http://www.epa.gov/pesticides/reregistration/REDs/ddvp_ired.pdf).

The IRED was based in part on an irrevocable request from Amvac Chemical Corporation (Amvac), the sole technical product registrant, to cancel certain uses and include additional pest strip label restrictions on the DDVP technical product labels. Pursuant to section 6(f) of FIFRA, 7 U.S.C. 136d(f)(1), on June 30, 2006, the Agency published a notice in the **Federal Register** that it had received the request and sought comment on EPA's intention to grant the request and cancel the specified uses. (71 FR 37570)(FRL–8075–2). On October 20, 2006, EPA issued the final cancellation order granting Amvac's request. (71 FR 61968)(FRL–8075–8).

Specifically on May 9, 2006, Amvac submitted to EPA a request for cancellation of several existing DDVP products, uses and application methods, including the 100 gram pest strip, the total release fogger, use on lawn, turf and ornamentals, residential crack and crevice use, and hand held fogger applications in mushroom houses, greenhouses, and warehouses. Amvac also requested several label amendments further restricting residential use of pest strips and adding personal protective equipment requirements and more protective re-entry intervals for mushroom and greenhouse uses. The added restrictions on the use of the pest strip products provided, among other things, that large pest strips could no longer be used in homes except for garages, attics, crawl spaces, and sheds that are occupied for less than 4 hours per day. For a full description of the registrant's request, see the May 9, 2006 letter from AMVAC to EPA in the DDVP Special Review docket (EPA–HQ–OPP–2006–0396).

Subsequently, in early March, 2007, Amvac also requested the voluntary cancellation of all its pet collar and bait registrations and deletion of those uses from its technical label. Pursuant to section 6(f) of FIFRA, Amvac's requests to cancel the pet collar and bait registrations as well as deleting such uses from the technical label were published in the **Federal Register** on March 23, 2007. (72 FR 13786)(FRL–8120–7). On June 27, 2007, EPA granted Amvac's request and issued a final cancellation order for the pet collar and bait registrations. (72 FR 35235)(FRL–8127–5).

This proposal to conclude Special Review is based upon the label amendments requested by Amvac (as set forth in the May 9, 2006 letter) and EPA's determination that DDVP is eligible for reregistration as set forth in the June 30, 2006 IRED as well as the section 6(f) cancellations discussed above.

In sum, the Agency has determined that potential liver and cancer effects are no longer risks of concern, and based on the IRED and subsequent label changes that the cholinesterase inhibition issues have been adequately addressed through cancellations and other mitigation actions which limit exposure to DDVP. This Notice therefore proposes to terminate the DDVP Special Review based on the Agency's determination that all risks of concern identified in the PD 1 and earlier PD 2/3 have been satisfactorily addressed. Again, termination of this Special Review does not prejudice the Agency's review of the petition to cancel DDVP registrations and revoke DDVP tolerances, which will proceed separately and, if the Agency were to agree with the petition in whole or in part, could result in changes to the terms and conditions of DDVP registrations. For a complete description of the toxicity endpoints and risk assessment, see the DDVP Revised Human Health Risk Assessment, dated June 22, 2006, available in the DDVP reregistration docket (EPA–HQ–OPP–2002–0302) at *http://www.regulations.gov.*

III. Evaluation of Comments to PD 1

See section III.G of the September 1995 PD 2/3 for the evaluation of public comments received on the PD 1. This document is available in the DDVP Special Review docket (EPA–HQ–OPP–2006–0396) at the OPP Regulatory Public Docket (7508P), Environmental Protection Agency, Rm. S–4400, One Potomac Yard (South Bldg), 2777 S. Crystal Dr., Arlington, VA.

List of Subjects

Environmental protection, Pesticides and pests.

Dated: September 19, 2007.

Peter Caulkins,

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

[FR Doc. E7–18861 Filed 9–25–07; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2007–0986; FRL–8144–8]

The Allethrins 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 allethrin series of pesticides (bioallethrin, esbiol, esbiothrin, and pynamin forte). The Agency's risk assessments and other related documents also are available in the allethrins docket. The allethrins are synthetic pyrethroids used as insecticides on both indoor (residential and commercial) and outdoor (residential, commercial, and recreational) use sites. EPA has reviewed the allethrins 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: Molly Clayton, Special Review and Reregistration Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 603–0522; fax number: (703) 308–7070; e-mail address: *clayton.molly@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–2007–0986. 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. 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 RED for the allethrins series of pesticides under section 4(g)(2)(A) of FIFRA. The allethrins are synthetic pyrethroids used as insecticides on both indoor (residential and commercial) and outdoor (residential, commercial, and recreational) use sites. EPA has determined that the database to support reregistration is substantially complete and that products containing the allethrins are eligible for reregistration, provided the risks are mitigated in the manner described in the RED. Upon submission of any required product specific data under section 4(g)(2)(B) of FIFRA 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) of FIFRA for products containing the allethrins.

Although the allethrins RED was signed on June 30, 2007, certain components of the document, which did not affect the final regulatory decision, were undergoing final editing at that time. These components, including the table of contents and the summary of labeling changes, have been added to the allethrins RED document. None of these additions or changes alter the conclusions documented in the June 30, 2007 allethrins RED.

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 their uses, risk, and other factors, the allethrins were 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 the allethrins.

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. Few comments were received during the earlier comment periods for this reregistration case, and the issues related to this pesticide were resolved through consultations with stakeholders. The Agency therefore is issuing the allethrins 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.''

List of Subjects

Environmental protection, Pesticides and pests.

Dated: August 21, 2007.

Margaret J. Rice,

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

[FR Doc. E7–18789 Filed 9–25–07; 8:45 am]

BILLING CODE 6560–50–S

EXPORT-IMPORT BANK OF THE UNITED STATES

Sunshine Act Meeting

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

TIME AND PLACE: Thursday, September 27, 2007 at 9:30 a.m. The meeting will be held at Ex-Im Bank in Room 1143, 811 Vermont Avenue, NW., Washington, DC 20571.

OPEN AGENDA ITEM: PEFCO Secured Note Issues Resolutions.

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

FOR FURTHER INFORMATION CONTACT: Office of the Secretary, 811 Vermont Avenue, NW., Washington, DC 20571 (Telephone 202–565–3957).

Howard A. Schweitzer,

General Counsel.

[FR Doc. 07–4758 Filed 9–24–07; 12:00 pm]

BILLING CODE 6690–01–M

FEDERAL COMMUNICATIONS COMMISSION

Notice of Public Information Collection(s) Being Reviewed by the Federal Communications Commission, Comments Requested

September 19, 2007.

SUMMARY: The Federal Communications Commission, as part of its continuing effort to reduce paperwork burden invites the general public and other Federal agencies to take this opportunity to comment on the following information collection(s), as required by the Paperwork Reduction Act (PRA) of 1995, Public Law 104–13. An agency may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the Paperwork Reduction Act (PRA) that does not display a valid control number. Comments are requested concerning (a) 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; (b) the accuracy of the Commission's burden estimate; (c) ways to enhance the quality, utility, and clarity of the information collected; and (d) 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.

DATES: Written PRA comments should be submitted on or before November 26, 2007. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: You may submit all PRA comments by e-mail or U.S. mail. To submit your comments by e-mail, send them to *PRA@fcc.gov.* To submit your comments by U.S. mail, send them to Jerry Cowden, Federal Communications Commission, Room 1–B135, 445 12th Street, SW., Washington, DC 20554.

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection(s), contact Jerry