Wildlife Service (FWS) and the National Marine Fisheries Service (NMFS) (jointly referred to as ''the Services'') regarding potential effects from these pesticides to federally listed threatened or endangered species (listed species) and habitat designated as critical to such species. The result of consultation will be a biological opinion issued by the Services that expresses whether they believe the pesticide's use is likely to jeopardize the continued existence of any listed species or destroy or adversely modify habitat designated as critical to any listed species. If the Services determine there is likely jeopardy or adverse modification, they will provide reasonable and prudent alternatives to the action. If the Services conclude the action will result in ''take'' of any individuals of a listed species, they will specify reasonable and prudent measures to minimize such impact. The Agency will review and consider both the public comments received on the draft ecological risk assessments and, if provided, the information in the Service's biological opinions when developing its proposed registration review decisions.

As described in detail in the ''Clomazone Summary Document Registration Review: Initial Docket (January 2007), Section IV—Human Health Effects Scoping Document'' (see docket ID number EPA–HQ–OPP–2006–0113), the Agency believes that the human health assessments completed prior to registration review are adequate, and there are no dietary risks that exceed the Agency's level of concern. In addition, there are no residential uses of clomazone and all worker margins of exposure (MOEs) are below the Agency's level of concern. Thus, no additional human health data are needed for the registration review of clomazone.

Also, as described in detail in the ''Fomesafen Summary Document Registration Review: Initial Docket (March 2007), Section IV – Human Health Effects Scoping Document'' (see docket ID number EPA–HQ–OPP–2006–0239), the Agency believes that the human health assessments completed prior to registration review are adequate and there are no dietary risks that exceed the Agency's level of concern. In addition, there are no residential uses of fomesafen. The occupational scenarios do not result in risk concerns, with the exception of inhalation risks to mixer/loaders for aerial application. This risk was mitigated below the Agency's level of concern with the following change that is currently on the label: ''In addition, for aerial applications, mixers and loaders handling more than 140 gallons of Reflex Herbicide in any single workday must wear dust/mist filtering NIOSH-approved respirator with any N, R, P, or HE filter.''

1. *Other related information.* More information on EPA's review of these cases is available on the Registration Review Status web page, *http://www.epa.gov/oppsrrd1/registration_review/reg_review_status.htm.* Information such as the active ingredients in each case, may be found 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 is available at *http://www.epa.gov/oppsrrd1/registration_review.*

2. *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.

List of Subjects

Environmental protection, Registration review, Pesticides and pests.

Dated: April 13, 2009.

Richard P. Keigwin, Jr.,

Director, Special Review and Reregistration Review Program.

[FR Doc. E9–9231 Filed 4–21–09; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2007–0513; FRL–8410–5]

Triclosan; Notice of Receipt of Requests for Amendments To Delete Uses in Certain Pesticide Registrations

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: In accordance with section 6(f)(1) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), as amended, EPA is issuing a notice of receipt of request for amendments by registrants to delete uses in certain pesticide registrations. 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 amended to delete one or more uses. FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of any request in the **Federal Register**.

DATES: The deletions are effective October 19, 2009, unless the Agency receives a written withdrawal request on or before October 19, 2009. The Agency will consider a withdrawal request postmarked no later than October 19, 2009.

Users of these products who desire continued use on sites being deleted should contact the applicable registrant on or before October 19, 2009.

ADDRESSES: Submit your withdrawal request, identified by docket identification (ID) number EPA–HQ–OPP–2007–0513, by one of the following methods:

• *Mail*: Office of Pesticide Programs (OPP) Regulatory Public Docket (7502P), Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001.

• *Delivery*: OPP Regulatory Public Docket (7502P), Environmental Protection Agency, Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr., Arlington, VA. Deliveries are only accepted during the Docket Facility's normal hours of operation (8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays). Special arrangements should be made for deliveries of boxed information. The Docket Facility telephone number is (703) 305–5805.

FOR FURTHER INFORMATION CONTACT: Heather Garvie, Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 308–0034; e-mail address: *garvie.heather@epa.gov*.

SUPPLEMENTARY INFORMATION:

I. General Information

A. Does This Action Apply to Me?

This action is directed to the public in general. Although this action may be of particular interest to persons who produce or use pesticides, 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 information in this notice, consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

B. How Can I Get Copies of this Document and Other Related Information?

1. *Docket*. EPA has established a docket for this action under docket ID number EPA–HQ–OPP–2007-0513. Publicly available docket materials are available either in the electronic docket at *http://www.regulations.gov*, or, if only available in hard copy, at the Office of Pesticide Programs (OPP) Regulatory Public Docket in Rm. S–4400, One Potomac Yard (South Bldg.), 2777 S. Crystal Dr., Arlington, VA. The hours of operation of this Docket Facility are from 8:30 a.m. to 4 p.m., Monday through Friday, excluding legal holidays. The Docket Facility telephone number is (703) 305–5805.

2. *Electronic access*. You may access this **Federal Register** document electronically through the EPA Internet under the ''**Federal Register**'' listings at *http://www.epa.gov/fedrgstr*.

II. What Action is the Agency Taking?

This notice announces receipt by the Agency of applications from registrants to delete uses in certain pesticide registrations. These registrations are listed in Table 1 of this unit by registration number, product name, active ingredient, and specific uses deleted:

TABLE 1.—REQUESTS FOR AMENDMENTS TO DELETE USES IN CERTAIN PESTICIDE REGISTRATIONS

EPA Registration No.	Product Name	Active Ingredient	Delete from Label
73951-1	VIV-20	Triclosan	Materials preservative used in paint formulations

Users of these products who desire continued use on sites being deleted should contact the applicable registrant before October 19, 2009 to discuss withdrawal of the application for amendment. This 180–day period will also permit interested members of the public to intercede with registrants prior to the Agency's approval of the deletion.

Table 2 of this unit includes the names and addresses of record for all registrants of the products listed in Table 1 of this unit, in sequence by EPA company number.

TABLE 2.—REGISTRANTS REQUESTING AMENDMENTS TO DELETE USES IN CERTAIN PESTICIDE REGISTRATIONS

EPA Company Number	Company Name and Address
73951	Har-Met International, Inc. 60 Houk Road Doylestown, PA 18901

III. 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 amended to delete one or more uses. The FIFRA further provides that, before acting on the request, EPA must publish a notice of receipt of any such request in the **Federal Register**. Thereafter, the Administrator may approve such a request.

IV. Procedures for Withdrawal of Request

Registrants who choose to withdraw a request for use deletion must submit the withdrawal in writing to Heather Garvie using the methods in **ADDRESSES**. The Agency will consider written withdrawal requests postmarked no later than October 19, 2009.

V. Provisions for Disposition of Existing Stocks

The Agency has authorized the registrants to sell or distribute product under the previously approved labeling for a period of 18 months after approval of the revision, unless other restrictions have been imposed, as in special review actions.

List of Subjects

Environmental protection, Pesticides and pests, Antimicrobials, Triclosan.

Dated: April 7, 2009.

Betty Shackleford,

Acting Director, Antimicrobials Division, Office of Pesticide Programs

[FR Doc. E9–8993 Filed 4–21–09; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2007–0513; FRL–8410–6]

Triclosan, Notice of Receipt of Requests To Voluntarily Cancel Certain Pesticide Registrations

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: In accordance with section 6(f)(1) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), as amended, EPA is issuing a notice of receipt of request by registrants to voluntarily cancel certain pesticide registrations.

DATES: Unless a request is withdrawn by May 22, 2009 for registrations for which the registrant requested a waiver of the 180–day comment period, orders will be issued canceling these registrations. The Agency will consider withdrawal requests postmarked no later than May 22, 2009, whichever is applicable. Comments must be received on or before May 22, 2009, for those registrations where the 180–day comment period has been waived.

ADDRESSES: Submit your comments and your withdrawal request, identified by docket identification (ID) number EPA–HQ–OPP–2007–0513, by one of the following methods:

- *Federal eRulemaking Portal*: *http://www.regulations.gov*. Follow the on-line instructions for submitting comments.
- *Mail*: Office of Pesticide Programs (OPP) Regulatory Public Docket (7502P),