sabproduct.nsf/fedrgstr\_activites/ CCL%204?OpenDocument.

Technical Contacts: Any technical questions concerning EPA's draft CCL 4 should be directed to Ms. Meredith Russell in the EPA Office of Water, by telephone at (202) 564–0814 or by email at Russell.Meredith@epa.gov.

Availability of Meeting Materials: Prior to the meeting, the review documents, agenda and other materials will be accessible through the calendar link on the blue navigation bar at <a href="http://www.epa.gov/sab/">http://www.epa.gov/sab/</a>. Materials may also be accessed at the URL provided above.

Procedures for Providing Public Input: Public comment for consideration by EPA's federal advisory committees and panels has a different purpose from public comment provided to EPA program offices. Therefore, the process for submitting comments to a federal advisory committee is different from the process used to submit comments to an EPA program office. Federal advisory committees and panels, including scientific advisory committees, provide independent advice to the EPA. Interested members of the public may submit relevant information on the topic of this advisory activity, and/or the group conducting the activity, for the SAB to consider during the advisory process. Input from the public to the SAB will have the most impact if it provides specific scientific or technical information or analysis for SAB committees and panels to consider or if it relates to the clarity or accuracy of the technical information. Members of the public wishing to provide comment should contact the DFO directly. Oral Statements: In general, individuals or groups requesting an oral presentation at the teleconference will be limited to three minutes. Interested parties wishing to provide comments should contact Ms. Sanzone, DFO, in writing (preferably via email) at the contact information noted above by July 27, 2015, to be placed on the list of public speakers for the meeting. Written Statements: Written statements will be accepted throughout the advisory process; however, for timely consideration by Committee members, statements should be supplied to the DFO (preferably via email) at the contact information noted above by July 27, 2015. It is the SAB Staff Office general policy to post written comments on the Web page for advisory meetings. Submitters are requested to provide an

unsigned version of each document because the SAB Staff Office does not publish documents with signatures on its Web sites. Members of the public should be aware that their personal contact information, if included in any written comments, may be posted to the SAB Web site. Copyrighted material will not be posted without explicit permission of the copyright holder.

Accessibility: For information on access or services for individuals with disabilities, please contact Ms. Sanzone at the contact information provided above. To request accommodation of a disability, please contact Ms. Sanzone preferably at least ten days prior to the meeting to give EPA as much time as possible to process your request.

Dated: June 30, 2015.

#### Thomas H. Brennan,

Deputy Director, EPA Science Advisory Board Staff Office.

[FR Doc. 2015-16721 Filed 7-7-15; 8:45 am]

BILLING CODE 6560-50-P

# ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2015-0393; FRL-9929-24]

# Registration Review Interim Decisions; Notice of Availability

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

**ACTION:** Notice.

**SUMMARY:** This notice announces the availability of EPA's interim registration review decisions for the pesticides listed in the table in Unit II of this notice. 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, that the pesticide can perform its intended function without causing unreasonable adverse effects on human health or the environment. 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. This document also announces the Agency's closure of the registration review docket diclofopmethyl. All pesticide products containing diclofop-methyl have been

**FOR FURTHER INFORMATION CONTACT:** For pesticide specific information, contact

the Chemical Review Manager identified in the table in Unit II. for the pesticide of interest.

For general information on the registration review program, contact: Richard Dumas, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 308–8015; email address: dumas.richard@epa.gov.

### SUPPLEMENTARY INFORMATION:

#### I. General Information

### A. 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 Chemical Review Manager identified in the table in Unit II. for the pesticide of interest.

# B. How can I get copies of this document and other related information?

The docket for this action, identified by docket identification (ID) number EPA-HQ-OPP-2015-0393, is available at http://www.regulations.gov or at the Office of Pesticide Programs Regulatory Public Docket (OPP Docket) in the **Environmental Protection Agency** Docket Center (EPA/DC), West William Jefferson Clinton Bldg., Rm. 3334, 1301 Constitution Ave. NW., Washington, DC 20460-0001. The Public Reading Room is open from 8:30 a.m. to 4:30 p.m., Monday through Friday, excluding legal holidays. The telephone number for the Public Reading Room is (202) 566-1744, and the telephone number for the OPP Docket is (703) 305-5805. Please review the visitor instructions and additional information about the docket available at http://www.epa.gov/dockets.

### II. What action is the agency taking?

Pursuant to 40 CFR 155.58(c), this notice announces the availability of EPA's interim registration review decisions for the pesticides in the following table:

| TABLE—REGIS | tration Revie | EW INTERIM DI | ECISIONS |
|-------------|---------------|---------------|----------|
|             |               |               |          |

| Registration review case name and No.               | Docket ID No.        | Chemical review manager and contact information                 |
|---|----------------------|---|
| Acetic acid and sodium diacetate, 4001              | EPA-HQ-OPP-2008-0016 | Cathryn Britton, britton.cathryn@epa.gov, (703) 308–0136.       |
| Fosetyl-Al, 0646                                    | EPA-HQ-OPP-2006-0379 | Ricardo Jones, jones.ricardo@epa.gov, (703) 347–0493.           |
| Picaridin, 7433                                     | EPA-HQ-OPP-2014-0341 | Ricardo Jones, jones.ricardo@epa.gov, (703) 347–0493.           |
| Sodium fluoride, 3132                               | EPA-HQ-OPP-2014-0655 | SanYvette Williams, williams.Sanyvette@epa.gov, (703) 305–7702. |
| Yellow mustard seed/sulfonic acid salts, 7619/7618. | EPA-HQ-OPP-2014-0762 | Roy Johnson, johnson.roy@epa.gov, (703) 347–0492.               |

The registration review final decisions for these cases are dependent on the assessments of listed species under the Endangered Species Act (ESA), determinations on the potential for endocrine disruption, and/or pollinator risk assessments.

Acetic acid and sodium diacetate (Interim Decision). The registration review docket for acetic acid and sodium diacetate opened in March 2008. Acetic acid and sodium diacetate are two different active ingredients; sodium diacetate is a salt of acetic acid. Acetic acid is used as a preservative for post-harvest stored grains and have intended for livestock feed. It is also applied as a non-selective herbicide for control of broadleaf weeds and grasses. Sodium diacetate is a fungicide and bactericide registered to control molds and bacteria. It is applied to hay to prevent spoilage and to silage as an aid in fermentation. The Agency has determined that previous human health assessments for acetic acid and sodium diacetate are sufficient for registration review and no human health risks of concern have been identified. The Agency completed a comprehensive ecological risk assessment for the nonselective herbicide use of acetic acid, including an endangered species assessment, and a qualitative ecological risk assessment for sodium diacetate. The Agency has made a No Effect determination for acetic acid used as a nonselective herbicide and all currently registered uses of sodium diacetate for all non-target organisms. EPA published the Acetic Acid and Sodium Diacetate Proposed Interim Decision in December 2014. One comment was received from the Center for Biological Diversity concurring with EPA's No Effect determination. No risk mitigation measures for human health or ecological effects are included in the interim

Fosetyl-Al (Interim Decision). Fosetyl-Al is a phosphonate fungicide registered for use to control various oomycyete

pathogens that cause fungal diseases on numerous crops. Fosetyl-Al is registered for use on agricultural crops including avocado, caneberries, citrus, grape, stone fruit, strawberry, certain tree nuts, tobacco, and certain vegetables. It is also registered for commercial use on ornamentals, turf, and conifer nurseries. EPA conducted quantitative risk assessments for both human health and ecological risk. The Agency also completed a partial screening level endangered species assessment, making a No Effects determination for listed species of fish, aquatic invertebrates, aquatic plans, and monocot plants. No human health risks of concern were identified. The ecological risk assessment indicated potential risks to birds, mammals, terrestrial and aquatic plants, and terrestrial invertebrates. In the Federal Register of December 24, 2014 (79 FR 77480) (FRL-9919-24), the Agency issued its *Proposed Interim* Registration Review Decision for fosetyl-Al for public comment. Fourteen comments were received, including comments on the Agency's proposal to increase the Restricted Entry Interval, require additional pollinator data and add pollinator advisory language to labels. The Agency's Interim Decision modifies the application directions for fosetyl-Al to mitigate ecological risks and updates labels to reflect current Agency policy for worker protection. It does not require additional pollinator data or pollinator advisory language on the product labels.

Picaridin (Interim Decision). Picaridin is a broad-spectrum insect repellent registered for use against biting flies, chiggers, fleas, mosquitoes, and ticks. Picaridin is labeled for use on human skin and footwear. The Agency completed qualitative ecological and human health risk assessments for picaridin, and found no risks of concern. The Agency has made a No Effects determination under the Endangered Species Assessment (ESA) for all listed species and No

Modification of designated critical habitat for such species. In the **Federal Register** of December 24, 2014 (79 FR 77480) (FRL—9919—24), EPA issued the *Combined Work Plan and Proposed Interim Registration Review Decision*, indicating that no additional data would be needed for the picaridin registration review. Two comments were received, neither of which modified the proposed decision or work plan for picaridin. No risk mitigation measures for human health or ecological effects are included in the interim decision.

Sodium fluoride (Interim Decision). Sodium fluoride is registered for use as a wood preservative to protect the groundline portion of existing wooden utility poles. It is formulated as an impregnated pole wrap material. This use is not expected to result in direct or indirect food or drinking water exposure. Occupational and residential exposure is minimal by the dermal and inhalation routes, and the Agency has determined that a human health risk assessment was not needed. Based on the lack of potential exposure and toxic effects to fish, aquatic invertebrates, and birds, the Agency has made a No Effect determination for federally listed species and designated critical habitat.

Yellow mustard seed/sulfonic acid salts (Interim Decision). Yellow mustard seed and sulfonic acid salts are coformulated as a rodenticide registered for the control of ground squirrels in rangelands, ornamental plantings, seed orchards and nurseries, golf courses, parks, and rights-of-way. The product is delivered into burrows occupied by these rodents through a modified handheld spray wand with an aspirating muzzle to facilitate foaming action. The Agency relied on qualitative assessments conducted for yellow mustard seed/sulfonic acid salts at the time of the initial registration, and the Agency did not believe that updated or quantitative assessments were needed for registration review. No human health risks of concern or risks of

concern to non-listed species have been identified. No risk mitigation measures for human health or ecological effects are included in the interim decision.

Case Closure for Diclofop-methyl (PC Code: 110902, Case: 2160). Diclofopmethyl is an herbicide which was labeled for use on wheat, barley, and golf course turf. On October 23, 2014, the Agency received a request for voluntary cancellation of diclofopmethyl from the technical and end-use registrants; Bayer CropScience and Bayer Environmental Science, respectively. EPA subsequently issued a Federal Register notice announcing receipt of the request (FRL-9396-04), and allowed a 30-day period for public comment on the request. No substantive comments were received, and on June 10, 2015, EPA issued the cancellation order for all remaining registrations of products containing diclofop-methyl (FRL-9968-03), which sets out the existing stocks policy for such products. With the cancellation of all remaining diclofop-methyl products, the Agency is announcing the closure of the registration review case for the active ingredient.

Pursuant to 40 CFR 155.57, a registration review decision is the Agency's determination whether a pesticide meets, or does not meet, the standard for registration in FIFRA. EPA has considered the pesticides listed in the table in this unit in light of the FIFRA standard for registration. The Interim Decision documents for these pesticides in the docket describe the Agency's rationale for issuing a registration review interim decision for this pesticide.

In addition to an interim registration review decision document, the registration review docket for each of these pesticides may also include other relevant documents related to the registration review of the case. A proposed interim registration review decision was previously posted to each docket and the public was invited to submit any comments or new information relevant to the proposal.

EPA has addressed the substantive comments and information received during the 60-day comment period in the discussion for each pesticide listed in this document. During the 60-day comment period, no public comments were received for any of these cases that resulted in changes in the Agency's interim decisions.

Pursuant to 40 CFR 155.58(c), the registration review case docket for each pesticide discussed in this notice will remain open until all actions required in the interim decision have been completed.

Background on the registration review program is provided at: http://www2.epa.gov/pesticide-reevaluation. Links to earlier documents related to the registration review of the pesticide cases identified in this notice are provided in the Pesticide Chemical Search data base accessible at: http://iaspub.epa.gov/apex/pesticides/f?p=chemicalsearch.

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

Dated: June 26, 2015.

### Richard P. Keigwin, Jr.,

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

[FR Doc. 2015–16406 Filed 7–7–15; 8:45 am]

BILLING CODE 6560-50-P

# ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2015-0386; FRL-9929-23]

Registration Review; Draft Human Health and Ecological Risk Assessments; Notice of Availability

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

ACTION: Notice.

**SUMMARY:** This notice announces the availability of EPA's draft human health and ecological risk assessments for the registration reviews of flufenacet, flurprimidol, propoxur, and sodium acifluorfen, and opens a public comment period on these documents. In addition, this notice announces both the opening of the registration review docket for thidiazuron and the availability of the registration review draft human health and ecological risk assessments for thidiazuron. The Agency is opening a public comment period on both the Preliminary Work Plan and the draft risk assessments for thidiazuron. 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.

**DATES:** Comments must be received on or before September 8, 2015.

ADDRESSES: Submit your comments, identified by docket identification (ID) number for the specific pesticide of interest provided in Table 1 of Unit III, 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: For pesticide specific information contact the Chemical Review Manager listed as the contact in Table 1 of Unit III.

For general questions on the registration review program, contact: Richard Dumas, Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460–0001; telephone number: (703) 308–8015; email address: dumas.richard@epa.gov.

### SUPPLEMENTARY INFORMATION:

#### I. General Information

A. 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 Chemical Review Manager for the case in question, listed in Table 1 of Unit III of this notice.

- 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 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