

TABLE 1.—LINDANE PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR CANCELLATION—Continued

Registration No.	Product name	Company
554-144	Lindane ST-40	AGSCO Inc
19713-61	Lindane Technical	Drexel Chemical Co
19713-191	Lindane Technical	Drexel Chemical Co
19713-387	Lindane Flowable	Drexel Chemical Co
19713-401	Lindane 30%	Drexel Chemical Co
82378-1	Lindane Technical	JLM International Inc.

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.

TABLE 2.—REGISTRANTS REQUESTING VOLUNTARY CANCELLATION

EPA Company No.	Company name and address
400	Chemtura USA Corporation 199 Benson Road Middlebury, Connecticut 06749
554	AGSCO Inc. PO Box 13458 Grand Forks North Dakota 58208-3458
19713	Drexel Chemical Co. 1700 Channel Avenue, PO Box 13327 Memphis, Tennessee 38113-0327
82378	JLM International Inc. 8675 Hidden River Parkway Tampa, Florida 33637

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**. Thereafter, following the public comment period,

the Administrator may approve such a request.

V. Procedures for Withdrawal of Request and Considerations for Reregistration of Lindane

Registrants who choose to withdraw a request for cancellation must submit such withdrawal in writing to the person listed under **FOR FURTHER INFORMATION CONTACT**, postmarked before September 22, 2006. This written withdrawal of the request for cancellation will apply only to the applicable FIFRA section 6(f)(1) request listed in this notice. If the products 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

If the request for voluntary cancellation is granted as discussed above, the effective date of cancellation will be the date of the cancellation order. EPA intends to issue a cancellation order that will allow continued sale and/or use of existing stocks until such stocks are exhausted, provided that such further sale and use is consistent with the terms of the previously approved labeling on, or that accompany, the applicable product. It is EPA's intent that the order will specifically prohibit any use of existing stocks that is not consistent with such previously approved labeling.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: August 14, 2006.

Debra Edwards,

Director, Special Review and Reregistration Division, Office of Pesticide Programs.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2005-0497; FRL-8084-5]

Propiconazole Reregistration Eligibility Decision

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the availability of EPA's Reregistration

Eligibility Decision (RED) for the pesticide propiconazole. The Agency's risk assessments and other related documents also are available in the propiconazole docket. Propiconazole is used as a conventional fungicide on agricultural crops, ornamentals, and turf and is used as an antimicrobial material preservative and wood preservative. EPA has reviewed propiconazole through the public participation process that the Agency uses to involve the public in developing pesticide reregistration and tolerance reassessment decisions. Through these programs, EPA is ensuring that all pesticides meet current health and safety standards.

FOR FURTHER INFORMATION CONTACT: Christina Scheltema, Special Review and Reregistration Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001; telephone number: (703) 308-2201; fax number: (703) 308-8005; e-mail address: scheltema.christina@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. If you have any questions regarding the applicability of this action to a particular entity, 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 identification (ID) number EPA-HQ-OPP-2005-0497. 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 Building), 2777 S. Crystal Drive 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. Background

A. What Action is the Agency Taking?

Under section 4 of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA), EPA is reevaluating existing pesticides to ensure that they meet current scientific and regulatory standards. EPA has completed a Reregistration Eligibility Decision (RED) for the pesticide propiconazole under section 4(g)(2)(A) of FIFRA. Propiconazole is used as a conventional fungicide on agricultural crops, ornamentals, and turf and is used as an antimicrobial material preservative and wood preservative. EPA has determined that the database to support reregistration is substantially complete and that currently registered products containing propiconazole are eligible for reregistration, provided the risks are mitigated either in the manner described in the RED or by another means that achieves equivalent risk reduction. Upon submission of any required product-specific data under section 4(g)(2)(B) and any necessary changes to the registration and labeling (either to address concerns identified in the RED or as a result of product-specific data), EPA will make a final reregistration decision under section 4(g)(2)(C) for products containing propiconazole.

EPA must review tolerances and tolerance exemptions that were in effect when the Food Quality Protection Act (FQPA) was enacted in August 1996, to ensure that these existing pesticide residue limits for food and feed commodities meet the safety standard established by the new law. Tolerances are considered reassessed once the safety finding has been made or a revocation occurs. EPA has reviewed and made the requisite safety finding for the propiconazole tolerances.

Although the Propiconazole RED was signed on July 18, 2006, certain components of the document, which did not affect the final regulatory decision, were undergoing final editing at that time. These components, including the list of additional generic data requirements, summary of labeling changes, appendices, and other relevant information, have been added to the Propiconazole RED document.

EPA is applying the principles of public participation to all pesticides

undergoing reregistration and tolerance reassessment. The Agency's Pesticide Tolerance Reassessment and Reregistration; Public Participation Process, published in the **Federal Register** on May 14, 2004 (69 FR 26819) (FRL-7357-9), explains that in conducting these programs, EPA is tailoring its public participation process to be commensurate with the level of risk, extent of use, complexity of issues, and degree of public concern associated with each pesticide. Due to its uses, risks, and other factors, propiconazole was reviewed through the modified 4-Phase public participation process. Through this process, EPA worked extensively with stakeholders and the public to reach the regulatory decisions for propiconazole.

The reregistration program is being conducted under Congressionally mandated time frames, and EPA recognizes the need both to make timely decisions and to involve the public. Because few substantive comments were received during the earlier comment period for this pesticide, and all issues related to this pesticide were resolved through consultations with stakeholders, no comment period is needed on this regulatory decision. The Agency therefore is issuing the Propiconazole RED without a comment period.

B. What is the Agency's Authority for Taking this Action?

Section 4(g)(2) of FIFRA as amended, directs that, after submission of all data concerning a pesticide active ingredient, "the Administrator shall determine whether pesticides containing such active ingredient are eligible for reregistration," before calling in product-specific data on individual end-use products and either reregistering products or taking other "appropriate regulatory action."

Section 408(q) of the Federal Food, Drug, and Cosmetic Act (FFDCA), 21 U.S.C. 346a(q), requires EPA to review tolerances and exemptions for pesticide residues in effect as of August 2, 1996, to determine whether the tolerance or exemption meets the requirements of section 408(b)(2) or (c)(2) of FFDCA. This review is to be completed by August 3, 2006.

List of Subjects

Environmental protection, Material preservatives, Pesticides and pests, Propiconazole, Triazole fungicides, Wood preservatives.

Dated: August 14, 2006.

Peter Caulkins,

Acting Director, Special Review and Reregistration Division, Office of Pesticide Programs.

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2006-0505; FRL-8073-2]

Notice of Filing of a Pesticide Petition for Establishment of Regulations for Residues of Myclobutanil in or on Soybean Commodities

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: This notice announces the initial filing of a pesticide petition proposing the establishment of regulations for residues of myclobutanil in or on soybean commodities.

DATES: Comments must be received on or before September 22, 2006.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA-HQ-OPP-2006-0505 and pesticide petition number (PP) 5F6997, 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), 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 Building), 2777 S. Crystal Drive, Arlington, VA. Deliveries are only accepted during the Docket'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 telephone number is (703) 305-5805.

Instructions: Direct your comments to docket ID number EPA-HQ-OPP-2006-0505. EPA's policy is that all comments received will be included in the docket without change and may be made available on-line at <http://www.regulations.gov>, including any personal information provided, unless the comment includes information claimed to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Do not submit information that you