BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP- 2014-0763; FRL-9918-44]

Registration Review; Pesticide Dockets Opened for Review and Comment

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: With this notice, EPA is opening the public comment period for several registration reviews. 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. Registration review dockets contain information that will assist the public in understanding the types of information and issues that the Agency may consider during the course of registration reviews. 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. For flufenpyr-ethyl, EPA is seeking comment on the preliminary work plan, the ecological problem formulation, and the human health draft risk assessment. For Sodium Fluoride, Yellow Mustard Seed and Sulfonic Acid, EPA is seeking comment on the Combined Work Plan, Summary Document, and Proposed Interim Registration Review Decision, which includes the human health and ecological risk assessments. This notice also announces a registration review case closure for thiacloprid.

DATES: Comments must be received on or before [*INSERT DATE 60 DAYS AFTER DATE OF PUBLICATION IN THE* **FEDERAL REGISTER**].

ADDRESSES: Submit your comments identified by the docket identification (ID) number for the specific pesticide of interest provided in the table in Unit III.A., 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** for the pesticide of interest identified in the table in Unit III.A.

For general information 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; fax number: (703) 308-8005; 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, farmworker, 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.

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 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*.

3. *Environmental justice*. EPA seeks to achieve environmental justice, the fair treatment and meaningful involvement of any group, including minority and/or low income populations, in the development, implementation, and enforcement of environmental laws, regulations, and policies. To help address potential environmental

justice issues, the Agency seeks information on any groups or segments of the population who, as a result of their location, cultural practices, or other factors, may have atypical or disproportionately high and adverse human health impacts or environmental effects from exposure to the pesticide(s) discussed in this notice, compared to the general population.

II. Authority

EPA is initiating its review of the pesticides identified in this notice pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) (7 U.S.C. 136a(g)) 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.

III. Registration Reviews

A. What Action is the Agency Taking?

As directed by FIFRA section 3(g), EPA is reviewing the pesticide registrations identified in the table in this unit to assure that they continue to satisfy the FIFRA standard for registration--that is, they can still be used without unreasonable adverse effects on human health or the environment. A pesticide's registration review begins when the Agency establishes a docket for the pesticide's registration review case and

opens the docket for public review and comment. At present, EPA is opening registration review dockets for the cases identified in the following table.

Table 1.--Registration Review Dockets Opening

Registration Review Case Name and Number	Docket ID Number	Chemical Review Manager or Regulatory Action Leader, Telephone Number, Email Address
3-methyl-cyclohexen-1-one (Case 6074)	EPA-HQ-OPP-2014-0671	Gina Burnett, (703) 605-0513 *burnett.gina@epa.gov*
Alkyl trimethylenediamines (ATMD) (Case 3014)	EPA-HQ-OPP-2014-0656	Donna Kamarei, (703) 347-0443 *kamarei.donna@epa.gov*
Boscalid (Case 7039)	EPA-HQ-OPP-2014-0199	Maria Piansay, (703) 308-8063 *piansay.maria@epa.gov*
Dikegulac sodium (Case 3061)	EPA-HQ-OPP-2014-0771	Matthew Manupella, (703) 347-0411 *manupella.matthew@epa.gov*
Ethoxyquin (Case 0003)	EPA-HQ-OPP-2014-0780	Khue Nguyen, (703) 347-0248 *nguyen.khue@epa.gov*
Fenpyroximate (Case 7432)	EPA-HQ-OPP-2014-0572	Miguel Zavala, (703) 347-0504 *zavala.miguel@epa.gov*
Flonicamid (Case 7436)	EPA-HQ-OPP-2014-0777	Ricardo Jones, (703) 347-0493 *jones.ricardo@epa.gov*
Fluazifop butyl, isomers (Case 2285)	EPA-HQ-OPP-2014-0779	Matthew Manupella, (703) 347-0411 *manupella.matthew@epa.gov*
Flufenpyr-ethyl (Case 7262)	EPA-HQ-OPP-2014-0768	Steven Snyderman, (703) 347-0249 *snyderman.steven@epa.gov*
HHT (Grotan) (Case 3074)	EPA-HQ-OPP-2014-0654	Tina Pham, (703) 308-0125 *pham.thao@epa.gov*
Metolachlor & s-Metolachlor (Case 0001)	EPA-HQ-OPP-2014-0772	Steven Snyderman, (703) 347-0249 *snyderman.steven@epa.gov*

Napthaleneacetic acid (Case 0379)	EPA-HQ-OPP-2014-0773	Christina Scheltema, (703) 308-2201 *scheltema.christina@epa.gov*
Oxadiazon (Case 2485)	EPA-HQ-OPP-2014-0782	Katherine St. Clair, (703) 347-8778 *stclair.katherine@epa.gov*
Oxyfluorfen (Case 2490)	EPA-HQ-OPP-2014-0778	Benjamin Askin, (703) 347-0503 *askin.benjamin@epa.gov*
Pentachlorophenol (Case 2505)	EPA-HQ-OPP-2014-0653	Sandra O'Neill, (703) 347-0141 *oneill.sandra@epa.gov*
Sodium fluoride (Case 3132)	EPA-HQ-OPP-2014-0655	SanYvette Williams, (703) 305-7702 *williams.sanyvette@epa.gov*
Sulfonic acid salts (Case 7619)	EPA-HQ-OPP-2014-0762	Roy Johnson, (703) 347-0492 *johnson.roy@epa.gov*
Triclopyr (Case 2710)	EPA-HQ-OPP-2014-0576	Brittany Pruitt, (703) 347-0289 *pruitt.brittany@epa.gov*
Yellow mustard seed (Case 7618)	EPA-HQ-OPP-2014-0762	Roy Johnson, (703) 347-0492 *johnson.roy@epa.gov*

For flufenpyr-ethyl (Case 7262), EPA is seeking comment on the preliminary work plan, the ecological problem formulation, and the human health draft risk assessment. For Sodium Fluoride (Case 3132), Yellow Mustard Seed (Case 7618) and Sulfonic Acid (Case 7619), EPA is seeking comment on the Combined Work Plan, Summary Document, and Proposed Interim Registration Review Decision, which includes the human health and ecological risk assessments. This notice also announces 1 case closure. On August 6, 2014, the Agency issued a product cancellation order in the **Federal Register** (79 FR 45798; FRL-9914-09) for all thiacloprid product registrations. Due to the cancellation of all registered thiacloprid products in the United States, the Agency closed the registration review case for thiacloprid. The "Notice of

Registration Review Case Closure for Thiacloprid" is available in docket EPA-HQ-OPP-2012-0218 at *http://www.regulations.gov*.

B. Docket Content

1. *Review dockets*. The registration review dockets contain information that the Agency may consider in the course of the registration review. The Agency may include information from its files including, but not limited to, the following information:

- An overview of the registration review case status.
- A list of current product registrations and registrants.
- **Federal Register** notices regarding any pending registration actions.
- **Federal Register** notices regarding current or pending tolerances.
- 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.

2. *Other related information*. More information on these cases, including the active ingredients for each case, may be located in the registration review schedule on the

Agency's website at *http://www.epa.gov/oppsrrd1/registration_review/schedule.htm*. Information on the Agency's registration review program and its implementing regulation may be seen at *http://www.epa.gov/oppsrrd1/registration_review*.

3. *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 audiographic or videographic 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: December 10, 2014.

Richard P. Keigwin, Jr.,
Director, Pesticide Re-Evaluation Division, Office of Pesticide Programs.

[FR Doc. 2014-29578 Filed 12/18/2014 at 8:45 am; Publication Date: 12/19/2014]