OMB Control Number 2070-0029; EPA ICR Number 0155.09

ICR ATTACHMENT D

"Guidance for Antimicrobial Pesticide Products With Anthrax-Related Claims"

Federal Register Notice of Availability (72 FR 31325)
and Draft Pesticide Registration (PR) Notice 2007-X

NOTE: The public comment period for the draft PR Notice closed on September 4, 2007. EPA is no longer accepting public comments on that document.

symptoms in patients with either spinal cord injury or multiple sclerosis. 4-aminopyridine is an active ingredient used in bird repellents that is currently undergoing reregistration.

• Extensive background materials concerning research to quantify the level of exposure received by people who mix, load, and apply pesticides. These materials, which were prepared by the Agricultural Handlers Exposure Task Force and by the Antimicrobial Exposure Assessment Task Force, generally explain the scope of the research programs being proposed by the Task Forces and describe the general scientific framework for conducting the research. In addition, each Task Force has provided Standard Operating Procedures which will guide the conduct of the studies.

The Board may also be reviewing draft HSRB reports for subsequent Board approval. Finally, the Board may also discuss planning for future HSRB meetings.

b. *Meeting Minutes and Reports.* Minutes of the meeting, summarizing the matters discussed and recommendations, if any, made by the advisory committee regarding such matters will be released within 90 calendar days of the meeting. Such minutes will be available at *http://www.epa.gov/osa/hsrb/* and *http://www.regulations.gov.* In addition, information concerning a Board meeting report, if applicable, can be found at *http://www.epa.gov/osa/hsrb/* or from the person listed under **FOR FURTHER INFORMATION CONTACT**.

Dated: May 31, 2007.

Kevin Teichman,

Acting EPA Science Advisor.

[FR Doc. E7–10859 Filed 6–5–07; 8:45 am]

BILLING CODE 6560–50–P

ENVIRONMENTAL PROTECTION AGENCY

[EPA–HQ–OPP–2006–1004; FRL–8113–5]

Pesticides; Draft Guidance for Pesticide Registrants on Antimicrobial Pesticide Products With Anthrax-Related Claims

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice of availability.

SUMMARY: The Agency is announcing the availability of, and seeking public comment on, a draft Pesticide Registration Notice entitled, ''Guidance for Antimicrobial Pesticide Products With Anthrax-Related Claims.'' PR notices are issued by the Office of Pesticide Programs (OPP) to inform pesticide registrants and other interested persons about important policies, procedures, and registration related decisions. This particular notice would, once final, provide guidance to prospective applicants of antimicrobial products that make labeling claims to inactivate Bacillus anthracis (anthrax) spores (hereafter referred to as ''anthrax-related products'').

DATES: Comments must be received on or before September 4, 2007.

ADDRESSES: Submit your comments, identified by docket identification (ID) number EPA–HQ–OPP–2006–1004, 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 Bldg.), 2777 S. Crystal Dr., 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 Facility telephone number is (703) 305–5805.

Instructions: Direct your comments to docket ID number EPA–HQ–OPP–2006–1004. 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 consider to be CBI or otherwise protected through regulations.gov or e-mail. The Federal regulations.gov website is an ''anonymous access'' system, which means EPA will not know your identity or contact information unless you provide it in the body of your comment. If you send an e-mail comment directly to EPA without going through regulations.gov, your e-mail address will be automatically captured and included as part of the comment that is placed in the docket and made available on the Internet. If you submit an electronic comment, EPA recommends that you include your name and other contact information. If EPA cannot read your comment due to technical difficulties and cannot contact you for clarification, EPA may not be able to consider your comment. Electronic files should avoid the use of special characters, any form of encryption, and be free of any defects or viruses.

Docket: All documents in the docket are listed in the docket index. Although listed in the index, some information is not publicly available, e.g., CBI or other information whose disclosure is restricted by statute. Certain other material, such as copyrighted material, is not placed on the Internet and will be publicly available only in hard copy form. 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 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.

FOR FURTHER INFORMATION CONTACT: Jeff Kempter, Antimicrobials Division, Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460–0001; telephone number: (703) 305–5448; fax number: (703) 308–6467; e-mail address: *kempter.carlton@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 those persons who are required to register pesticides and federal, state, and local government agencies and private institutions or organizations who are interested in bio-decontamination chemicals. Since other entities may also 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 information in this notice, consult the person listed under **FOR FURTHER INFORMATION CONTACT**.

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 or e-mail. 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.

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–OPPT–2006–1004. 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 Guidance Does this PR Notice Provide?

This draft PR Notice provides guidance to the registrant concerning antimicrobial products that make labeling claims to ''inactivate Bacillus anthracis (anthrax) spores'' (hereafter referred to as ''anthrax-related products''). In summary, this notice specifies that in order for a product to qualify for a claim of inactivating anthrax spores, an anthrax-related product should be:

1. Supported by specific sporicidal efficacy studies that are acceptable to EPA; and

2. Subject to specific terms and conditions of registration that limit the use of these products to certain trained persons. Prospective applicants are encouraged to follow the guidance in this notice and consult with EPA prior to applying for registration or amendment of a product when seeking such a claim. This guidance should help the United States be better prepared to respond to the intentional, accidental or natural introduction of anthrax spores by helping to assure that anthrax-related products bear appropriate labeling and are effective when used as directed.

In October 2001, when several letters containing Bacillus anthracis (anthrax) spores were introduced into the U.S. Postal Service mail system causing extensive contamination to dozens of buildings, no antimicrobial products were specifically registered for inactivating this particular pathogen. Since that time, the EPA has conducted extensive research and coordinated across the federal government to determine which efficacy test methods would be appropriate for demonstrating the effectiveness of antimicrobial products for inactivating B. anthracis spores. Guidance on acceptable efficacy test methods will be made available in a separate document. EPA's Office of Pesticide Programs has also developed guidance on the terms and conditions of registration for the labeling of these products. EPA intends to limit the use of these products to certain groups of trained persons. This notice is aimed primarily at applicants and registrants, but may also be of interest to other federal, state, and local government agencies, academic institutions, and other interested parties.

III. Do PR Notices Contain Binding Requirements?

The PR Notice discussed in this notice is intended to provide guidance to EPA personnel and to pesticide registrants. While the requirements in the statutes and Agency regulations are binding on EPA and the applicants, this PR Notice is not binding on pesticide registrants, and EPA may depart from the guidance where circumstances warrant and without prior notice. Likewise, pesticide registrants may assert that the guidance is not appropriate generally or not applicable to a specific pesticide or situation.

List of Subjects

Environmental protection, Administrative practice and procedure, Agricultural commodities, Pesticides and pests, Decontamination.

Dated: May 21, 2007.

Debra Edwards,

Director, Office of Pesticide Programs.

[FR Doc. E7–10694 Filed 6–5–07; 8:45 am]

BILLING CODE 6560–50–S

OFFICE OF SCIENCE AND TECHNOLOGY POLICY

Meeting of the President's Council of Advisors on Science and Technology

ACTION: Notice of meeting.

SUMMARY: This notice sets forth the schedule and summary agenda for a meeting of the President's Council of Advisors on Science and Technology (PCAST), and describes the functions of the Council. Notice of this meeting is required under the Federal Advisory Committee Act (FACA).

Dates and Place: June 25, 2007, Arlington, VA. The meeting will be held in Room 1235 of the National Science Foundation at 4201 Wilson Boulevard, Arlington, Virginia 22230.

Note that due to security requirements at the National Science Foundation, anyone planning to attend must pre-register no later than close of business on Thursday, June 21, 2007 by going to the PCAST Web site at: *http://www.ostp.gov/PCAST/pcast.html* or by calling 703–536–4996.

Type of Meeting: Open. Further details on the meeting agenda will be posted on the PCAST web site given above.

Proposed Schedule and Agenda: The President's Council of Advisors on Science and Technology (PCAST) is scheduled to meet in open session on Monday, June 25, 2007, at approximately 9 a.m. The PCAST subcommittee on nanotechnology has convened a group of experts from academia, industry, and non-governmental organizations to provide an overview of nanotechnology applications and implications. The

PESTICIDE REGISTRATION (PR) NOTICE 2007-X [DRAFT]

NOTICE TO MANUFACTURERS, PRODUCERS, FORMULATORS AND REGISTRANTS OF PESTICIDES

ATTENTION: **Persons Responsible for Registration of Pesticide Products**

SUBJECT: **Guidance for Antimicrobial Pesticide Products With Anthrax-Related Claims**

This notice provides guidance to prospective applicants of antimicrobial products that make labeling claims to "inactivate *Bacillus anthracis* (anthrax) spores" (hereafter referred to as "anthrax-related products"). This guidance should help the United States be prepared to respond to incidents involving anthrax spores by assuring that anthrax-related products are made available as registered products, that they bear appropriate labeling, and that they are effective when used as directed. In addition, this guidance will help protect public health from the potential risks of anthrax spores by limiting the use of anthrax-related products to those who are properly trained and qualified in their use and who can safely and effectively remediate contaminated facilities and their contents.

In summary, this notice specifies that in order for a product to qualify for a claim of inactivating anthrax spores, an anthrax-related product should be: (a) supported by specific sporicidal efficacy studies that are acceptable to EPA; and (b) subject to specific terms and conditions of registration that limit the use of these products to certain trained persons. Prospective applicants are encouraged to follow the guidance in this notice and consult with EPA prior to applying for registration or amendment of a product when seeking such a claim.

I. Why This Notice is Being Issued

In October 2001, when several letters containing *Bacillus anthracis* (anthrax) spores were introduced into the U.S. Postal Service mail system, no antimicrobial products were specifically registered for inactivating this particular pathogen. Since that time, EPA's Office of Pesticide Programs (EPA/OPP) has developed guidance on the terms and conditions of registration for the labeling of these products, which will limit their use to certain groups of trained persons, as described below. In addition, the EPA/OPP has developed guidance on efficacy test methods for demonstrating the effectiveness of antimicrobial products for inactivating *B. anthracis* spores on inanimate surfaces. Such efficacy testing guidance will be issued separately from this PR Notice, as described below. It is EPA's intent that registrants will follow these guidance documents when applying for registration of anthrax-related products, and that such products, when registered, will be made available to trained civilian and military personnel responsible for preparing for and responding to incidents involving anthrax spores.

II. Registration Guidance for Anthrax-related Products

A. Efficacy Test Methods

Currently, no EPA guidance is available concerning test methods to demonstrate the efficacy of anthrax-related products. EPA/OPP intends to add this guidance to Section 810.2100 of OPPTS' Pesticide Assessment Guidelines (PAG), which currently describes efficacy test methods that are recommended to demonstrate the efficacy of sterilants and sporicides. EPA/OPP also plans to present the draft efficacy test method guidance to the FIFRA Scientific Advisory Panel (SAP) for review and comment, and then issue the guidance in final form at about the same time that this PR Notice is issued. Until EPA issues such efficacy test method guidance, registrants are encouraged to submit their proposed efficacy test protocols to EPA for review prior to conducting such testing. If an application for registration is received with efficacy data to support a claim to inactivate *B. anthracis* spores in the absence of efficacy test method guidance, EPA will evaluate such data on a case-by-case basis.

B. Terms and Conditions of Registration

Due to the high potential risk to human health posed by *B. anthracis* spores (especially spores processed in a manner designed to enhance the potential for inhalation exposure), the Agency intends to establish terms and conditions of registration that limit the use of anthrax-related products to persons who are trained in their use. These limitations will help assure that the registered antimicrobial products are used safely and effectively. To implement such limitations, the registrant should agree in writing to specific Terms and Conditions of Registration that include items 1-5 below. Once the registrant has agreed to the terms and conditions, and once the Agency has issued a registration, the registrant would need to meet the terms and conditions. The consequences of not meeting the terms and conditions are described in section II.B.5. below. Further, the terms and conditions will be inserted into the Notice of Registration and will apply to all uses on the product's labeling. Accordingly, EPA strongly recommends a separate product registration for the uses that fall within the scope of this PR Notice.

1. Sale and Distribution Limitations

The registrant commits not to sell or distribute the product except to:

- Federal On-Scene Coordinators (FOSC), and contractors and other trained federal/state/local response personnel under the FOSC's supervision;
- Trained U.S. Military personnel and contractors under their supervision;
- Persons who, within the preceding 24 months, have been trained and determined to be competent by the registrant (or its contractor) in each of the topics described under item 2 below.

2. Training

Where a registrant proposes to sell or distribute anthrax-related products to persons other than FOSCs, trained U.S. Military personnel, and contractors and other response personnel under

the supervision of FOSCs or trained U.S. Military personnel, the registrant commits to providing training for such persons and to determining their competency. The registrant may itself perform the training or enter into a contract with a qualified third party to conduct the training on the registrant's behalf. The registrant further commits that the training curriculum will provide for refresher training at least every two years and, at a minimum, include instruction concerning:

- Characteristics of and human health hazards posed by *B. anthracis* spores;
- Personal Protective Equipment (PPE) appropriate for protection against both *B. anthracis* spores and the use of the pesticide product itself;
- Detailed instructions for safe and effective use of the pesticide product and any associated equipment;
- Detailed review of all steps involved in the decontamination process as provided in guidance from federal agencies (e.g., National Response Team Technical Assistance for Anthrax Response, Interim Final Draft, July 2005) as well as review of applicable federal statutory and regulatory requirements and guidance; and
- An assessment of the trained applicator's competency on the above issues through a written exam to be developed by the registrant.

EPA intends to review and approve any training program proposed in an application for registration or amended registration before determining the product's eligibility for registration.

3. Records and Reporting

The registrant commits to the following, commencing upon the effective date of registration or, if the anthrax-related claim is added to an existing registration by amendment, upon the effective date of that amendment:

- Maintain accurate, up-to-date records of its required activities under these Terms and Conditions, including:
  --Names of trainers, a listing of their qualifications, and copies of all training materials used.
  --Names and addresses of persons who have been trained and the dates and locations of such training.
  --Names and addresses of persons to whom the pesticide has been sold or distributed as well as the date and location of such sale or distribution and the product and quantity sold or distributed.
- Agree to maintain such records for three years, and to provide copies upon request of any authorized employee of the EPA, or of any State or political subdivision, duly designated by the Administrator.
- Agree that records and reports made or maintained in connection with these Terms and Conditions will not be claimed as confidential business information or trade secrets.

4. Labeling and Sale/Distribution/Use Limitations

The labeling of the affected anthrax-related product will include both a primary label on the container of the product and supplemental labeling that is similar to a technical manual. All primary and supplemental product labeling will be submitted to and reviewed by EPA as part of

the application for registration.

- The primary label and the technical manual will bear the following statements on the front panel:

  a. Directly above the product name, within a black outline, and in a font size no smaller than 12 points:

> ***For use only by:***
> - Federal On-Scene Coordinators and contractors and other trained federal/state/local response personnel under the FOSC's supervision;
> - Trained U.S. Military personnel and contractors under their supervision;
> - Persons who, within the preceding 24 months, have been trained and determined to be competent by the registrant (or its contractor) following completion of the required training.
>
> ***Under the terms and conditions of this product's registration, this product may only be sold or distributed by the registrant directly to the persons identified above.***

  b. Directly under the Directions for Use heading, in a font size no smaller than the heading, a statement that refers to the supplemental labeling, such as: **"See the accompanying technical manual for this product for complete use directions and safety precautions for inactivating *Bacillus anthracis* spores."** The technical manual is supplemental labeling under FIFRA and must comply with all pesticide product labeling requirements and bear the same boxed statement as specified above.

5. Consequence of Non-Compliance with The Terms and Conditions of Registration

The registrant agrees that failure to comply with any terms and conditions of registration specified above, may, at the sole discretion of EPA, result in the issuance of an order canceling the affected registration(s) without a hearing. Before issuing any such order, the Agency will notify the registrant in writing its intention to cancel the registration(s) and specify in such notification the basis for its conclusion that the registrant has failed to comply with the terms and conditions of registration. EPA will allow the registrant ten business days from the receipt of such notification to submit in writing a request to meet with the Director of the Office of Pesticide Programs ("Office Director") before a cancellation order is issued. The Agency will not issue a cancellation order before providing the registrant an opportunity to meet with the Office Director to discuss whether cancellation is appropriate. The registrant agrees that the decision of the Office Director will be final.

III. Implementation

To amend the registration of a currently registered product, the registrant will need to submit: (a) an Application for Registration form (EPA Form 8570-1) marked "Amendment" (for currently registered sterilants/sporicides), (b) appropriate efficacy data in proper format, (c) three copies of the revised labeling with changes clearly circled, and (d) a signed copy of the

agreement to the Terms and Conditions of Registration described in section II.B. to be effective in the event the registration is approved.

For new products, the registrant will submit: (a) an Application for Registration form marked "Registration" (EPA Form 8570-1), (b) appropriate efficacy data, (c) three copies of the draft labeling, (d) other forms and data required for a new product, and (e) a signed copy of the agreement to Terms and Conditions of registration described in section II.B. to be effective in the event the amendment is granted.

Registrants should note that the Terms and Conditions of registration will be inserted into the Notice of Registration and will apply to all uses on the product's labeling. Accordingly, EPA strongly recommends a separate product registration for the uses that fall within the scope of this PR Notice.

All applications for registration are subject to the Pesticide Registration Improvement Act of 2003 with regard to fees charged for applications for registration of pesticide products. Further information can be found at http://www.epa.gov/pesticides/fees/.

Submissions via the U.S. Postal Service: Use the official mailing address below for all submissions directed to the OPP regulatory divisions by mail:

Document Processing Desk (AMEND or REG)
Office of Pesticide Programs (7504PY)
U. S. Environmental Protection Agency
Ariel Rios Building
1200 Pennsylvania Avenue, N.W.
Washington, DC 20460

PLEASE NOTE: Do not address mail to be sent through the U.S. Postal Service (USPS) to the Arlington, Virginia address below. USPS will return it to you causing delay in processing your actions. There is no U.S. Postal Service delivery at the Virginia address.

Submissions via Personal/Courier Delivery: Deliveries in person or by a commercial courier for the regulatory divisions will be accepted at OPP's Document Processing Desk (7504C). Couriers and delivery personnel must present a valid picture identification card to gain access to the building. Hours of operation for the Document Processing Desk are 8:00 A.M. to 4:30 P.M., Monday through Friday, excluding Federal holidays. Personal and courier deliveries should be directed to:

Document Processing Desk (AMEND or REG)
Office of Pesticide Programs (7504PY)
U. S. Environmental Protection Agency
2777 S. Crystal Drive
Arlington, VA 22202

IV. Scope of This Notice

This PR Notice provides general guidance to EPA and to pesticide registrants and applicants, and the public. This guidance is not binding on either EPA or any outside parties, and the EPA may depart from the guidance where circumstances warrant and without prior notice. In their submissions, registrants and applicants may propose alternatives to the recommendations described in this notice, and the Agency will assess them for appropriateness on a case-by-case basis and will respond in writing.

V. Paperwork Reduction Act Notice

The information collection activities associated with the activities described in this PR Notice are already approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act (PRA), 44 U.S.C. 3501 *et seq*. The corresponding Information Collection Request (ICR) document for the training of pesticide applicators has been assigned EPA ICR number 0155 and is approved OMB control number 2070-0029. The ICR document for the pesticide application process has been assigned EPA ICR number 0277, and is approved OMB control number 2070-0060. The total estimated respondent paperwork burden associated with the training and certification of a pesticide applicator is an annual average of 3.1 burden hours. The annual average reporting and recordkeeping burden for a registration applicant respondent are estimated to range from 14 hours to 646 hours, depending upon the type of activity. For "Type A" activities, which include new active ingredients and new uses, the estimated annual applicant burden average is 194 hours per application. For "Type B" activities, which include amendments and notifications, the estimated annual applicant burden average is 14 hours per application. The respondent burden estimate for "Type C" reduced risk products is an average of 646 hours per product.

Under the PRA, "burden" means the total time, effort, or financial resources expended by persons to generate, maintain, retain, or disclose or provide information to or for a Federal agency. For this collection, it is the time reading the regulations, planning the necessary data collection activities, conducting tests, analyzing data, generating reports and completing other required paperwork, and storing, filing, and maintaining the data.

Under the PRA, an agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. The OMB control numbers for EPA's regulations codified in Chapter 40 of the CFR, after appearing in the preamble of the final rule, are listed in 40 CFR part 9, are displayed either by publication in the Federal Register or by other appropriate means, such as on the related collection instrument or form, if applicable. The display of OMB control numbers in certain EPA regulations is consolidated in 40 CFR part 9. For the ICR activity contained in this PR Notice, EPA is displaying the applicable OMB control number in the PR Notice above, and the applicable OMB control number also appears on the pesticide application.

In addition to commenting on the substance of this PR Notice, EPA welcomes your comments on the information collected related activities and the provided burden estimates, as well as any suggested methods for minimizing respondent burden, including the use of automated collection techniques. A copy of the most recent version of EPA ICR # 0155 is available under Docket ID No. EPA-HQ-OPP-2003-0357, and a copy of the most recent version of EPA ICR #0277 is available under Docket ID No. EPA-HQ-OPP-2004-0419. Both dockets may be accessed at www.regulations.gov. Please submit any comments on these ICRs as part of

your comments on the PR Notice, using the docket established for the PR Notice and following the instructions for submitting comments that are provided in the Federal Register document accompanying the draft PR Notice.

VI. For Further Information

If you have questions or need further information about this notice, you may contact the Antimicrobials Division, (703) 308-6411.