the notice indicates that EPA anticipated allowing registrants to sell and distribute existing stocks of chlorpyrifos for one year after the publication of the cancellation order and to allow others to sell, distribute, and use existing stocks until exhausted. Earthjustice asserts that EPA cannot allow for existing stocks of chlorpyrifos to be sold, distributed, or used in this way as a result of the revocation of all chlorpyrifos tolerances. See the August 30, 2021 (86 FR 48315) (FRL–5993–04–OCSPP) publication in the **Federal Register**.

The applicability of the existing stocks language in the April 2022 notice of receipt to chlorpyrifos products was an oversight and is being corrected in this cancellation order. The existing stocks section of the April 2022 notice of receipt contained specific language pertaining to one specific product (not the chlorpyrifos products) and other language ''for all other voluntary cancellations listed in the notice.'' 87 FR 25259. That language provided broad existing stocks provisions due to the Agency's conclusion that there were ''no significant potential risk concerns associated with those pesticide products.'' *Id.* In using somewhat standard language for a voluntary cancellation notice, EPA failed to specify a different existing stocks provision for the chlorpyrifos products in the notice, for which tolerances were revoked due to EPA's conclusion that chlorpyrifos tolerances were not safe. See 87 FR 11222. FIFRA section 6(a)(1) gives EPA the discretion to permit the continued sale and use of existing stocks, where doing so is determined to be consistent with the purposes of FIFRA. 7 U.S.C. 136d(a)(1). In the case of the chlorpyrifos registrations subject to this order, without tolerances in place to cover residues from use of these products, these products may not be used on food, nor may they be sold or distributed. Allowing for continued use or sale would not be consistent with FIFRA; therefore, EPA is not allowing for continued sale, distribution, or use of chlorpyrifos products listed above.

III. The Cancellation Order

Pursuant to FIFRA section 6(f) (7 U.S.C. 136d(f)(1)), EPA hereby approves the requested cancellations of the registrations identified in Table 1 of Unit I. Accordingly, the Agency hereby orders that the product registrations identified in Table 1 of Unit I. are cancelled.

The cancellations and amendments addressed in this Order are effective August 31, 2022. Any distribution, sale, or use of existing stocks of the products identified in Table 1 of Unit I. in a manner inconsistent with any of the provisions for disposition of existing stocks set forth in Unit VI. will be a violation of FIFRA.

IV. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products that are currently in the United States and that were packaged, labeled, and released for shipment prior to the effective date of the cancellation action. The language regarding the intended disposition of existing stocks that published in the **Federal Register** on April 28, 2022 (87 FR 25256) (FRL–9723–01–OCSPP) on page 25259 is not appropriate for application to the pesticide products subject to this Order.

None of the registrants listed in this order have requested any continued sale or distribution of existing stocks of the registrations subject to this cancellation order nor have they requested special provisions to relabel the products listed in this order. Because of that and because chlorpyrifos tolerances have been revoked and use of chlorpyrifos renders food adulterated, all sale, distribution, and use of the chlorpyrifos products identified in Table 1 of Unit I. is prohibited, except for export consistent with FIFRA section 17, 7 U.S.C. 136o or for proper disposal.

Dated: August 26, 2022.

Mary Elissa Reaves,

Director, Pesticide Re-Evaluation Division, Office of Pesticide Programs.

[FR Doc. 2022–18838 Filed 8–30–22; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)).

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than September 30, 2022.

A. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. *Sword Financial Corporation, Horicon, Wisconsin;* to acquire Community Bancshares Wisconsin and thereby indirectly acquire Cornerstone Community Bank, both of Grafton, Wisconsin.

B. Federal Reserve Bank of Kansas City (Jeffrey Imgarten, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198–0001:

1. *First National Buffalo Bankshares, Inc., Buffalo, Wyoming;* to acquire First State Bank of Newcastle, Newcastle, Wyoming.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2022–18808 Filed 8–30–22; 8:45 am]

BILLING CODE P

FEDERAL RETIREMENT THRIFT INVESTMENT BOARD

Senior Executive Service Performance Review Board

AGENCY: Federal Retirement Thrift Investment Board.

ACTION: Notice.

SUMMARY: This notice announces the appointment of the members of the Senior Executive Service Performance Review Board for the Federal Retirement Thrift Investment Board. The purpose of the Performance Review Board is to make written recommendations on each executive's annual summary ratings, performance-based pay adjustment, and performance awards to the appointing authority.

DATES: This notice is applicable on August 31, 2022.

FOR FURTHER INFORMATION CONTACT: Kelly Powell, HR Specialist, at 202–942–1681.

SUPPLEMENTARY INFORMATION: Title 5, U.S. Code, 4314(c)(4), requires that the appointment of Performance Review Board members be published in the **Federal Register** before Board service commences. The following persons will serve on the Federal Retirement Thrift Investment Board's Performance Review Board which will review initial summary ratings to ensure the ratings are consistent with established performance requirements, reflect meaningful distinctions among senior executives based on their relative performance and organizational results and provide recommendations for ratings, awards, and pay adjustments in a fair and equitable manner: Thomas Brandt, Jim Courtney, Sean McCaffrey, and Kim Weaver.

Dharmesh Vashee,

General Counsel, Federal Retirement Thrift Investment Board.

[FR Doc. 2022–18784 Filed 8–30–22; 8:45 am]

BILLING CODE 6760–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket Nos. FDA–2020–E–2259; FDA–2020–E–2260; and FDA–2020–E–2261]

Determination of Regulatory Review Period for Purposes of Patent Extension; THEROX DOWNSTREAM SYSTEM

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or the Agency) has determined the regulatory review period for THEROX DOWNSTREAM SYSTEM and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of applications to the Director of the U.S. Patent and Trademark Office (USPTO), Department of Commerce, for the extension of a patent which claims that medical device.

DATES: Anyone with knowledge that any of the dates as published (see **SUPPLEMENTARY INFORMATION**) are incorrect must submit either electronic or written comments and ask for a redetermination by October 31, 2022. Furthermore, any interested person may petition FDA for a determination regarding whether the applicant for extension acted with due diligence during the regulatory review period by February 27, 2023. See ''Petitions'' in the **SUPPLEMENTARY INFORMATION** section for more information.

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 31, 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 Nos. FDA–2020–E–2259; FDA–2020–E–2260; and FDA–2020–E–2261 for ''Determination of Regulatory Review Period for Purposes of Patent Extension; THEROX DOWNSTREAM SYSTEM.'' 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 § 10.20 (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 numbers, 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: Beverly Friedman, Office of Regulatory Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 51, Rm. 6250, Silver Spring, MD 20993, 301–796–3600.

SUPPLEMENTARY INFORMATION: