- Risk assessments.
- Bibliographies concerning current registrations.
- Summaries of incident data.
- Any other pertinent data or information.

Each docket contains a document summarizing what the agency currently knows about the pesticide case and a preliminary work plan for anticipated data and assessment needs. Additional documents provide more detailed information. During this public comment period, the agency is asking that interested persons identify any additional information they believe the agency should consider during the registration reviews of these pesticides. The agency identifies in each docket the areas where public comment is specifically requested, though comment in any area is welcome.

The registration review final rule at 40 CFR 155.50(b) provides for a minimum 60-day public comment period on all preliminary registration review work plans. This comment period is intended to provide an opportunity for public input and a mechanism for initiating any necessary changes to a pesticide's workplan. All comments should be submitted using the methods in **ADDRESSES** and must be received by the EPA on or before the closing date. These comments will become part of the docket for the pesticides included in Table 1 in Unit IV. Comments received after the close of the comment period will be marked ''late.'' The EPA is not required to consider these late comments.

The agency will carefully consider all comments received by the closing date and may provide a ''Response to Comments Memorandum'' in the docket. The final registration review work plan will explain the effect that any comments had on the final work plan and provide the agency's response to significant comments.

Background on the registration review program is provided at: *https://www.epa.gov/pesticide-reevaluation.*

Authority: 7 U.S.C. 136 *et seq.*

Dated: October 19, 2022.

Mary Elissa Reaves,

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

[FR Doc. 2022–23186 Filed 10–24–22; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2017–0751; FRL–10274–01–OCSPP]

Pesticide Registration Review; Decisions and Case Closures for Several Pesticides; Notice of Availability

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's interim or final registration review decisions for the following chemicals: *pasteuria* species, *pseudomonas syringae;* and trimethylamine and trimethylamine hydrochloride. In addition, this notice announces the closure of the registration review case for cryolite, furfural, mefluidide, and sabadilla alkaloids because the last U.S. registrations for these pesticides have been canceled.

ADDRESSES: The docket for this action, identified under docket identification (ID) number EPA–HQ–OPP–2017–0751, is available online at *https://www.regulations.gov.* Additional instructions on visiting the docket, along with more information about dockets generally, is available at *https://www.epa.gov/dockets.*

FOR FURTHER INFORMATION CONTACT:

For pesticide specific information, contact: The Chemical Review Manager for the pesticide of interest identified in Table 1 in Unit IV.

For general information on the registration review program, contact: Melanie Biscoe, Pesticide Re-evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460–0001; telephone number: 202–566–0701; email address: *biscoe.melanie@epa.gov.*

SUPPLEMENTARY INFORMATION:

I. General Information

Does this action apply to me?

This action is directed to the public in general and may be of interest to a wide range of stakeholders including environmental, human health, farm worker, and agricultural advocates; the chemical industry; pesticide users; and members of the public interested in the sale, distribution, or use of pesticides. Since others also may be interested, the Agency has not attempted to describe all the specific entities that may be affected by this action. If you have any questions regarding the applicability of this action to a particular entity, consult the pesticide specific contact person listed under **FOR FURTHER INFORMATION CONTACT**: *For pesticide specific information, contact:* The Chemical Review Manager for the pesticide of interest identified in Table 1 in Unit IV.

II. Background

Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed interim or final decisions for all pesticides listed in Table 1 in Unit IV. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment.

III. Authority

EPA is conducting its registration review of the chemicals listed in Table 1 in Unit IV pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review at 40 CFR part 155, subpart C. Section 3(g) of FIFRA provides, among other things, that the registrations of pesticides are to be reviewed every 15 years. Under FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5) (7 U.S.C. 136a(c)(5)). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

IV. What action is the Agency taking?

Pursuant to 40 CFR 155.58, this notice announces the availability of EPA's interim or final registration review decisions for the pesticides shown in Table 1. The registration review decisions are supported by rationales included in the docket established for each chemical.

TABLE 1—REGISTRATION REVIEW INTERIM AND FINAL DECISIONS BEING ISSUED

Registration review case name and No.	Docket ID No.	Chemical review manager and contact information
Pasteuria; species Case Numbers: 6526, 6527, 6535	EPA–HQ–OPP–2021–0614	Andrew Queen *queen.andrew@epa.gov,* (202) 566–1539.
Pseudomonas syringae; Case Number 6007	EPA–HQ–OPP–2022–0088	Bibiana Oe, *oe.bibiana@epa.gov,* (202) 566–1538.
Trimethylamine and Trimethylamine Hydrochloride; Case No. 6304.	EPA–HQ–OPP–2021–0852	Monica Thapa, *thapa.monica@epa.gov,* (202) 566–1543.

The proposed decisions and proposed interim registration review decisions for the chemicals in the table above were posted to the docket and the public was invited to submit any comments or new information. EPA addressed the comments or information received during the 60-day comment period for the proposed interim decisions in the discussion for each pesticide listed in the table. Comments from the 60-day comment period that were received may or may not have affected the Agency's interim or final decision. Pursuant to 40 CFR 155.58(c), the registration review case docket for the chemicals listed in the Table will remain open until all actions required in the decision have been completed.

This document also announces the closure of the registration review case for cryolite (Case Number 0087, Docket ID Number EPA–HQ–OPP–2011–0173), furfural (Case Number 7050, Docket ID Number EPA–HQ–OPP–2014–0764), mefluidide (Case Number 2370, Docket ID Number EPA–HQ–2015–0786), and sabadilla alkaloids (Case Number 3128, Docket ID Number EPA–HQ–OPP–2015–0063) because the last U.S. registrations for these pesticides have been canceled.

Background on the registration review program is provided at: *https://www.epa.gov/pesticide-reevaluation.*

Authority: 7 U.S.C. 136 *et seq.*

Dated: October 19, 2022.

Mary Elissa Reaves,

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

[FR Doc. 2022–23210 Filed 10–24–22; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL MEDIATION AND CONCILIATION SERVICE

Senior Executive Service Performance Review Board

AGENCY: Federal Mediation and Conciliation Service (FMCS).

ACTION: Notice of Senior Executive Service Performance Review Board.

SUMMARY: The Federal Mediation and Conciliation Service (FMCS) is issuing this notice to inform the public of the names of the members of the Agency's Senior Executive Service (SES) Performance Review Board.

DATES: This SES Performance Review Board is effective October 25, 2022.

FOR FURTHER INFORMATION CONTACT: Alisa Zimmerman, Acting General Counsel, 202–606–5488, *ogc@fmcs.gov,* 250 E St. SW, Washington, DC 20427.

SUPPLEMENTARY INFORMATION: Sec. 4314(c)(1) through (5) of title 5, U.S.C., requires each agency to establish, in accordance with regulations prescribed by the Office of Personnel Management, one or more performance review boards. The board shall review and evaluate the initial appraisal of a senior executive's performance by the supervisor, along with any recommendations to the appointing authority relative to the performance of the senior executive.

The members of FMCS's Performance Review Board are:

1. Marla Hendriksson, Deputy Director for the Office of Partnership and Operational Policy, Office of Regulatory Affairs, Food and Drug Administration, Department of Health and Human Services
2. Javier Ramirez, Deputy Director Field Operations, Federal Mediation and Conciliation Service
3. Angie Titcombe, Director of Human Resources, Federal Mediation and Conciliation Services
4. Josh Flax, Deputy Director for Policy and Strategy, Federal Mediation and Conciliation Service

Dated: October 20, 2022.

Alisa Zimmerman,

Acting General Counsel.

[FR Doc. 2022–23209 Filed 10–24–22; 8:45 am]

BILLING CODE 6732–01–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Savings and Loan Holding Company

The notificants listed below have applied under the Change in Bank Control Act (''Act'') (12 U.S.C. 1817(j)) and of the Board's Regulation LL (12 CFR 238.31) to acquire shares of a savings and loan holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

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 paragraph 7 of the Act.

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 November 8, 2022.

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

1. *The Vanguard Group, Inc., Malvern, Pennsylvania; on behalf of itself, its subsidiaries and affiliates, including investment companies registered under the Investment Company Act of 1940, other pooled investment vehicles, and institutional accounts that are sponsored, managed, or advised by Vanguard;* to acquire additional voting shares of Capitol Federal Financial, Inc., and thereby indirectly acquire additional voting shares of Capitol Federal Savings Bank, both of Topeka, Kansas.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2022–23212 Filed 10–24–22; 8:45 am]

BILLING CODE P