Notices

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This section of the FEDERAL REGISTER contains documents other than rules or proposed rules that are applicable to the public. Notices of hearings and investigations, committee meetings, agency decisions and rulings, delegations of authority, filing of petitions and applications and agency statements of organization and functions are examples of documents appearing in this section.

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. 03–033–1]

National Wildlife Services Advisory Committee; Meeting

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice of meeting.

SUMMARY: Pursuant to the Federal Advisory Committee Act, we are giving notice of a meeting of the National Wildlife Services Advisory Committee.

DATES: The meeting will be held on June 24, 2003, from 8 a.m. to 5 p.m. and June 25, 2003, from 8 a.m. to noon.

ADDRESSES: The meeting will be held at the USDA Center at Riverside, 4700 River Road, Riverdale, MD.

FOR FURTHER INFORMATION CONTACT: Mrs. Joanne Garrett, Director, Operational Support Staff, WS, APHIS, 4700 River Road, Unit 87, Riverdale, MD 20737–1234, (301) 734–7921.

SUPPLEMENTARY INFORMATION: The National Wildlife Services Advisory Committee (Committee) advises the Secretary of Agriculture concerning policies, program issues, and research needed to conduct the Wildlife Services (WS) program. The Committee also serves as a public forum enabling those affected by the WS program to have a voice in the program's policies.

The meeting will focus on operational and research activities and will be open to the public. Due to time constraints, the public will not be able to participate in the Committee's discussions. However, written statements concerning meeting topics may be filed with the Committee before or after the meeting by sending them to Mrs. Joanne Garrett at the address listed under **FOR FURTHER INFORMATION CONTACT,** or may be filed at the meeting. Please refer to Docket No. 03–033–1 when submitting your statements.

This notice of meeting is given pursuant to section 10 of the Federal Advisory Committee Act (5 U.S.C. App. II).

Done in Washington, DC, this 2nd day of April, 2003.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 03–8600 Filed 4–8–03; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. 03–035–1]

Determination of Regulatory Review Period for Purposes of Patent Extension; Poulvac® ST Vaccine

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice.

SUMMARY: We are advising the public that the Animal and Plant Health Inspection Service has determined the regulatory review period for Poulvac® ST Vaccine and is publishing this notice of that determination as required by law. We have made this determination in response to the submission of an application to the Commissioner of Patents and Trademarks, Department of Commerce, for the extension of a patent that claims that veterinary biologic.

DATES: We will consider all requests for revision of the regulatory review period determination that we receive on or before May 9, 2003. We will consider all due diligence petitions that we receive on or before October 6, 2003.

ADDRESSES: You may submit revision requests and due diligence petitions by postal mail/commercial delivery or by e-mail. If you use postal mail/commercial delivery, please send four copies of your request or petition (an original and three copies) to: Docket No. 03–035–1, Regulatory Analysis and Development, PPD, APHIS, Station 3C71, 4700 River Road Unit 118, Riverdale, MD 20737–1238. Please state that your request or petition refers to Docket No. 03–035–1. If you use e-mail, address your request or petition to *regulations@aphis.usda.gov.* Your request or petition must be contained in the body of your message; do not send attached files. Please include your name and address in your message and ''Docket No. 03–035–1'' on the subject line.

You may request a copy of the regulatory review period determination by writing to Dr. Patricia L. Foley, USDA, APHIS, VS, CVB–LPD, 510 South 17th Street, Suite 104, Ames, IA 50010–8197, or by calling (515) 232–5785. Please refer to the docket number, date, and complete title of this notice when requesting copies.

A copy of the regulatory review period determination and any revision requests or due diligence petitions that we receive on this determination are available for public inspection in our reading room. The reading room is located in room 1141 of the USDA South Building, 14th Street and Independence Avenue SW., Washington, DC. Normal reading room hours are 8 a.m. to 4:30 p.m., Monday through Friday, except holidays. To be sure someone is there to help you, please call (202) 690–2817 before coming.

APHIS documents published in the **Federal Register**, and related information, including the names of organizations and individuals who have commented on APHIS dockets, are available on the Internet at *http://www.aphis.usda.gov/ppd/rad/webrepor.html.*

FOR FURTHER INFORMATION CONTACT: Dr. Albert P. Morgan, Chief Staff Officer, Operational Support Section, Center for Veterinary Biologics, Licensing and Policy Development, VS, APHIS, 4700 River Road Unit 148, Riverdale, MD 20737–1231; phone (301) 734–8245; fax (301) 734–4314. For information regarding the regulatory review period determination, contact Dr. Patricia L. Foley, APHIS, VS, CVB–LPD, 510 South 17th Street, Suite 104, Ames, IA 50010–8197; phone (515) 232–5785.

SUPPLEMENTARY INFORMATION: The provisions of 35 U.S.C. 156, ''Extension of patent term,'' provide, generally, that a patent for a product may be extended for a period of up to 5 years as long as the patent claims a product that, among other things, was subject to a regulatory review period before its commercial marketing or use. (The term ''product'' is defined in that section as ''a drug product'' [which includes veterinary