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.

AGENCY FOR INTERNATIONAL DEVELOPMENT

Bureau for Democracy, Conflict and Humanitarian Assistance; Office of Food for Peace; Announcement of Draft Food for Peace Pub. L. 480 Title II Program Policies and Proposal Guidelines (FY 07)

Pursuant to the Agricultural Trade Development and Assistance Act of 1954 (Public Law 480, as amended), notice is hereby given that the Draft Food for Peace Pub. L. 480 Title II Program Policies and Proposal Guidelines (FY 07) are being made available to interested parties for the required thirty (30) day comment period.

Individuals who wish to receive a copy of these draft guidelines should contact: Office of Food for Peace, U.S. Agency for International Development, RRB 7.06–102, 1300 Pennsylvania Avenue, NW., Washington, DC 20523–7600. The draft guidelines may also be found at *http://www.usaid.gov/our_work/ humanitarian_assistance/ffp/fy07_myap.html.* Individuals who have questions or comments on the draft guidelines should contact Lisa Witte at the above address, at (202) 712–5162 or *lwitte@usaid.gov.* The thirty-day comment period will begin on the date that this announcement is published in the **Federal Register**.

Lisa Witte,

Acting Chief, Policy and Technical Division, Office of Food for Peace, Bureau for Democracy, Conflict and Humanitarian Assistance.

[FR Doc. 06–1933 Filed 3–1–06; 8:45 am]

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DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2006–0031]

Availability of an Environmental Assessment for Field Testing Marek's Disease-Newcastle Disease Vaccine, Serotypes 2 and 3, Live Virus, Live Marek's Disease Vector

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 prepared an environmental assessment concerning authorization to ship for the purpose of field testing, and then to field test, an unlicensed Marek's Disease-Newcastle Disease Vaccine, Serotypes 2 and 3, Live Virus, Live Marek's Disease Vector. The environmental assessment, which is based on a risk analysis prepared to assess the risks associated with the field testing of this vaccine, examines the potential effects that field testing this veterinary vaccine could have on the quality of the human environment. Based on the risk analysis, we have reached a preliminary determination that field testing this veterinary vaccine will not have a significant impact on the quality of the human environment, and that an environmental impact statement need not be prepared. We intend to authorize shipment of this vaccine for field testing following the close of the comment period for this notice unless new substantial issues bearing on the effects of this action are brought to our attention. We also intend to issue a U.S. Veterinary Biological Product license for this vaccine, provided the field test data support the conclusions of the environmental assessment and the issuance of a finding of no significant impact and the product meets all other requirements for licensing.

DATES: We will consider all comments that we receive on or before April 3, 2006.

ADDRESSES: You may submit comments by either of the following methods:

- Federal eRulemaking Portal: Go to *http://www.regulations.gov* and, in the ''Search for Open Regulations'' box, select ''Animal and Plant Health Inspection Service'' from the agency drop-down menu, then click on ''Submit.'' In the Docket ID column, select APHIS–2006–0031 to submit or view public comments and to view supporting and related materials available electronically. After the close of the comment period, the docket can be viewed using the ''Advanced Search'' function in Regulations.gov.
- Postal Mail/Commercial Delivery: Please send four copies of your comment (an original and three copies) to Docket No. APHIS–2006–0031, Regulatory Analysis and Development, PPD, APHIS, Station 3A–03.8, 4700 River Road Unit 118, Riverdale, MD 20737–1238. Please state that your comment refers to Docket No. APHIS–2006–0031.

Reading Room: You may read any comments that we receive on this docket 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.

Other Information: Additional information about APHIS and its programs is available on the Internet at *http://www.aphis.usda.gov.*

FOR FURTHER INFORMATION CONTACT: Dr. Albert P. Morgan, Section Leader, Operational Support Section, Center for Veterinary Biologics, Policy, Evaluation, and Licensing, VS, APHIS, 4700 River Road Unit 148, Riverdale, MD 20737–1231; (301) 734–8245.

For information regarding the environmental assessment or the risk analysis, or to request a copy of the environmental assessment (as well as the risk analysis with confidential business information removed), contact Dr. Patricia L. Foley, Risk Manager, Center for Veterinary Biologics, Policy, Evaluation, and Licensing VS, APHIS, 510 South 17th Street, Suite 104, Ames, IA 50010; phone (515) 232–5785, fax (515) 232–7120.

SUPPLEMENTARY INFORMATION: Under the Virus-Serum-Toxin Act (21 U.S.C. 151 *et seq.*), a veterinary biological product must be shown to be pure, safe, potent, and efficacious before a veterinary biological product license may be issued. A field test is generally necessary to satisfy prelicensing