FEDERAL RESERVE SYSTEM

Agency Information Collection Activities: Announcement of Board Approval Under Delegated Authority and Submission to OMB

AGENCY: Board of Governors of the Federal Reserve System.

SUMMARY: The Board of Governors of the Federal Reserve System (Board) is adopting a proposal to extend for three years, with revision, the Request for Extension of Time to Dispose of Assets Acquired in Satisfaction of Debts Previously Contracted (FR 4006; OMB No. 7100–0129).

FOR FURTHER INFORMATION CONTACT: Federal Reserve Board Clearance Officer—Nuha Elmaghrabi—Office of the Chief Data Officer, Board of Governors of the Federal Reserve System, *nuha.elmaghrabi@frb.gov,* (202) 452–3884.

Office of Management and Budget (OMB) Desk Officer for the Federal Reserve Board, 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.

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. Board-approved collections of information are incorporated into the official OMB inventory of currently approved collections of information. The OMB inventory, as well as copies of the PRA Submission, supporting statements, and approved collection of information instrument(s) are available at *https://www.reginfo.gov/public/do/PRAMain.* These documents are also available on the Federal Reserve Board's public website at *https://www.federalreserve.gov/apps/reportforms/review.aspx* or may be requested from the agency clearance officer, whose name appears above.

Final Approval Under OMB Delegated Authority of the Extension for Three Years, With Revision, of the Following Information Collection

Collection title: Request for Extension of Time to Dispose of Assets Acquired in Satisfaction of Debts Previously Contracted.

Collection identifier: FR 4006.

OMB control number: 7100–0129.

Effective Date: The revisions are applicable as of August 2, 2022.

Frequency: On occasion.

Respondents: Bank holding companies (BHCs).

Estimated number of respondents: Section 225.12(b), 1; Section 225.22(d)(1), 20; Section 225.140(c) and (d), 12.

Estimated average hours per response: Section 225.12(b), 5; Section 225.22(d)(1), 5; Section 225.140(c) and (d), 2.

Estimated annual burden hours: Section 225.12(b), 5; Section 225.22(d)(1), 100; Section 225.140(c) and (d), 24.

General description of report: The Bank Holding Company Act of 1956 (BHC Act) and the Board's Regulation Y (12 CFR part 225) require a bank holding company that, either through foreclosure or otherwise in the ordinary course of collecting a debt previously contracted (DPC), acquired voting securities of a bank or BHC or the securities or assets of a company engaged in a nonbanking activity to seek prior Board approval in order to retain ownership of those shares or assets for more than two years.

Legal authorization and confidentiality: The FR 4006 is authorized pursuant to sections 3(a) and 4(c)(2) of the BHC Act [1] and sections 225.12(b) and 225.22(d) of the Board's Regulation Y, which permit a BHC to acquire securities or assets in the ordinary course of collecting a DPC in good faith without seeking prior Board approval if such securities or assets (DPC property) are divested within two years of acquisition. To hold the DPC property beyond this two-year period, a BHC must seek the Board's approval.[2] The FR 4006 is required to obtain this benefit.

The information contained on the FR 4006 is not considered confidential unless an applicant requests confidential treatment in accordance with the Board's Rules Regarding Availability of Information.[3] Requests for confidential treatment of information are reviewed on a case-by-case basis. To the extent information provided on the FR 4006 is nonpublic commercial or financial information, which is both customarily and actually treated as private by the respondent, such information may be protected from disclosure pursuant to exemption 4 of the Freedom of Information Act.[4]

Current actions: On April 6, 2022, the Board published a notice in the **Federal Register** (87 FR 19926) requesting public comment for 60 days on the extension, with revision, of the FR 4006. The Board has revised the FR 4006 to account for the voluntary reporting provisions set forth in sections 225.140(c) and 225.140(d) of Regulation Y. These sections state, respectively, that a BHC that holds nonbanking DPC assets past the two-year statutory holding period should report annually to the appropriate Reserve Bank on its efforts to accomplish divestiture of such assets; and that a BHC that holds real estate acquired as DPC property for longer than five years should keep the appropriate Reserve Bank advised on a regular basis concerning its efforts to dispose of the property. The comment period for this notice expired on June 6, 2022. The Board did not receive any comments. The revisions will be implemented as proposed.

Board of Governors of the Federal Reserve System, July 27, 2022.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

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[1] 12 U.S.C. 1842(a) and 1843(c)(2).

[2] The two-year period may be extended by the Board for up to three years in one-year increments (12 CFR 225.12(b); 12 CFR 225.22(d)(1)). The Board may provide up to five additional one-year extensions (for a total of ten years) if the DPC property is shares, real estate, or other assets where the holding company demonstrates that each extension would not be detrimental to the public interest and either the bank holding company has made good faith attempts to dispose of such shares, real estate or other assets or disposal of the shares, real estate or other assets during the initial period would have been detrimental to the company (12 CFR 225.22(d)(1)(ii)).

[3] 12 CFR 261.17.

[4] 5 U.S.C. 552(b)(4).

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–0879]

Agency Information Collection Activities; Proposed Collection; Comment Request; Procedures for the Safe Processing and Importing of Fish and Fishery Products

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA, Agency, or we) 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 associated with safe and sanitary processing and importing of fish and fishery products.

DATES: Either electronic or written comments on the collection of information must be submitted by October 3, 2022.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of October 3, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received 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–0879 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Procedures for the Safe Processing and Importing of Fish and Fishery Products.'' 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, 240–402–7500.

- 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.govinfo.gov/content/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, 240–402–7500.

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.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3521), 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 Agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Procedures for the Safe Processing and Importing of Fish and Fishery Products—21 CFR Part 123

OMB Control Number 0910–0354—Extension

This information collection supports regulations in part 123 (21 CFR part 123), which mandate the application of hazard analysis and critical control point (HACCP) principles to the processing of seafood. HACCP is a preventive system of hazard control designed to help ensure the safety of foods. The regulations were issued under FDA's statutory authority to regulate food safety, including section 402(a)(1) and (4) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 342(a)(1) and (4)). Certain provisions in

part 123 require that processors and importers of seafood collect and record information.

The HACCP records compiled and maintained by a seafood processor primarily consist of the periodic observations recorded at selected monitoring points during processing and packaging operations, as called for in a processor's HACCP plan (*e.g.,* the values for processing times, temperatures, acidity, etc., as observed at critical control points). The primary purpose of HACCP records is to permit a processor to verify that products have been produced within carefully established processing parameters (critical limits) that ensure that hazards have been avoided.

HACCP records are normally reviewed by appropriately trained employees at the end of a production lot or at the end of a day or week of production to verify that control limits have been maintained, or that appropriate corrective actions were taken if the critical limits were not maintained. Such verification activities are essential to ensure that the HACCP system is working as planned. A review of these records during the conduct of periodic plant inspections also permits FDA to determine whether the products have been consistently processed in conformance with appropriate HACCP food safety controls.

Section 123.12 requires that importers of seafood products take affirmative steps and maintain records that verify that the fish and fishery products they offer for import into the United States were processed in accordance with the HACCP and sanitation provisions set forth in part 123. These records are also to be made available for review by FDA as provided in § 123.12(c).

The time and costs of these recordkeeping activities will vary considerably among processors and importers of fish and fishery products, depending on the type and number of products involved, and on the nature of the equipment or instruments required to monitor critical control points. The burden estimate in table 1 includes only those collections of information under the seafood HACCP regulations that are not already required under other statutes and regulations. The estimate also does not include collections of information that are a usual and customary part of businesses' normal activities. For example, the tagging and labeling of molluscan shellfish (21 CFR 1240.60) is a customary and usual practice among seafood part of processors. Consequently, the estimates in table 1 account only for information collection and recording requirements attributable to part 123.

Description of Respondents: Respondents to this collection of information include processors and importers of seafood.

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

TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]

21 CFR section; [2] activity	Number of recordkeepers	Number of records per recordkeeper [3]	Total annual records	Average burden per recordkeeping [4]	Total hours
123.6(a), (b), and (c); Prepare hazard analysis and HACCP plan	50	1	50	16	800
123.6(c)(5); Undertake and prepare records of corrective actions	15,000	4	60,000	0.30 (18 minutes)	18,000
123.8(a)(1) and (c); Reassess hazard analysis and HACCP plan	15,000	1	15,000	4	60,000
123.12(a)(2)(ii); Verify compliance of imports and prepare records of verification activities.	4,100	80	328,000	0.20 (12 minutes)	65,600
123.6(c)(7); Document monitoring of critical control points	15,000	280	4,200,000	0.30 (18 minutes)	1,260,000
123.7(d); Undertake and prepare records of corrective actions due to a deviation from a critical limit.	6,000	4	24,000	0.10 (6 minutes)	2,400
123.8(d); Maintain records of the calibration of process-monitoring instruments and the performing of any periodic end-product and in-process testing.	15,000	47	705,000	0.10 (6 minutes)	70,500
123.11(c); Maintain sanitation control records	15,000	280	4,200,000	0.10 (6 minutes)	420,000
123.12(c); Maintain records that verify that the fish and fishery products they offer for import into the United States were processed in accordance with the HACCP and sanitation provisions set forth in part 123.	4,100	80	328,000	0.10 (6 minutes)	32,800
123.12(a)(2); Prepare new written verification procedures to verify compliance of imports.	41	1	41	4	164
Total					1,930,264

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] These estimates include the information collection requirements in the following sections:
§ 123.16—Smoked Fish—process controls (see § 123.6(b));
§ 123.28(a)—Source Controls—molluscan shellfish (see § 123.6(b));
§ 123.28(c) and (d)—Records—molluscan shellfish (see § 123.6(c)(7).
[3] Based on an estimated 280 working days per year.
[4] Estimated average time per 8-hour workday unless one-time response.

Based on a review of the information collection since our last OMB approval, we have made no adjustments to our burden estimate. We base this hour burden estimate on our experience with the application of HACCP principles in food processing. Further, the burdens have been estimated using typical small seafood processing firms as a model because these firms represent a significant proportion of the industry. The hour burden of HACCP recordkeeping activities will vary considerably among processors and importers of fish and fishery products, depending on the size of the facility and complexity of the HACCP control scheme (*i.e.,* the number of products and the number of hazards controlled); the daily frequency that control points are monitored and values recorded; and also on the extent that data recording time and cost are minimized by the use of automated data logging technology. The burden estimate does not include burden hours for activities that are a usual and customary part of businesses' normal activities. For example, the tagging and labeling of molluscan shellfish (§ 1240.60) is a customary and usual practice among seafood processors.

Dated: July 20, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

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