

Prevention Act (PPA), 42 U.S.C. 13101 *et seq.*, and other applicable statutes. The 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 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 SACC currently consists of 26 members. When needed, the committee will be assisted in their reviews by ad hoc participants with specific expertise in the topics under consideration.

B. Purpose of This Public Meeting

EPA is announcing the rescheduled meeting dates for the TSCA SACC's public meeting to review PV29 that was previously announced in the **Federal Register** on November 30, 2018 (83 FR 61629) (FRL-9983-08). As previously announced, the public is invited to comment on the draft risk evaluation for PV29 and related documents, including the draft charge questions, in advance of and during this peer review meeting. The TSCA SACC will consider these comments during their discussions. See 83 FR 57473, November 15, 2018 (FRL-9986-45) and 84 FR 16011, April 17, 2019 (FRL-9990-36), as corrected by 84 FR 16485, April 19, 2019.

The focus of the TSCA SACC meeting is to peer review the Agency's draft risk evaluation of PV29. After the peer review process, EPA will consider reviewer comments and recommendations, and public comments to finalize the risk evaluation. The discussion of charge questions, for scope and clarity, originally planned for January 8, 2019, will be folded into the rescheduled in-person meeting.

As previously announced, approximately one hour of the TSCA SACC's in-person meeting will be closed to the public for the TSCA SACC to consider and discuss material that has been claimed as CBI provided to the TSCA SACC as background for the draft risk evaluation for PV29. In accordance with FACA section 10(d), 5 U.S.C. App. 2, and section (c)(4) of the Government in the Sunshine Act, 5 U.S.C. 552b, this approximately one-hour session of the

TSCA SACC will be closed to the public to avoid the potential disclosure of CBI, which is protected from disclosure by statute. The Administrator's determination for a closed meeting is available in the docket.

C. TSCA SACC Documents and Meeting Minutes

EPA's draft risk evaluation for PV29, related supporting materials, and draft charge/questions to the TSCA SACC are available on the TSCA SACC website and in the meeting docket. In addition, the Agency 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/tsc-peer-review>.

TSCA SACC will prepare the Meeting Minutes and Final Report document summarizing its recommendations to the Agency approximately 90 days after the meeting. The meeting minutes will be posted on the TSCA SACC website and placed in the meeting docket.

Authority: 15 U.S.C. 2625(o) *et seq.*; 5 U.S.C. Appendix 2 *et seq.*

Dated: May 1, 2019.

Hayley Hughes,

Director, Office of Science Coordination and Policy.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2019-0008; FRL-9991-15]

Pesticide Product Registration; Receipt of Applications for New Uses

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA has received applications to register new uses for pesticide products containing currently registered active ingredients. Pursuant to the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), EPA is hereby providing notice of receipt and opportunity to comment on these applications.

DATES: Comments must be received on or before June 10, 2019.

ADDRESSES: Submit your comments, identified by the Docket Identification (ID) Number, and the File Symbol or EPA Registration Number of interest as

shown in the body of this document, 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:** 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 <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:

Michael Goodis, Registration Division (7505P), main telephone number: (703) 305-7090, email address:

RDfrNotices@epa.gov. The mailing address for each contact person is:

Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW, Washington, DC 20460-0001. As part of the mailing address, include the contact person's name, division, and mail code. The division to contact is listed at the end of each application summary.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

You may be potentially affected by this action if you are an agricultural producer, food manufacturer, or pesticide manufacturer. The following list of North American Industrial Classification System (NAICS) codes is not intended to be exhaustive, but rather provides a guide to help readers determine whether this document applies to them. Potentially affected entities may include:

- Crop production (NAICS code 111).
- Animal production (NAICS code 112).
- Food manufacturing (NAICS code 311).
- Pesticide manufacturing (NAICS code 32532).

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

1. **Submitting CBI.** Do not submit this information to EPA through [regulations.gov](http://www.regulations.gov) or email. Clearly mark the part or all of the information that you claim to be CBI. For CBI

information in a disk or CD-ROM that you mail to EPA, mark the outside of the disk or CD-ROM as CBI and then identify electronically within the disk or CD-ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

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

II. Registration Applications

EPA has received applications to register new uses for pesticide products containing currently registered active ingredients. Pursuant to the provisions of FIFRA section 3(c)(4) (7 U.S.C. 136a(c)(4)), EPA is hereby providing notice of receipt and opportunity to comment on these applications. Notice of receipt of these applications does not imply a decision by the Agency on these applications.

III. New Uses

1. *EPA registration number(s):* 100–1609, 100–1601, 100–1602, 100–1603, 100–1605, and new seed treatment product 100–RAUI. *Docket ID number:* EPA–HQ–OPP–2018–0688. *Applicant:* Syngenta Crop Protection, LLC, P.O. Box 18300, Greensboro, NC 27419. *Active ingredient:* Pydiflumetofen. *Product type:* Fungicide. *Proposed use(s):* Foliar use proposed on Berries, Low Growing Crop Subgroup 13–07G (except cranberry and blueberry); *Brassica* Head and Stem Crop Group 5–16; *Brassica* Leafy Greens Subgroup 4–16B; Bulb Vegetable Crop Group 3–07A; Bulb Vegetable Crop Group 3–07B; Bushberry Crop Subgroup 13–07B; Citrus Crop Subgroup 10–10; Cottonseed Subgroup 20C; Edible-podded Legume Vegetables Subgroup 6A; Leaves of Root and Tuber Vegetables, Crop Group 2; Succulent Shelled Pea and Bean Subgroup 6B; Pome Fruit Crop Group 11–10; Root Vegetable Crop Group 1A; Sorghum; Stone Fruit Crop Group 12–12 (Subgroups 12A, 12B and 12C); Sunflower, Oilseed Subgroup 20B; and Tree Nuts Crop Group 14–12. Seed treatment proposed for use on Rapeseed Subgroup 20A and soybean. Contact: RD.

2. *EPA registration number:* 7969–262. *Docket ID number:* EPA–HQ–OPP–

2019–0060. *Applicant:* BASF Corporation, 26 Davis Drive, Research Triangle Park, NC 27709. *Active ingredient:* Topramezone. *Product type:* Herbicide. *Proposed use:* African Marigold. Contact: RD.

3. *EPA registration number:* 10163–357. *Docket ID number:* EPA–HQ–OPP–2017–0565. *Applicant:* Gowan Company LLC, P.O. Box 5569, Yuma, AZ 85366–5569. *Active ingredient:* Extract of *Swinglea glutinosa*. *Product type:* Biochemical fungicide. *Proposed use:* Addition of aerial and chemigation applications and residential uses. Contact: BPPD.

4. *EPA registration numbers:* 67690–6 and 67690–78. *Docket ID number:* EPA–HQ–OPP–2019–0074. *Applicant:* SePRO Corporation, 11550 North Meridian Street, Suite 600, Carmel, IN 46032. *Active ingredient:* Fluridone. *Product type:* Herbicide. *Proposed uses:* Avocado, mandarin, pistachio, pomegranate, and stone fruit crop group 12–12. Contact: RD.

5. *EPA registration number(s):* 71512–2 and 71512–3. *Docket ID number:* EPA–HQ–OPP–2018–0832. *Applicant:* ISK Biosciences Corporation, 7470 Auburn Road, Suite A, Concord, OH 44077. *Active ingredient:* Cyazofamid. *Product type:* Fungicide. *Proposed use(s):* New use on ginseng and greenhouse cucumber; conversion from *Brassica*, leafy greens, subgroup 5B to *Brassica*, leafy greens, subgroup 4–16B; conversion from leafy greens subgroup 4A to Leafy greens subgroup 4–16A; conversion from *Brassica*, head and stem, subgroup 5A to Vegetable, *Brassica*, head and stem group 5–16 and kohlrabi. Contact: RD

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

Dated: May 1, 2019.

Delores Barber,

Director, Information Technology and Resources Management Division, Office of Pesticide Programs.

[FR Doc. 2019–09490 Filed 5–8–19; 8:45 am]

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EXECUTIVE OFFICE OF THE PRESIDENT

Office of National Drug Control Policy

Paperwork Reduction Act; Proposed Collection; Comment Request

AGENCY: Office of National Drug Control Policy.

ACTION: 60-Day notice and request for comments. Revisions of Currently Approved Collection: Drug-Free Communities Support Program National Evaluation.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, the Office of National Drug Control Policy (ONDCP) announces it will submit to the Office of Management and Budget (OMB) and Office of Information and Regulatory Affairs (OIRA) an information collection request.

DATES: ONDCP encourages and will accept public comments on or before 60 days after the date of this publication.

ADDRESSES: Address all comments in writing within 60 days to Helen Hernandez. Email is the most reliable means of communication. Ms. Hernandez's email address is HHernandez@ondcp.eop.gov. Mailing address is: Executive Office of the President, Office of National Drug Control Policy, Drug-Free Communities (DFC) Support Program, 1800 G Street NW, Suite 9110, Washington, DC 20006. Copies of documents submitted to OMB and other information is available from Ms. Hernandez who may be contacted at 202–395–6665.

SUPPLEMENTARY INFORMATION:

Abstract: ONDCP administers the Drug-Free Communities (DFC) Support Program and Community-Based Coalition Enhancement Grants to Address Local Drug Crisis (CARA Local Drug Crisis) Programs. The DFC Program has two primary goals: To reduce youth substance abuse, and to support community anti-drug coalitions by establishing, strengthening, and fostering collaboration among public and private agencies. The CARA Local Drug Crisis grant program funds current or former DFC grant award recipients to focus on preventing and reducing the abuse of opioids or methamphetamines and the abuse of prescription medications among youth ages 12–18 in communities throughout the United States.

Congress mandates an evaluation of the DFC program to determine its effectiveness in meeting objectives (see 21 U.S.C. 1521 *et al.*). Under the CARA Local Drug Crisis program statute, CARA Local Drug Crisis data collection is authorized and required by Public Law 114–198 Sec 103, “a grant under this section shall be subject to the same evaluation requirements and procedures as the evaluation requirements and procedures imposed on the recipients of a grant under the Drug-Free Communities Act of 1997, and may also include an evaluation of the effectiveness at reducing abuse of opioids or methamphetamines”. ONDCP awarded a contract for a DFC grant oversight system at the end of 2014, following a competitive request for proposals process. The DFC