II. Background

The USDA's APHIS is applying for an EUP for the use of GonaCon™ Immunocontraceptive Vaccine, containing the active ingredient GnRH, to investigate the efficacy of reproductive control in fallow deer (*Dama dama*) at Point Reyes National Seashore in Marin County, CA. There are 90,000 acres in the park, although the treated area will be much less than this. Total quantity of active ingredient to be used is two pounds (70 pounds of the formulated product). The proposed period of shipment/use is July-August 2007.

III. What Action is the Agency Taking?

Following the review of the USDA APHIS application and any comments and data received in response to this notice, EPA will decide whether to issue or deny the EUP request for this EUP program, and if issued, the conditions under which it is to be conducted. Any issuance of an EUP will be announced in the **Federal Register**.

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

The Agency's authority for taking this action is under FIFRA section 5.

List of Subjects

Environmental protection, Experimental use permits.

Dated: March 30, 2007.

Lois Rossi,

Director, Registration Division, Office of Pesticide Programs.

[FR Doc. E7–6850 Filed 4–10–07; 8:45 am]

BILLING CODE 6560–50–S

ENVIRONMENTAL PROTECTION AGENCY

[EPA-HQ-OPP-2007-0037; FRL-8121-4]

Pesticide Registration Review; New Dockets Opened for Review and Comment

AGENCY: Environmental Protection Agency (EPA).

ACTION: Notice.

SUMMARY: EPA has established registration review dockets for the pesticides listed in the table in Unit III.A. With this document, EPA is opening the public comment period for these registration reviews. Registration review is EPA's periodic review of pesticide registrations to ensure that each pesticide continues to satisfy the statutory standard for registration, that is, the pesticide can perform its intended function without unreasonable adverse effects on human health or the environment. Registration review dockets contain information that will assist the public in understanding the types of information and issues that the Agency may consider during the course of registration reviews. Through this program, EPA is ensuring that each pesticide's registration is based on current scientific and other knowledge, including its effects on human health and the environment.

DATES: Comments must be received on or before June 11, 2007.

ADDRESSES: Submit your comments identified by the docket identification (ID) number for the specific pesticide of interest provided in the table in Unit III.A., 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 the assigned docket ID number listed in the table in Unit III.A. for the pesticides you are commenting on. 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 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 in the body of your comment and with any disk or CD-ROM you submit. 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 available in regulations.gov. To access the electronic docket, go to *http://www.regulations.gov*, select ''Advanced Search,'' then ''Docket Search.'' Insert the docket ID number where indicated and select the ''Submit'' button. Follow the instructions on the regulations.gov website to view the docket index or access available documents. 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 electronically 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: For information about the pesticides included in this notice, contact the specific Chemical Review Managers for these pesticides as identified in the table in Unit III.A..

For general questions on the registration review program, contact Kennan Garvey, Special Review and Reregistration Division (7508P), Office of Pesticide Programs, Environmental Protection Agency, 1200 Pennsylvania Ave., NW., Washington, DC 20460-0001; telephone number: (703) 305-7106; fax number: (703) 308-8090; e-mail address: *garvey.kennan@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, farmworker, 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. 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.

II. Authority

EPA is initiating its reviews of the pesticides identified in this document pursuant to section 3(g) of the Federal Insecticide, Fungicide, and Rodenticide Act (FIFRA) and the Procedural Regulations for Registration Review published in the **Federal Register** issue of August 9, 2006, and effective on October 10, 2006 (71 FR 45719) (FRL-8080-4), you may also access this document on EPA's Internet at *http://www.epa.gov/fedrgstr/EPA-PEST/2006/August/Day-09/p12904.htm.* Section 3(g) of FIFRA provides, among other things, that the registrations of pesticides are to be periodically reviewed. The goal is a review of a pesticide's registration every 15 years. Under section 3(a) of FIFRA, a pesticide product may be registered or remain registered only if it meets the statutory standard for registration given in FIFRA section 3(c)(5). When used in accordance with widespread and commonly recognized practice, the pesticide product must perform its intended function without unreasonable adverse effects on the environment; that is, without any unreasonable risk to man or the environment, or a human dietary risk from residues that result from the use of a pesticide in or on food.

III. Registration Reviews

A. What Action is the Agency Taking?

As directed by FIFRA section 3(g), EPA is periodically reviewing pesticide registrations to assure that they continue to satisfy the FIFRA standard for registration—that is, they can still be used without unreasonable adverse effects on human health or the environment. The implementing regulations establishing the procedures for registration review appear at 40 CFR part 155. A pesticide's registration review begins when the Agency establishes a docket for the pesticide's registration review case and opens the docket for public review and comment. At present, EPA is opening registration review dockets for the cases identified in the following table.

TABLE—REGISTRATION REVIEW DOCKETS OPENING

Registration Review Case Name and Number	Pesticide Docket ID Number	Chemical Review Managers Name, Phone Number, E-mail Address
Case 6050 Trichoderma species	EPA-HQ-OPP-2006-0245	Shanaz Bacchus, Chemist/RAL; (703) 308-8097; bacchus.shanaz@epa.gov
Case 6058 Linalool	EPA-HQ-OPP-2006-0356	Stephen Morrill, Special Assistant; (703) 308-8319; morrill.stephen@epa.gov

B. Docket Content

1. *Review dockets.* The registration review dockets contain information that the Agency may consider in the course of the registration review. The Agency may include information from its files including, but not limited to, the following information:

- An overview of the registration review case status.
- A list of current product registrations and registrants.
- FR notices regarding any pending registration actions.
- FR notices regarding current or pending tolerances.
- Risk assessments.
- Bibliographies concerning current registrations.
- Summaries of incident data.
- Any other pertinent data or information.

Each docket contains a document summarizing what the Agency currently knows about the pesticide case and a preliminary work plan for anticipated data and assessment needs. Additional documents provide more detailed information. During this public comment period, the Agency is asking that interested persons identify any additional information they believe the Agency should consider during the registration reviews of these pesticides. The Agency identifies in each docket the areas where public comment is specifically requested, though comment in any area is welcome.

2. *Other related information.* More information on these cases, including the active ingredients for each case, may be located 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 may be seen at *http://www.epa.gov/oppsrrd1/registration_review*.

3. *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 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, Pesticides and pests.

Dated: March 30, 2007.

James Jones,

Acting Assistant Administrator, Office of Prevention, Pesticides and Toxic Substances.

[FR Doc. E7–6626 Filed 4–10–07; 8:45 am]

BILLING CODE 6560–50–S

EXPORT-IMPORT BANK OF THE U.S.

[Public Notice 98]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Export-Import Bank of the U.S.

ACTION: Notice and request for comments.

SUMMARY: The Export-Import Bank (''Ex-Im Bank'') is seeking approval of the proposed information collection described below. The collection comprises certain applications and forms relating to Ex-Im Bank's insurance program. As part of its continuing effort to reduce paperwork and respondent burden, Ex-Im Bank invites the general public and other Federal Agencies to comment on the proposed information collection as required by the Paperwork Reduction Act of 1995.

SUPPLEMENTARY INFORMATION: This notice is soliciting comments from the public concerning the proposed collection of information to: (1) Evaluate whether the proposed collection is necessary for the performance of the functions of the agency, including whether the information will have practical utility; (2) evaluate the accuracy of the agency's estimate of the burden of the proposed information collection; (3) enhance the quality, utility, and clarity of the information to be collected; and (4) minimize the burden of collection of information on those who are to respond, including through the use of appropriated automated collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses.

DATES: Written comments should be received on or before May 11, 2007 to be assured of consideration.

ADDRESSES: Direct all comments to David Rostker, Office of Management and Budget, Office of Information and Regulatory Affairs, NEOB, Room 10202, Washington, DC 20503, (202) 395–3897. Direct all requests for information, including copies of the proposed collection of information and supporting documentation to Solomon Bush, Office of Information and Records Management, Export-Import Bank of the United States, 811 Vermont Avenue, NW., Room 776, Washington, DC 20571, (202) 565–3353 or (800) 565–3946, x3353.

Titles and Form Numbers:

Application for Letter of Credit Insurance Policy, EIB 92–34;

Beneficiary Certificate and Agreement, EIB 92–37;

Short-Term Multi-Buyer Export Credit Insurance Policy Application, EIB 92–50;

Broker Registration Form, EIB 92–79.

OMB Number: 3048–0009.

Type of Review: Regular.

Need and Use: The information requested provides Ex-Im Bank with information necessary to determine legislatively required reasonable assurance of repayment and fulfills other statutory requirements. These forms are used in connection with Ex-Im Bank's insurance program.

Affected Public: The forms affect entities involved in Ex-Im Bank's programs supporting the export of U.S. goods and services, including exporters, banks, insurance brokers and non-profit or state and local governments acting as facilitators.

	EIB 92–34	EIB 92–37	EIB 92–50	EIB 92–79
Estimated Annual Respondents	10	10	398	50
Estimated Time Per Respondent	1 Hour	20 Minutes	1 Hour	2 Hours
Estimated Annual Burden	10 Hours	3.3 Hours	368 Hours	100 Hours
Frequency of Reporting or Use	Applications submitted one time.			