the search function. The OMB control number for this information collection is 0910-0816. Also include the FDA docket number found in brackets in the heading of this document.

#### **FOR FURTHER INFORMATION CONTACT:**

Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A-12M, 11601 Landsdown St., North Bethesda, MD 20852, 301-796-8867, [PRAStaff@fda.hhs.gov](mailto:PRAStaff@fda.hhs.gov).

#### **SUPPLEMENTARY INFORMATION:**

In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

#### **Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption; 21 CFR Part 112**

*OMB Control Number 0910-0816—Extension*

To minimize the risk of serious adverse health consequences or death from consumption of contaminated produce, we have established science-based minimum standards for the safe growing, harvesting, packing, and holding of produce, meaning fruits and vegetables grown for human consumption. The standards are codified in part 112 (21 CFR part 112) and set forth procedures and processes that include information collection activities such as establishing monitoring and sampling plans, documenting data and training, and ensuring disclosure that produce for human consumption meets these requirements. The regulations also provide for certain exemptions and variances to qualified respondents. The information collection continues to implement provisions of the FDA Food Safety Modernization Act, while certain requirements for covered produce other than sprouts associated with pre-harvest agricultural water testing are being amended through rulemaking (RIN 0910-AI49). We use the information to verify that the standards established by the regulations are followed such that produce entering the marketplace is reasonably unlikely to be associated with foodborne illness.

In addition to the referenced regulations, we have developed two draft guidance documents: “Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption” and “Compliance with and Recommendations for Implementation of the Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption for Sprout Operations;” both are available at <https://www.fda.gov/Food/GuidanceRegulation/GuidanceDocumentsRegulatoryInformation/default.htm>. The former was developed to help covered farms comply with the requirements of the Produce Safety regulation. This draft guidance, when finalized, will not create any additional burden not already considered as part of the Produce Safety regulation.

The latter (the Sprouts draft guidance) was developed to assist sprout operations also subject to the Produce Safety regulation. Sprouts represent a special food safety concern because the conditions under which they are produced (time, temperature, water activity, pH, and available nutrients) are ideal for the growth of pathogens, if present. The Sprouts draft guidance, when finalized, will assist sprout operations subject to the regulations in part 112 in complying with the sprout-specific requirements in subpart M.

#### *Description of Respondents:*

Respondents to this information collection include farms that grow, harvest, pack, or hold produce for human consumption, meaning fruits and vegetables such as berries, tree nuts, herbs, and sprouts. Respondents are from the private sector (for-profit businesses).

In the **Federal Register** of December 3, 2021 (86 FR 68673), FDA published a 60-day notice requesting public comment on the proposed collection of information. One comment was received and appears to pertain to rulemaking that has already concluded, rather than to this renewal. Significantly, this comment did not suggest that we revise the currently approved estimate. To the extent that the comment relates to ongoing rulemaking, we have posted the comment to the docket at FDA-2021-N-0471 and will ensure it is considered and addressed appropriately.

We estimate the burden of this collection of information as follows: