unreasonable adverse effects on human health or the environment. As part of the registration review process, the Agency has completed comprehensive draft human health and/or ecological risk assessments for all pesticides listed in the Table in Unit IV. After reviewing comments received during the public comment period, EPA may issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments and may request public input on risk mitigation before completing a proposed registration review decision for the pesticides listed in the Table 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 the Table 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 human health and/or ecological risk assessments for the pesticides shown in the following table and opens a 60-day public comment period on the risk assessments.

<table>
<thead>
<tr>
<th>Registration review case and No.</th>
<th>Docket ID No.</th>
<th>Chemical review manager and contact information</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chlorothalonil; Case 0097</td>
<td>EPA–HQ–OPP–2011–0840</td>
<td>Jon Williams, <a href="mailto:williams.jonathanr@epa.gov">williams.jonathanr@epa.gov</a>, (703) 347–0670.</td>
</tr>
</tbody>
</table>

Pursuant to 40 CFR 155.53(c), EPA is providing an opportunity, through this notice of availability, for interested parties to provide comments and input concerning the Agency’s draft human health and/or ecological risk assessments for the pesticides listed in the Table in Unit IV. The Agency will consider all comments received during the public comment period and make changes, as appropriate, to a draft human health and/or ecological risk assessment. EPA may then issue a revised risk assessment, explain any changes to the draft risk assessment, and respond to comments.

**Information submission requirements.** Anyone may submit data or information in response to this document. To be considered during a pesticide’s registration review, the submitted data or information must meet the following requirements:

- To ensure that EPA will consider data or information submitted, interested persons must submit the data or information during the comment period. The Agency may, at its discretion, consider data or information submitted at a later date.

- The data or information submitted must be presented in a legible and useable form. For example, an English translation must accompany any material that is not in English and a written transcript must accompany any information submitted as an audio-graphic or video-graphic record. Written material may be submitted in paper or electronic form.

  - Submitters must clearly identify the source of any submitted data or information.

  - Submitters may request the Agency to reconsider data or information that the Agency rejected in a previous review. However, submitters must explain why they believe the Agency should reconsider the data or information in the pesticide’s registration review.

As provided in 40 CFR 155.58, the registration review docket for each pesticide case will remain publicly accessible through the duration of the registration review process; that is, until all actions required in the final decision on the registration review case have been completed.

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

**Dated:** April 23, 2021.

**Mary Reaves,**

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

[FR Doc. 2021–10715 Filed 5–20–21; 8:45 am]

**BILLING CODE P**

### ENVIRONMENTAL PROTECTION AGENCY


**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 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 new members to serve on the panel because of the upcoming expirations of membership terms. Current members of the SAP are eligible for reappointment during this period. Therefore, the appointments completed over the next year may include a mix of newly appointed and reappointed members. As additional background, the biographies of current SAP members are available on the FIFRA SAP website at: [https://www.epa.gov/sap](https://www.epa.gov/sap). 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 SAP.

DATES: Comments identified by docket ID number EPA–HQ–OPP–2021–0293, must be received on or before June 21, 2021.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2021–0293, using the Federal eRulemaking Portal at 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. Comments submitted to the EPA, including any personal information that is in the body of the submission, will be publicly posted to https://www.regulations.gov and are also made available for in-person viewing at the EPA Docket Center’s Reading Room. There are some exceptions. Please see additional instructions on commenting or visiting the docket, along with more information about dockets generally, available at http://www.epa.gov/dockets.

Please note that due to the public health concerns related to COVID–19, the EPA Docket Center (EPA/DC) and Public Reading Room are closed to visitors with limited exceptions. The EPA/DC staff continue to provide remote customer service via email, phone, and webform. For the latest status information on EPA/DC services and docket access, visit https://www.epa.gov/dockets.

FOR FURTHER INFORMATION CONTACT: Steven Knott, M.S., Designated Federal Officer (DFO), Office of Program Support, Environmental Protection Agency; telephone number: (202) 564–0103; email address: 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 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/commenting-epa-dockets#tips.

II. Background

The FIFRA SAP serves as a scientific peer review mechanism of EPA’s Office of Chemical Safety and Pollution Prevention (OCPPP) 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). Members serve staggered terms of appointment, generally of three to six years duration. 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.

III. Charter

A Charter for the FIFRA SAP, dated October 17, 2020, 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. 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. 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). 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. Each nominee selected by the Administrator, before being formally appointed, is required to submit a Confidential Financial Disclosure Form, which shall fully disclose, among other financial interests, the nominee’s sources of research support, if any.

C. Process of Obtaining Nominees

The Agency, at this time, anticipates selecting new members to serve on the panel to address upcoming expirations of membership terms.

On February 5, 2021, in accordance with the provisions of section 25(d) of FIFRA, EPA requested that the NIH and NSF nominate scientists to fill vacancies occurring on the FIFRA SAP. The Agency requested that NIH and NSF nominate experts in the fields of ecological and human exposure assessment, including expertise in environmental fate and transport of chemicals, mathematical biostatistics, New Approach Methodologies (NAMs), including in vitro to in vivo extrapolation (IVIVE), and expertise in toxicology, including carcinogenicity and physiologically-based pharmacokinetic modeling (PBPK).
Experts with specific experience in risk assessment, dose response analysis, cheminformatics, bioinformatics, and genomics are preferred. Nominees should be well published and current in their fields of expertise.

NIH and NSF responded, providing the Agency with a total of 25 nominees. One nominee, Dr. Jeffrey Bloomquist, Ph.D., of the University of Florida is a member of the FIFRA SAP with a current term ending in 2022. Therefore, he is not considered further at this time. Of the remaining 24 nominees, 10 are interested and available to actively participate in FIFRA SAP meetings (see Unit IV.D.).

The following 14 individuals are not available to be considered further for membership at this time:

1. Joseph Braun, Ph.D., Brown University, Providence, Rhode Island.
2. Brenda Eskanazi, Ph.D., University of California-Berkeley, Berkeley, California.
5. Paul Hollenberg, Ph.D., University of Michigan Medical School, Ann Arbor, Michigan.
6. Mary K. O’Rourke, Ph.D., University of Arizona, Tucson, Arizona.
7. Angela Peace, Ph.D., Texas Tech University, Lubbock, Texas.
8. Kenneth Portier, Ph.D., Independent Consultant, Atlanta, Georgia.
9. Jason Rohr, Ph.D., University of Notre Dame, Notre Dame, Indiana.
11. Ronald Tjoerdema, Ph.D., University of California-Davis, Davis, California.
12. Tim Verslycke, Ph.D., Gradient Corp., Boston, Massachusetts.
13. Christopher P. Weis, Ph.D., National Institute of Environmental Health Sciences, Bethesda, Maryland.
14. William Wuest, Ph.D., Emory University, Atlanta, Georgia.

D. Interested and Available Nominees

The 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 (docket identification (ID) number EPA-HQ-OPP-2021-0293) and through the FIFRA SAP website at https://www.epa.gov/sap. The Agency, at this time, anticipates selecting new members to fill upcoming vacancies occurring on the Panel:

1. Dana Barr, Ph.D., Emory University, Atlanta, Georgia.
2. Veronica Berrocal, Ph.D., University of California, Irvine, California.
3. Assa Bradman, Ph.D., University of California, Berkeley, California.
4. Glenn Allen Burton, Ph.D., University of Michigan, Ann Arbor, Michigan.
5. Celia Chen, Ph.D., Dartmouth College, Hanover, New Hampshire.
7. Valery Forbes, Ph.D., University of Minnesota, Minneapolis, Minnesota.
8. Cheryl A. Murphy, Ph.D., Michigan State University, East Lansing, Michigan.


Michal Freedhoff.
Principal Deputy Assistant Administrator, Office of Chemical Safety and Pollution Prevention.

FEDERAL COMMUNICATIONS COMMISSION

Next Meeting of the North American Numbering Council

AGENCY: Federal Communications Commission.

ACTION: Notice.

SUMMARY: In this document, the Commission released a public notice announcing the meeting of the North American Numbering Council (NANC), which will be held via video conference and available to the public via live internet feed.

DATES: Wednesday, June 23, 2021. The meeting will come to order at 2:00 p.m.

ADDRESSES: The meeting will be conducted via video conference and available to the public via the internet at http://www.fcc.gov/live.


SUPPLEMENTARY INFORMATION: The NANC meeting is open to the public on the internet via live feed from the FCC’s web page at http://www.fcc.gov/live. Open captioning will be provided for this event. Other reasonable accommodations for people with disabilities are available upon request. Requests for such accommodations should be submitted via email to fcc504@fcc.gov or by calling the Consumer & Governmental Affairs Bureau at (202) 418–0530 (voice), (202) 418–0432 (TTY). Such requests should include a detailed description of the accommodation needed. In addition, please include a way for the FCC to contact the requester if more information is needed to fill the request. Please allow at least five days’ advance notice for accommodation requests; last minute requests will be accepted but may not be possible to accommodate.

Members of the public may submit comments to the NANC in the FCC’s Electronic Comment Filing System, ECFS, at www.fcc.gov/ecfs. Comments to the NANC should be filed in CC Docket No. 92–237. This is a summary of the Commission’s document in CC Docket No. 92–237, DA 21–578, released May 14, 2021.

Proposed Agenda: At the June 23 meeting, the NANC will consider and vote on recommendations from the Numbering Administration Oversight working group on the North American Numbering Plan Billing & Collection Fund Size Projections and Contributions Factor, as well as an evaluation of the performance of the Billing & Collection Agent, Welch LLP. The NANC will also hear reports from the Billing & Collection Agent, Welch LLP, the Numbering Administration Oversight working group on the Reassigned Numbers Database and Administrator, and routine status reports from the North American Portability Management, LLC and the Secure Telephone Identity Governance Authority. This agenda may be modified at the discretion of the NANC Chair and the Designated Federal Officers (DFO) (5 U.S.C. App 2 § 10(a)(2))

Federal Communications Commission.

Daniel Kahn, Associate Bureau Chief, Wireline Competition Bureau.

BILLING CODE 6712–01–P

FEDERAL DEPOSIT INSURANCE CORPORATION

Agency Information Collection Activities: Proposed Collection Renewal; Comment Request [OMB No. 3064–0082; and –0084]

AGENCY: Federal Deposit Insurance Corporation (FDIC).

ACTION: Agency information collection activities: Submission for OMB review; comment request.

SUMMARY: The FDIC, as part of its obligations under the Paperwork