burden estimates; (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: Submit comments on or before July 17, 2007.

FOR FURTHER INFORMATION CONTACT: Beverly Johnson, Bureau for Management, Office of Administrative Services. Information and Records Division, U.S. Agency for International Development, Room 2.07–106, RRB, Washington, DC 20523, (202) 712–1365 or via e-mail *bjohnson@usaid.gov.*

SUPPLEMENTARY INFORMATION:

OMB No.: OMB 0412–0506.

Form No.: N/A.

Title: USAID Small Business Resource Database (formerly known as the Vendor Database).

Type of Review: Renewal of information collection.

Purpose: This information collection is to assist U.S. small businesses, small disadvantaged businesses, women-owned small businesses, HUBZone small businesses, service-disabled Veteran-owned small businesses and Veteran-owned small businesses (hereinafter referred to as ''U.S. small businesses'') to participate in the U.S. Agency for International Development (USAID) in supporting long-term and equitable economic growth and advances U.S. foreign policy objectives by supporting sustainable development program activities.

Annual Reporting Burden:

Respondents: 1.100.

Total annual responses: 2.200.

Total annual hours requested: 550 hours.

Dated: May 11, 2007.

Joanne Paskar,

Chief, Information and Records Division, Office of Administrative Services, Bureau for Management.

[FR Doc. 07–2470 Filed 5–17–07; 8:45 am]

BILLING CODE 6116–01–M

DEPARTMENT OF AGRICULTURE

Agricultural Marketing Service

[Docket No. AMS–FV–07–0075, FV–07–376]

Fruit and Vegetable Industry Advisory Committee

AGENCY: Agricultural Marketing Service, USDA.

ACTION: Notice of public meeting.

SUMMARY: The purpose of this notice is to notify all interested parties that the Agricultural Marketing Service (AMS) will hold a Fruit and Vegetable Industry Advisory Committee (Committee) meeting that is open to the public. The U.S. Department of Agriculture (USDA) established the Committee to examine the full spectrum of issues faced by the fruit and vegetable industry and to provide suggestions and ideas to the Secretary of Agriculture on how USDA can tailor its programs to meet the fruit and vegetable industry's needs. This notice sets forth the schedule and location for the meeting.

DATES: Monday, June 4, 2007, from 8 a.m. to 5 p.m., and Tuesday, June 5, 2007, from 8 a.m. to 2 p.m.

ADDRESSES: The Committee meeting will be held at the Holiday Inn Central, 1501 Rhode Island Avenue NW, Washington, DC Conference Room: Mayors I and II.

FOR FURTHER INFORMATION CONTACT: Andrew Hatch, Designated Federal Official, USDA, AMS, Fruit and Vegetable Programs. Telephone: (202) 690–0182. Facsimile: (202) 720–0016. E-mail: *andrew.hatch@usda.gov.*

SUPPLEMENTARY INFORMATION: Pursuant to the Federal Advisory Committee Act (FACA) (5 U.S.C. App. II), the Secretary of Agriculture established the Committee in August 2001 to examine the full spectrum of issues faced by the fruit and vegetable industry and to provide suggestions and ideas to the Secretary on how USDA can tailor its programs to meet the fruit and vegetable industry's needs. The Committee was re-chartered in July 2003 and again in June 2005 with new members appointed by USDA from industry nominations.

AMS Deputy Administrator for Fruit and Vegetable Programs, Robert C. Keeney, serves as the Committee's Executive Secretary. Representatives from USDA mission areas and other government agencies affecting the fruit and vegetable industry are called upon to participate in the Committee's meetings as determined by the Committee Chairperson. AMS is giving notice of the Committee meeting to the public so that they may attend and present their recommendations. Reference the date and address section of this announcement for the time and place of the meeting.

Topics of discussion at the advisory committee meeting will include: updates on State audit requirements and marketing agreements, food safety initiatives, the Food Stamp program, and invasive pest and disease safety precautions.

Those parties that would like to speak at the meeting should register on or before May 28, 2007. To register as a speaker, please e-mail your name, affiliation, business address, e-mail address, and phone number to Mr. Andrew Hatch at: *andrew.hatch@usda.gov* or facsimile to (202) 720–0016. Speakers who have registered in advance will be given priority. Groups and individuals may submit comments for the Committee's consideration to the same e-mail address. The meeting will be recorded, and information about obtaining a transcript will be provided at the meeting.

The Secretary of Agriculture selected a diverse group of members representing a broad spectrum of persons interested in providing suggestions and ideas on how USDA can tailor its programs to meet the fruit and vegetable industry's needs. Equal opportunity practices were considered in all appointments to the Committee in accordance with USDA policies.

If you require special accommodations, such as a sign language interpreter, please use either contact name listed above.

Dated: May 15, 2007.

Lloyd Day,

Administrator, Agricultural Marketing Service.

[FR Doc. E7–9575 Filed 5–17–07; 8:45 am]

BILLING CODE 3410–02–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2007–0057]

Notice of Request for Extension of Approval of an Information Collection; Importation of Animals and Poultry, Animal and Poultry Products, Certain Animal Embryos, Semen, and Zoological Animals

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Extension of approval of an information collection; comment request.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, this notice announces the Animal and Plant Health Inspection Service's intention to request an extension of approval of an information collection associated with the importation of animals and poultry, animal and poultry products, and animal germplasm.

DATES: We will consider all comments that we receive on or before July 17, 2007.

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

• *Federal eRulemaking Portal:* Go to *http://www.regulations.gov,* select ''Animal and Plant Health Inspection Service'' from the agency drop-down menu, then click ''Submit.'' In the Docket ID column, select APHIS–2007–0057 to submit or view public comments and to view supporting and related materials available electronically. Information on using Regulations.gov, including instructions for accessing documents, submitting comments, and viewing the docket after the close of the comment period, is available through the site's ''User Tips'' link.

• *Postal Mail/Commercial Delivery:* Please send four copies of your comment (an original and three copies) to Docket No. APHIS–2007–0057, 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–2007–0057.

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: For information on an information collection associated with the importation of animals and poultry, animal and poultry products, and animal germplasm, contact Dr. James Davis, Senior Staff Veterinarian, National Center for Import and Export, VS, APHIS, 4700 River Road Unit 39, Riverdale, MD 20737–1231; (301) 734–0694. For copies of more detailed information on the information collection, contact Mrs. Celeste Sickles, APHIS' Information Collection Coordinator, at (301) 734–7477.

SUPPLEMENTARY INFORMATION: *Title:* Importation of Animals and Poultry, Animal and Poultry Products, Certain Animal Embryos, Semen, and Zoological Animals.

OMB Number: 0579–0040.

Type of Request: Extension of approval of an information collection.

Abstract: The Animal and Plant Health Inspection Service (APHIS) of the United States Department of Agriculture is authorized, among other things, to prohibit or restrict the importation of animals, animal products, and other articles into the United States to prevent the introduction of animal diseases and pests. In connection with this mission, APHIS collects pertinent information from persons who import animals or poultry, animal or poultry products, or animal germplasm into the United States.

This information includes data such as the origins of the animals or animal products to be imported, the health status of the animals or the processing methods used to produce animal products to be imported, and whether the animals or animal products were temporarily offloaded in another country during their transit to the United States. We need this information to help ensure that these imports do not introduce exotic animal diseases into the United States.

We use a variety of information collection procedures, devices, and forms including, but not limited to: Health certificates, import permits, ear tags, leg bands, specimen submission forms, inspection reports, cooperative and trust fund agreements, and certification statements.

We are asking the Office of Management and Budget (OMB) to approve our use of these information collection activities for an additional 3 years.

The purpose of this notice is to solicit comments from the public (as well as affected agencies) concerning our information collection. These comments will help us:

(1) Evaluate whether the collection of information is necessary for the proper performance of the functions of the Agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of our estimate of the burden of the information collection, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the information collection on those who are to respond, through use, as appropriate, of automated, electronic, mechanical, and other collection technologies, e.g., permitting electronic submission of responses.

Estimate of burden: The public reporting burden for this collection of information is estimated to average 0.41688779 hours per response.

Respondents: Entities that import animals or poultry, animal or poultry products, and animal embryos, germplasm, and semen.

Estimated annual number of respondents: 77,976.

Estimated annual number of responses per respondent: 162.4330306.

Estimated annual number of responses: 12,665,878.

Estimated total annual burden on respondents: 528,025 hours. (Due to averaging, the total annual burden hours may not equal the product of the annual number of responses multiplied by the reporting burden per response.)

All responses to this notice will be summarized and included in the request for OMB approval. All comments will also become a matter of public record.

Done in Washington, DC, this 11th day of May 2007.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. E7–9551 Filed 5–17–07; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2007–0056]

Draft Guidelines on Pharmacovigilance of Veterinary Medicinal Products: Controlled Lists of Terms

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Notice of availability and request for comments.

SUMMARY: The International Cooperation on Harmonization of Technical Requirements for the Registration of Veterinary Medicinal Products (VICH) has developed a draft guideline titled ''Pharmacovigilance of Veterinary Medicinal Products: Controlled Lists of Terms.'' This draft guideline provides guidance for the development and maintenance of the controlled lists of terms required to complete the controlled data fields contained in adverse event reports concerning the use of veterinary medicinal products. Because the draft guideline applies to pharmacovigilance and adverse event reporting on veterinary vaccines regulated by the Animal and Plant Health Inspection Service under the Virus-Serum-Toxin Act, we are requesting comments on the scope of the guideline and its provisions so that we may include any relevant public input on the draft in the Agency's