

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 submitting comments, remember to:

i. Identify the document by docket ID number and other identifying information (subject heading, **Federal Register** date and page number).

ii. Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.

iii. Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.

iv. Describe any assumptions and provide any technical information and/or data that you used.

v. If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.

vi. Provide specific examples to illustrate your concerns and suggest alternatives.

vii. Explain your views as clearly as possible, avoiding the use of profanity or personal threats.

viii. Make sure to submit your comments by the comment period deadline identified.

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

A copy of the draft white paper is available in the docket under docket ID number EPA-HQ-OPP-2013-0442.

II. Background

A. What action is the agency taking?

EPA uses PCA adjustment factors to modify modeled concentrations of pesticides in surface waters in accordance with land cover types (i.e., crops) associated with a pesticide's uses. PCA-adjusted concentrations are used as estimated drinking water concentrations in human health risk assessments. Previously, PCAs were generated for Hydrologic Unit Code 8 (HUC-8) regions (part of a hierarchical system for classifying and mapping drainage areas in the United States). In this current update, PCAs have been generated for watersheds delineated based on surface-source drinking water intakes (DWI) of community water systems (CWSs) across the United States

The new PCAs are an improvement over previously calculated PCAs in terms of their relevance to human health

risk assessment because they were derived for known drinking water sources. Out of 6,550 DWI locations, which both met the selection criteria for watershed delineation and passed a Quality Assurance screen, 74% (4,840) had delineated watersheds that also passed a Quality Assurance screen. Summary values are presented in the draft white paper along with detailed descriptions of their development and suggested procedures for their routine use in pesticide risk assessment.

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

The Agency's authority is FIFRA (7 U.S.C. 136-136y).

List of Subjects

Environmental protection, Community water systems, Drinking water exposure assessments, Health and safety, Percent cropped area, Pesticides and pests, Surface water intakes.

Dated: May 1, 2014.

Donald J. Brady,

Director, Environmental Fate and Effects Division, Office of Pesticide Programs.

[FR Doc. 2014-10693 Filed 5-8-14; 8:45 am]

BILLING CODE 6560-50-P

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2009-1017; FRL-9909-95]

Iprodione, Pendimethalin, and Permethrin; Notice of Receipt of Requests To Voluntarily Cancel and Amend Registrations To Terminate Certain Uses

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: In accordance with the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), EPA is issuing a notice of receipt of requests by the registrants to voluntarily amend their Iprodione, Pendimethalin, and Permethrin product registrations to delete one or more uses. The requests would delete Iprodione use on rice, Pendimethalin use on alfalfa, corn, garlic, onions, peanuts, sorghum, sugarcane and sunflower, and Permethrin use on dogs. The requests would not terminate the last Iprodione, Pendimethalin, or Permethrin products registered for use in the United States. EPA intends to grant these requests at the close of the comment period for this announcement unless the Agency receives substantive comments within the comment period that would merit its further review of the requests, or unless

the registrants withdraw its requests. If these requests are granted, any sale, distribution, or use of products listed in this notice will be permitted after the use has been deleted only if such sale, distribution, or use is consistent with the terms as described in the final order.

DATES: Comments must be received on or before June 9, 2014.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA-HQ-OPP-2009-1017, 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: John W. Pates, Jr., Pesticide Re-Evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460-0001; telephone number: (703) 308-8195; email address: pates.john@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, 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 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 submitting comments, remember to:

- i. Identify the document by docket ID number and other identifying information (subject heading, **Federal Register** date and page number).
- ii. Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.
- iii. Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.
- iv. Describe any assumptions and provide any technical information and/or data that you used.
- v. If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.

vi. Provide specific examples to illustrate your concerns and suggest alternatives.

vii. Explain your views as clearly as possible, avoiding the use of profanity or personal threats.

viii. Make sure to submit your comments by the comment period deadline identified.

II. Background on the Receipt of Requests To Cancel and/or Amend Registrations To Delete Uses

This notice announces receipt by EPA of requests from registrants FMC Agricultural Products, Drexel Chemical Company and Farnam Companies, Inc. to delete certain uses of Iprodione, Pendimethalin, and Permethrin product registrations. In letters dated April 3, 2014, February 6, 2014 and November 19, 2013, FMC Agricultural Products, Drexel Chemical Company and Farnam, respectively, requested EPA to delete certain uses of pesticide product registrations identified in Table 1 of Unit III. Specifically, FMC Agricultural Products Group voluntarily requested the deletion of Iprodione use on rice from several registrations listed in Table 1 of Unit II. The registrant also requested a 30-day comment period, and waived the 180-day comment period. Drexel Chemical Company voluntarily requested the deletion of Pendimethalin use on alfalfa, corn

(field, pop, sweet), garlic, onions (dry bulb, green, welsh), peanuts, sorghum (grain), sugarcane and sunflower from the Drexel Pendimethalin Technical registration. The registrant requested a 30-day comment period, and waived the 180-day comment period. Farnam Companies, Inc. voluntarily requested the deletion of use on dogs from the Permethrin Farnam Purge Insecticide registration. The registrant waived any comment period associated with their request for voluntary use deletion. The requests would not terminate the last Iprodione, Pendimethalin, or Permethrin products registered in the United States, or the last Iprodione, Pendimethalin, or Permethrin pesticide products registered in the United States for these uses.

III. What action is the agency taking?

This notice announces receipt by EPA of requests from registrants to delete certain uses of Iprodione, Pendimethalin, or Permethrin product registrations. The affected products and the registrants making the requests are identified in Tables 1 and 2 of this unit.

Unless a request is withdrawn by the registrant or if the Agency determines that there are substantive comments that warrant further review of this request, EPA intends to issue an order amending the affected registrations.

TABLE 1—IPRODIONE, PENDIMETHALIN, AND PERMETHRIN PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR AMENDMENT

Registration No.	Product name	Company	Uses to be deleted
000270–00279	Farnam Purge Pesticide	Farnam Companies, Inc	Use on dogs.
000279–09562	Iprodione Technical	FMC Corp. Agricultural Products Group.	Use on rice.
000279–09564	Rovral® brand 4 Flowable Fungicide	FMC Corp. Agricultural Products Group.	Use on rice.
000279–09565	Rovral® R Flowable Fungicide	FMC Corp. Agricultural Products Group.	Use on rice.
000279–09566	Rovral® brand WG Fungicide	FMC Corp. Agricultural Products Group.	Use on rice.
000279–09567	Rovral® 50 SP Fungicide	FMC Corp. Agricultural Products Group.	Use on rice.
000279–09569	Rovral® brand 75WG Fungicide	FMC Corp. Agricultural Products Group.	Use on rice.
019713–00600	Drexel Pendimethalin Technical	Drexel Chemical Company	Use on alfalfa, corn (field, pop, sweet), garlic, onions (dry bulb, green, welsh), peanuts, sorghum (grain), sugarcane and sunflower.

Table 2 of this unit includes the names and addresses of record for the registrants of the products listed in Table 1 of this unit, in sequence by EPA company number. This number corresponds to the first part of the EPA registration numbers of the products listed in Table 1 of this unit.

TABLE 2—REGISTRANTS REQUESTING VOLUNTARY AMENDMENTS

EPA company No.	Company name and address
270	Farnam Companies, Inc., 301 West Osborn Rd., Phoenix, AZ 85013.

TABLE 2—REGISTRANTS REQUESTING VOLUNTARY AMENDMENTS—Continued

EPA company No.	Company name and address
279	FMC Corp. Agricultural Products Group, 1735 Market St., RM 1978, Philadelphia, PA 19103.

TABLE 2—REGISTRANTS REQUESTING VOLUNTARY AMENDMENTS—Continued

EPA company No.	Company name and address
19713	Drexel Chemical Company, P.O. Box 13327, Memphis, TN 38113-0327.

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

Section 6(f)(1) of FIFRA provides that a registrant of a pesticide product may at any time request that any of its pesticide registrations be canceled or amended to terminate one or more uses. FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of any such request in the **Federal Register**.

Section 6(f)(1)(B) of FIFRA requires that before acting on a request for voluntary cancellation, EPA must provide a 30-day public comment period on the request for voluntary cancellation or use termination. In addition, FIFRA section 6(f)(1)(C) requires that EPA provide a 180-day comment period on a request for voluntary cancellation or termination of any minor agricultural use before granting the request, unless:

1. The registrants request a waiver of the comment period, or
2. The EPA Administrator determines that continued use of the pesticide would pose an unreasonable adverse effect on the environment.

The Iprodione, Pendimethalin, and Permethrin registrants have requested that EPA waive the 180-day comment period. Accordingly, EPA will provide a 30-day comment period on the proposed requests.

V. Procedures for Withdrawal of Requests

Registrants who choose to withdraw a request for use deletion should submit the withdrawal in writing to the person listed under **FOR FURTHER INFORMATION CONTACT**. If the products(s) have been subject to a previous cancellation action, the effective date of cancellation and all other provisions of any earlier cancellation action are controlling.

VI. Provisions for Disposition of Existing Stocks

Existing stocks are those stocks of registered pesticide products that are currently in the United States and that were packaged, labeled, and released for shipment prior to the effective date of the action. If the requests for voluntary amendments to delete uses are granted, the Agency intends to publish the

cancellation order in the **Federal Register**.

In any order issued in response to these requests for amendments to delete uses, EPA proposes to include the following provisions for the treatment of any existing stocks of the products listed in Table 1 of Unit III.

Once EPA has approved product labels reflecting the requested amendments to delete uses, registrants will be permitted to sell or distribute products under the previously approved labeling for a period of 18 months after the date of **Federal Register** publication of the cancellation order, unless other restrictions have been imposed. Thereafter, registrants will be prohibited from selling or distributing the products whose labels include the deleted uses identified in Table 1 of Unit III., except for export consistent with FIFRA section 17 or for proper disposal.

Persons other than the registrant may sell, distribute, or use existing stocks of products whose labels include the deleted uses until supplies are exhausted, provided that such sale, distribution, or use is consistent with the terms of the previously approved labeling on, or that accompanied, the deleted uses.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: May 2, 2014.

Michael Goodis,

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

[FR Doc. 2014-10694 Filed 5-8-14; 8:45 am]

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DEPARTMENT OF ENERGY

FEDERAL COMMUNICATIONS COMMISSION

Information Collection Being Submitted for Review and Approval to the Office of Management and Budget

AGENCY: Federal Communications Commission (FCC).

ACTION: Notice; request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burden and as required by the Paperwork Reduction Act (PRA) of 1995 (44 U.S.C. 3502-3520), the FCC invites the general public and other Federal agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: Whether the proposed collection of information is necessary for the proper performance of the functions of the

Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimates; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid OMB Control Number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid OMB Control Number.

DATES: Written PRA comments should be submitted on or before June 9, 2014. If you anticipate that you will be submitting PRA comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the FCC contact listed below as soon as possible.

ADDRESSES: Submit your PRA comments to Nicholas A. Fraser, Office of Management and Budget (OMB), via fax at 202-395-5167, or via the Internet at *Nicholas_A.Fraser@omb.eop.gov* and to Leslie F. Smith, Office of Managing Director (OMD), Federal Communications Commission (FCC), via the Internet at *Leslie.Smith@fcc.gov*. To submit your PRA comments by email, please send them to: *PRA@fcc.gov*.

FOR FURTHER INFORMATION CONTACT: Leslie F. Smith, Office of Managing Director (OMD), Federal Communications Commission (FCC), at 202-418-0217, or via the Internet at: *Leslie.Smith@fcc.gov*.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060-0972.
Title: Multi-Association Group (MAG) Plan Order, Parts 54 and 69 Filing Requirements for Regulation of Interstate Services of Non-Price Cap Incumbent Local Exchange Carriers and Interexchange Carriers.

Form Number(s): N/A.
Type of Review: Revision of a currently approved collection.

Respondents: Business or other for-profit.

Number of Respondents and Responses: 202 respondents; 69 responses.

Estimated Time per Response: 20 to 90 hours.

Frequency of Response: On occasion and three year reporting requirements.

Obligation to Respond: Required to obtain or retain benefits.