

Agency's determination whether a pesticide meets, or does not meet, the standard for registration in FIFRA. EPA has considered tralomethrin and fenarimol in light of the FIFRA standard for registration. The tralomethrin and fenarimol final decision documents in the docket describe the Agency's rationale for issuing a registration review final decision for these pesticides.

In addition to the final registration review decision documents, the registration review docket for tralomethrin and fenarimol also includes other relevant documents related to the registration review of these cases. The proposed registration review decisions were posted to the docket and the public was invited to submit any comments or new information. During the 60-day comment period, no public comments were received. Pursuant to 40 CFR 155.58(c), the registration review case docket for tralomethrin and fenarimol will remain open until all actions required in the final decision have been completed.

Background on the registration review program is provided at: http://www.epa.gov/oppsrrd1/registration_review. Links to earlier documents related to the registration review of these pesticides are provided at: <http://www.epa.gov/pesticides/chemicalsearch/>.

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

Section 3(g) of FIFRA and 40 CFR part 155, subpart C, provide authority for this action.

List of Subjects

Environmental protection, Registration review, Pesticides and pests, Tralomethrin and Fenarimol.

Dated: November 9, 2012.

Richard P. Keigwin, Jr.,

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

[FR Doc. 2012-28213 Filed 11-20-12; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2005-0252; FRL-9370-2]

Iodomethane; Notice of Receipt of Request to Voluntarily Cancel Iodomethane Pesticide Registrations and Amend a Registration

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 a request by the registrant to voluntarily cancel the registrations of products containing the pesticide iodomethane. In addition, the registrant has amended the terms and conditions of registration for their iodomethane technical product so that as of January 1, 2013, Arysta LifeScience North America, LLC (Arysta) will not sell or distribute this product unless it bears a label statement. The registrant's request would terminate the last iodomethane products registered for use in the United States. EPA intends to grant this request 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 request. If EPA issues a final order granting this request, the sale, distribution, or use of the products listed in this notice will be permitted only in accordance with the terms as described in the final order.

DATES: Comments must be received on or before December 21, 2012.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA-HQ-OPP-2005-0252, 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.htm>.

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:

Andrea Mojica, Pesticide Re-evaluation Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave. NW., Washington, DC 20460-0001; telephone number: (703) 308-0122; fax number: (703) 308-8090; email address: mojica.andrea@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 [regulations.gov](http://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

This notice announces receipt by EPA of a request from Arysta to cancel all of its iodomethane product registrations. Iodomethane is a pre-plant soil fumigant used to control pests in soil where fruits, vegetables, ornamental plants, and turf will be grown. In a Memorandum of Agreement (MOA), Arysta and EPA agreed to cancel and amend the pesticide product registrations identified in Tables 1 and 2 of Unit III. Specifically, the MOA contains Arysta's irrevocable request that its end-use products, EPA Registration Numbers 66330-43, 66330-57, 66330-58, 66330-59, and 66330-60,

will be canceled effective December 31, 2012, and that its iodomethane technical product, EPA Registration Number 66330-44 will be canceled effective December 1, 2015. The MOA also adds a condition of registration to the technical product's registration that as of January 1, 2013, Arysta will not sell or distribute this product unless its label bears the following statement:

It is unlawful to use this product for any purpose in the United States, except for formulation of products intended for export consistent with the requirements of FIFRA section 17.

(The request for amendment is conditioned on the issuance of a cancellation order including the requested effective dates and existing

stocks provisions.) Granting the registrant's cancellation request would terminate the last iodomethane products registered in the United States.

III. What action is the agency taking?

This notice announces receipt by EPA of the request to cancel the iodomethane product registrations described in Unit II. The affected products and the registrant making the requests are identified in Tables 1-3 of this unit.

Unless the Agency receives substantive comments in response to this notice that warrant further review of this request, EPA intends to issue an order canceling the affected registrations on the requested effective dates.

TABLE 1—IODOMETHANE PRODUCT REGISTRATIONS WITH PENDING REQUESTS FOR CANCELLATION

Registration No.	Product name	Company
66330-43	Midas 98:2	Arysta LifeScience North America, LLC.
66330-44	Iodomethane Technical	Arysta LifeScience North America, LLC.
66330-57	Midas 50:50	Arysta LifeScience North America, LLC.
66330-58	Midas EC Bronze	Arysta LifeScience North America, LLC.
66330-59	Midas 33:67	Arysta LifeScience North America, LLC.
66330-60	Midas EC Gold	Arysta LifeScience North America, LLC.

TABLE 2—IODOMETHANE PRODUCT REGISTRATION WITH PENDING REQUESTS FOR AMENDMENT

Registration No.	Product name	Company
66330-44	Iodomethane Technical	Arysta LifeScience North America, LLC.

Table 3 of this unit includes the name and address of record for the registrant of the products listed in Table 1 and Table 2 of this unit. This number corresponds to the first part of the EPA registration numbers of the products listed in Table 1 and Table 2 of this unit.

TABLE 3—REGISTRANT REQUESTING VOLUNTARY CANCELLATION AND AMENDMENTS

EPA Company No.	Company name and address
66330	Arysta LifeScience North America, 15401 Weston Parkway, Suite 150, Cary, NC 27513.

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 iodomethane registrant has requested that EPA waive the 180-day comment period. Accordingly, EPA will provide a 30-day comment period on the proposed requests.

V. 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 request for voluntary cancellation is granted, the Agency intends to publish the cancellation order in the **Federal Register**. EPA intends to include in any such final order the following provisions for the treatment of any existing stocks of the product(s) listed in Tables 1 and 2 of Unit III.

In any final order granting Arysta's request for voluntary cancellation of its iodomethane technical/manufacturing-use product registration, as of the effective date of the cancellation order, all sale and distribution of existing stocks of Arysta's iodomethane technical/manufacturing-use product by Arysta shall be prohibited unless the sale or distribution is for proper disposal or is solely for purposes of export consistent with the requirements of section 17 of FIFRA. In any final order granting Arysta's request for voluntary cancellation of end-use product registrations:

1. As of the effective date of the cancellation order, Arysta is prohibited from distributing or selling existing stocks of end-use products, unless the

sale or distribution is for proper disposal, or is solely for export consistent with the requirements of FIFRA section 17;

2. As of the effective date of the cancellation order, persons other than Arysta are prohibited from distributing or selling existing stocks of Arysta's end-use products, unless the sale or distribution is for proper disposal, return to Arysta, or is intended solely for export consistent with the requirements of FIFRA section 17; and

3. As of the effective date of the cancellation order, no person may use any existing stocks of any of Arysta's end-use products.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: November 14, 2012.

Richard P. Keigwin, Jr.

*Director, Pesticide Re-evaluation Division,
Office of Pesticide Programs.*

[FR Doc. 2012-28210 Filed 11-20-12; 8:45 am]

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EXPORT-IMPORT BANK OF THE UNITED STATES

Sunshine Act Meeting

ACTION: Notice of a Partially Open Meeting of the Board of Directors of the Export-Import Bank of the United States.

TIME AND PLACE: Thursday, November 29, 2012 at 9:30 a.m. The meeting will be held at Ex-Im Bank in Room 321, 811 Vermont Avenue NW, Washington, DC 20571.

OPEN AGENDA ITEMS: Item No. 1: Ex-Im Bank Advisory Committee for 2013 (Additional New Member).

PUBLIC PARTICIPATION: The meeting will be open to public observation for Item No. 1 only.

FURTHER INFORMATION: For further information, contact: Office of the Secretary, 811 Vermont Avenue NW., Washington, DC 20571 (202) 565-3336.

Lisa V. Terry,

Assistant General Counsel.

[FR Doc. 2012-28417 Filed 11-19-12; 4:15 pm]

BILLING CODE 6690-01-P

FEDERAL COMMUNICATIONS COMMISSION

Information Collections Approved by the Office of Management and Budget (OMB)

AGENCY: Federal Communications Commission.

ACTION: Notice.

SUMMARY: The Federal Communications Commission has received Office of Management and Budget (OMB) approval for the following public information collections pursuant to the Paperwork Reduction Act of 1995 (44 U.S.C. 3501-3520). An agency may not conduct or sponsor a collection of information unless it displays a currently valid OMB control number, and no person is required to respond to a collection of information unless it displays a currently valid OMB control number. Comments concerning the accuracy of the burden estimates and any suggestions for reducing the burden should be directed to the person listed in the **FOR FURTHER INFORMATION CONTACT** section below.

FOR FURTHER INFORMATION CONTACT:

Nakesha Woodward, Wireline Competition Bureau, Telecommunications Access Policy Division at 202-418-7400 or email at Kesha.Woodward@fcc.gov.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060-0824.

OMB Approval Date: November 1, 2012.

OMB Expiration Date: November 30, 2015.

Title: Service Provider Identification Number (SPIN) and Contact Information Form, Report and Order, GN Docket No. 09-191 and WC Docket No. 07-52.

Form Number: FCC Form 498.

Estimated Annual Burden: 5,000 respondents; 5,000 responses; 1.5 hours per response; 7,500 burden hours per year; total annual cost burden N/A.

Obligation to Respond: Required to obtain or retain benefits. The statutory authority for this collection is contained in sections 1-4 and 254 of the Communications Act of 1934, as amended, 47 U.S.C. 151-154, 254, and Part 54 of the Commission's rules.

Nature and Extent of Confidentiality: The Commission notes that USAC must preserve the confidentiality of all data obtained from respondents and contributors to the universal service programs, must not use the data except for purposes of administering the universal service programs, and must not disclose data in company-specific form unless directed to do so by the Commission. With respect to the Service Provider Identification Number and Contact Information Form (FCC Form 498), USAC shall publish each participant's name, SPIN, and contact information via USAC's Web site. All other information, including financial institution account numbers or routing information, shall remain confidential.

Needs and Uses: The information collected by FCC Form 498 is used by USAC to disburse federal universal service support consistent with the specifications of eligible participants in the universal service programs. FCC Form 498 submissions also provide USAC with updated contact information so that USAC can contact universal service fund participants when necessary. Without such information, USAC would not be able to distribute support to the proper entities and this would prevent the Commission from fulfilling its statutory responsibilities under the Act to preserve and advance universal service.

Federal Communications Commission.

Bulah P. Wheeler,

Associate Secretary.

[FR Doc. 2012-28347 Filed 11-20-12; 8:45 am]

BILLING CODE 6712-01-P

FEDERAL MARITIME COMMISSION

Notice of Agreement Filed

The Commission hereby gives notice of the filing of the following agreement under the Shipping Act of 1984. Interested parties may submit comments on the agreement to the Secretary, Federal Maritime Commission, Washington, DC 20573, within ten days of the date this notice appears in the **Federal Register**. Copies of the agreements are available through the Commission's Web site (www.fmc.gov) or by contacting the Office of Agreements at (202) 523-5793 or tradeanalysis@fmc.gov.

Agreement No.: 012057-008.

Title: CMA CGM/Maersk Line Space Charter, Sailing and Cooperative Working Agreement Asia to USEC and PNW-Suez/PNW & Panama Loops.

Parties: A.P. Moller-Maersk A/S and CMA CGM S.A.

Filing Party: Mark J. Fink, Esq.; Cozen O'Connor; 1627 I Street, NW Suite 1100; Washington, DC 20006.

Synopsis: The amendment would provide for the deployment of the seventeenth vessel and revise the space allocations of the parties accordingly. The parties have requested expedited review.

By Order of the Federal Maritime Commission.

Dated: November 16, 2012.

Karen V. Gregory,

Secretary.

[FR Doc. 2012-28344 Filed 11-20-12; 8:45 am]

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