

TSCA, the Pollution Prevention Act (PPA), 42 U.S.C. 13101 *et seq.*, and other applicable statutes. The TSCA SACC provides expert independent scientific advice and recommendations to the EPA on the scientific and technical aspects of risk assessments, methodologies, and pollution prevention measures and approaches for chemicals regulated under TSCA.

The TSCA SACC is comprised of experts in: Toxicology; human health and environmental risk assessment; exposure assessment; and related sciences (e.g., synthetic biology, pharmacology, biotechnology, nanotechnology, biochemistry, biostatistics, PBPK modeling, computational toxicology, epidemiology, environmental fate, and environmental engineering and sustainability). The TSCA SACC currently consists of 24 members. When needed, the committee will be assisted in their reviews by ad hoc participants with specific expertise in the topics under consideration.

#### D. TSCA SACC Documents and Meeting Minutes

EPA's background paper, related supporting materials, and draft charge questions to TSCA SACC are available on the TSCA SACC website and in the docket established for the specific chemical. In addition, the EPA will provide additional background documents (e.g., TSCA SACC members participating in this meeting and the meeting agenda) as the materials become available. You may obtain electronic copies of these documents, and certain other related documents that might be available, at <http://www.regulations.gov> and the TSCA SACC website at <https://www.epa.gov/tsca-peer-review>.

TSCA SACC will prepare meeting minutes summarizing its recommendations to the EPA. The meeting minutes will be posted on the TSCA SACC website and in the relevant docket.

**Authority:** 15 U.S.C. 2601 *et seq.*; 5 U.S.C. Appendix 2 *et seq.*

Dated: August 5, 2019.

**Andrew R. Wheeler,**  
Administrator.

[FR Doc. 2019-17222 Filed 8-9-19; 8:45 am]

**BILLING CODE 6560-50-P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2019-0448; FRL-9997-71]

### Nominations to the Federal Insecticide, Fungicide, and Rodenticide Act Scientific Advisory Panel; Request for Comments

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** This notice provides the names, addresses, and professional affiliations of persons recently nominated to serve on the Scientific Advisory Panel (SAP) established under section 25(d) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA). The Panel was created on November 28, 1975, and made a permanent Panel by amendment to FIFRA, dated October 25, 1988. The Agency, at this time, anticipates selecting three new members to serve on the panel because of membership terms that will expire during the next year. Public comments on the current nominations are invited, as these comments will be used to assist the Agency in selecting the new members for the chartered Scientific Advisory Panel.

**DATES:** Comments identified by docket ID number EPA-HQ-OPP-2019-0448, must be received on or before September 11, 2019.

**ADDRESSES:** Submit your comments, identified by docket identification (ID) number EPA-HQ-OPP-2019-0448, by one of the following methods:

- **Federal eRulemaking Portal:** <https://www.regulations.gov>. Follow the online instructions for submitting comments. Do not electronically submit any information you consider to be Confidential Business Information (CBI) or information whose disclosure is restricted by statute.

- **Mail:** OPP Docket, Environmental Protection Agency Docket Center (EPA/DC), (28221T), 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001.

- **Hand Delivery:** To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at <https://www.epa.gov/dockets/contacts.html>.

**FOR FURTHER INFORMATION CONTACT:** Steven Knott, M.S., Designated Federal Officer (DFO), Office of Science Coordination and Policy (7201M), Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001; telephone number: (202) 564-0103; email address: [knott.steven@epa.gov](mailto:knott.steven@epa.gov).

## SUPPLEMENTARY INFORMATION:

### I. General Information

#### A. Does this action apply to me?

This action is directed to the public in general. This action may, however, be of interest to persons who are or may be required to conduct testing of chemical substances under the Federal Food, Drug, and Cosmetic Act (FFDCA) and FIFRA. Given other entities may also 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 DFO listed under **FOR FURTHER INFORMATION CONTACT**.

#### B. What should I consider as I prepare my comments for EPA?

1. **Submitting CBI.** Do not submit CBI information to EPA through [regulations.gov](https://www.regulations.gov) or email. If your comments contain any information that you consider to be CBI or otherwise protected, please contact the DFO listed under **FOR FURTHER INFORMATION CONTACT** to obtain special instructions before submitting your comments.

2. **Tips for preparing your comments.** When preparing and submitting your comments, see the commenting tips at <https://www.epa.gov/dockets/comments.html>.

### II. Background

The FIFRA SAP serves as a scientific peer review mechanism of EPA's Office of Chemical Safety and Pollution Prevention (OCSP) and is structured to provide independent scientific advice, information, and recommendations to the EPA Administrator on pesticides and pesticide-related issues as to the impact of regulatory actions on health and the environment. The FIFRA SAP is a federal advisory committee established in 1975 under FIFRA that operates in accordance with requirements of the Federal Advisory Committee Act (5 U.S.C. Appendix). The FIFRA SAP is composed of a permanent panel consisting of seven members who are appointed by the EPA Administrator from nominees provided by the National Institutes of Health (NIH) and the National Science Foundation (NSF). FIFRA established a Science Review Board (SRB) consisting of at least 60 scientists who are available to the FIFRA SAP on an ad hoc basis to assist in reviews conducted by the FIFRA SAP. As a scientific peer review mechanism, the FIFRA SAP provides comments, evaluations, and recommendations to improve the

effectiveness and quality of analyses made by Agency scientists. Members of the FIFRA SAP are scientists who have sufficient professional qualifications, including training and experience, to provide expert advice and recommendations to the Agency.

The Agency, at this time, anticipates selecting three new members to serve on the panel because of membership terms that will expire during the next year. The Agency requested that NIH and NSF nominate experts for selection from the fields of ecological and human health risk assessment with specific expertise in mathematical biostatistics, ecotoxicology and environmental fate and transport of chemicals, and neurotoxicity (including developmental neurotoxicity). The Agency also noted that experts with specific experience in cheminformatics, bioinformatics, and genomics are preferred. Nominees should be well published and current in their fields of expertise.

### III. Charter

A Charter for the FIFRA SAP, dated October 17, 2018, was issued in accordance with the requirements of the Federal Advisory Committee Act, Public Law 92-463, 86 Stat. 770 (5 U.S.C. Appendix). The Charter provides for open meetings with opportunities for public participation.

### IV. Nominees

#### A. Qualifications of Members

Members are scientists who have sufficient professional qualifications, including training and experience, to provide expert comments on the impact of pesticides on human health and the environment. No persons shall be ineligible to serve on the FIFRA SAP by reason of their membership on any other advisory committee to a federal department or agency, or their employment by a federal department or agency (except the EPA). The Administrator appoints individuals to serve on the Panel for staggered terms of up to 3 years. Panel members are subject to the provisions of the Standards of Ethical Conduct for Employees of the Executive Branch at 5 Code of Federal Regulations Part 2635, conflict of interest statutes in Title 18 of the United States Code, and related regulations. Each nominee selected by the Administrator, before being formally appointed, is required to submit a confidential statement of employment and financial interest, which shall fully disclose, among other financial interests, the nominee's sources of research support, if any.

In accordance with section 25(d)(1) of FIFRA, the Administrator shall require nominees to the FIFRA SAP to furnish information concerning their professional qualifications, including educational background, employment history, and scientific publications. FIFRA further stipulates that the Agency publish the name, address, and professional affiliation of the nominees in the **Federal Register**.

#### B. Applicability of Existing Regulations

With respect to the requirements of section 25(d) of FIFRA that the Administrator promulgate regulations regarding conflicts of interest, FIFRA SAP members are subject to the provisions of the Standards of Ethical Conduct for Employees of the Executive Branch at 5 Code of Federal Regulations Part 2635, conflict of interest statutes in Title 18 of the United States Code, and related regulations.

#### C. Process of Obtaining Nominees

In accordance with the provisions of section 25(d) of FIFRA, EPA, on February 25, 2019, requested that the NIH and the NSF nominate scientists to fill vacancies occurring on the FIFRA SAP. The Agency requested nominations of experts in the fields of ecological and human health risk assessment with specific expertise in mathematical biostatistics, ecotoxicology and environmental fate and transport of chemicals, and neurotoxicity (including developmental neurotoxicity). The Agency also noted that experts with specific experience in cheminformatics, bioinformatics, and genomics are preferred. NIH and NSF responded, providing the Agency with a total of 35 nominees. Two nominees, Dr. Dana Barr of Emory University and Dr. Marion Ehrich of Virginia Tech, are recent members of the FIFRA SAP and, therefore, were not considered further for membership at this time. Of the remaining 33 nominees, 15 are interested and available to actively participate in FIFRA SAP meetings (see Section IV.D.). The following 18 individuals are not available:

1. *David Allison, Ph.D.*, Indiana University, Bloomington, Indiana.
2. *Asa Bradman, Ph.D.*, University of California, Berkeley, California.
3. *Alex Buerkle, Ph.D.*, University of Wyoming, Laramie, Wyoming.
4. *Atul Butte, M.D., Ph.D.*, University of California, San Francisco, California.
5. *Lucio Costa, Ph.D.*, University of Washington, Seattle, Washington.
6. *Rebecca Doerge, Ph.D.*, Carnegie Mellon University, Pittsburgh, Pennsylvania.

7. *Elaine Faustman, Ph.D.*, University of Washington, Seattle, Washington.

8. *Jodi Flaws, Ph.D.*, University of Illinois, Urbana-Champaign, Illinois.

9. *Brandon Gaut, Ph.D.*, University of California, Irvine, California.

10. *Phillipe Grandjean, M.D.*, Harvard University, Boston, Massachusetts.

11. *Linda Lee, Ph.D.*, Purdue University, West Lafayette, Indiana.

12. *Mary Kay O'Rourke, Ph.D.*, University of Arizona, Tucson, Arizona.

13. *Virginia Rauh, Ph.D.*, Columbia University, New York, New York.

14. *Rick Relyea, Ph.D.*, Rensselaer Polytechnic Institute, Troy, New York.

15. *Diane Rohlman, Ph.D.*, University of Iowa, Iowa City, Iowa.

16. *Jason Rohr, Ph.D.*, University of Notre Dame, Notre Dame, Indiana.

17. *Caroline Tanner, Ph.D.*, University of California, San Francisco, California.

18. *Cari Vanderpool, Ph.D.*, University of Illinois, Urbana-Champaign, Illinois.

#### D. Interested and Available Nominees

Following are the names, addresses, and professional affiliations of current nominees being considered for membership on the FIFRA SAP. Selected biographical data for each nominee is available in the public docket at [www.regulations.gov](http://www.regulations.gov) (ID number EPA-HQ-OPP-2019-0448) and through the FIFRA SAP website at <https://www.epa.gov/sap>. The Agency anticipates selecting three individuals to fill vacancies occurring during the next year.

1. *Jeffrey Bloomquist, Ph.D.*: Professor, Entomology and Nematology Department, Emerging Pathogens Institute, University of Florida, Gainesville, Florida.

2. *Maria Braga, D.D.S., Ph.D.*: Professor, Department of Anatomy, Physiology and Genetics, Uniformed Services University of the Health Sciences, Bethesda, Maryland.

3. *Joseph Braun, R.N., M.S.P.H., Ph.D.*: Associate Professor, Department of Epidemiology, Brown University, Providence, Rhode Island.

4. *Celia Chen, Ph.D.*: Director, Dartmouth Toxic Metals Superfund Research Program and Research Professor, Department of Biological Sciences, Dartmouth College, Hanover, New Hampshire.

5. *Susan Fisher, Ph.D.*: Professor Emerita, Department of Entomology, Ohio State University, Columbus, Ohio.

6. *Jean Harry, M.S., Ph.D.*: Group Leader, Neurotoxicology Laboratory, National Toxicology Program, National Institute of Environmental Health Sciences, Research Triangle Park, North Carolina.

7. *Tyrone Hayes, Ph.D.*: Professor, Department of Integrative Biology, University of California, Berkeley, California.

8. *Lucille Lange, Ph.D.*: Research Psychologist, US Army Medical Research Institute of Chemical Defense, Aberdeen Proving Ground, Maryland.

9. *James Lauderdale, Ph.D.*: Associate Professor, Department of Cellular Biology, University of Georgia, Athens, Georgia.

10. *Maureen Lichtveld, M.D., M.P.H.*: Professor and Chair, Department of Global Environmental Health Sciences, School of Public Health and Tropical Medicine, Tulane University, New Orleans, Louisiana.

11. *Lorenz Neuwirth, Ph.D.*: Assistant Professor, Department of Psychology and the Neuroscience Research Institute, State University of New York (SUNY), Old Westbury, New York.

12. *Edna Pereira, Ph.D.*: Associate Professor, Department of Epidemiology and Public Health and Director, Division of Translational Toxicology, University of Maryland School of Medicine, Baltimore, Maryland.

13. *Rebecca Smith, D.V.M., M.S., Ph.D.*: Assistant Professor, Department of Pathobiology, College of Veterinary Medicine, University of Illinois, Urbana-Champaign, Illinois.

14. *John Swaddle, Ph.D.*: Professor and Chair, Biology Department, College of William & Mary, Williamsburg, Virginia.

15. *Christopher Weis, Ph.D.*: Toxicology Liaison, National Toxicology Program, National Institute of Environmental Health Sciences, Bethesda, Maryland.

**Authority:** 7 U.S.C. 136 *et seq.*; 21 U.S.C. 301 *et seq.*

Dated: August 2, 2019.

**Hayley Hughes,**

*Director, Office of Science Coordination and Policy.*

[FR Doc. 2019-17150 Filed 8-9-19; 8:45 am]

**BILLING CODE 6560-50-P**

## ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPPT-2019-0075; FRL-9992-80]

### Certain New Chemicals; Receipt and Status Information for June 2019

**AGENCY:** Environmental Protection Agency (EPA).

**ACTION:** Notice.

**SUMMARY:** EPA is required under the Toxic Substances Control Act (TSCA), as amended by the Frank R. Lautenberg Chemical Safety for the 21st Century

Act, to make information publicly available and to publish information in the **Federal Register** pertaining to submissions under TSCA Section 5, including notice of receipt of a Premanufacture notice (PMN), Significant New Use Notice (SNUN) or Microbial Commercial Activity Notice (MCAN), including an amended notice or test information; an exemption application (Biotech exemption); an application for a test marketing exemption (TME), both pending and/or concluded; a notice of commencement (NOC) of manufacture (including import) for new chemical substances; and a periodic status report on new chemical substances that are currently under EPA review or have recently concluded review. This document covers the period from 06/01/2019 to 06/30/2019.

**DATES:** Comments identified by the specific case number provided in this document must be received on or before September 11, 2019.

**ADDRESSES:** Submit your comments, identified by docket identification (ID) number EPA-HQ-OPPT-2019-0075, and the specific case number for the chemical substance related to your comment, by one of the following methods:

- *Federal eRulemaking Portal:* <http://www.regulations.gov>. Follow the online instructions for submitting comments. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute.

- *Mail:* Document Control Office (7407M), Office of Pollution Prevention and Toxics (OPPT), Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001.

- *Hand Delivery:* To make special arrangements for hand delivery or delivery of boxed information, please follow the instructions at <http://www.epa.gov/dockets/contacts.html>. Additional instructions on commenting or visiting the docket, along with more information about dockets generally, is available at <http://www.epa.gov/dockets>.

#### FOR FURTHER INFORMATION CONTACT:

*For technical information contact:* Jim Rahai, Information Management Division (MC 7407M), Office of Pollution Prevention and Toxics, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001; telephone number: (202) 564-8593; email address: [rahai.jim@epa.gov](mailto:rahai.jim@epa.gov).

*For general information contact:* The TSCA-Hotline, ABVI-Goodwill, 422

South Clinton Ave., Rochester, NY 14620; telephone number: (202) 554-1404; email address: [TSCA-Hotline@epa.gov](mailto:TSCA-Hotline@epa.gov).

#### SUPPLEMENTARY INFORMATION:

##### I. Executive Summary

###### A. What action is the Agency taking?

This document provides the receipt and status reports for the period from 06/01/2019 to 06/30/2019. The Agency is providing notice of receipt of PMNs, SNUNs and MCANs (including amended notices and test information); an exemption application under 40 CFR part 725 (Biotech exemption); TMEs, both pending and/or concluded; NOCs to manufacture a new chemical substance; and a periodic status report on new chemical substances that are currently under EPA review or have recently concluded review.

EPA is also providing information on its website about cases reviewed under the amended TSCA, including the section 5 PMN/SNUN/MCAN and exemption notices received, the date of receipt, the final EPA determination on the notice, and the effective date of EPA's determination for PMN/SNUN/MCAN notices on its website at: <https://www.epa.gov/reviewing-new-chemicals-under-toxic-substances-control-act-tsca/status-pre-manufacture-notices>. This information is updated on a weekly basis.

###### B. What is the Agency's authority for taking this action?

Under the Toxic Substances Control Act (TSCA), 15 U.S.C. 2601 *et seq.*, a chemical substance may be either an "existing" chemical substance or a "new" chemical substance. Any chemical substance that is not on EPA's TSCA Inventory of Chemical Substances (TSCA Inventory) is classified as a "new chemical substance," while a chemical substance that is listed on the TSCA Inventory is classified as an "existing chemical substance." (See TSCA section 3(11).) For more information about the TSCA Inventory go to: <https://www.epa.gov/tsca-inventory>.

Any person who intends to manufacture (including import) a new chemical substance for a non-exempt commercial purpose, or to manufacture or process a chemical substance in a non-exempt manner for a use that EPA has determined is a significant new use, is required by TSCA section 5 to provide EPA with a PMN, MCAN or SNUN, as appropriate, before initiating the activity. EPA will review the notice, make a risk determination on the chemical substance or significant new