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 the Virus-Serum-Toxin Act and regulations, contact Dr. Albert Morgan, Section Leader, Operational Support Staff, Center for Veterinary Biologics, VS, APHIS, 4700 River Road Unit 148, Riverdale MD 20737, (301) 734–8245. 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: Virus-Serum-Toxin Act and Regulations.

OMB Number: 0579–0013.

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

Abstract: The Animal and Plant Health Inspection Service (APHIS) of the U.S. Department of Agriculture is responsible for ensuring that veterinary biological products are pure, safe, potent, and effective. This program is conducted under the Virus-Serum-Toxin Act (21 U.S.C. 151 *et seq.*) and the regulations in 9 CFR, chapter I, subchapter E. Veterinary biological products are defined as all viruses, serums, toxins (excluding substances that are selectively toxic to microorganisms, e.g., antibiotics), or analogous products at any stage of production, shipment, distribution, or sale, which are intended for use in the treatment of animals and which act primarily through the direct stimulation, supplementation, enhancement, or modulation of the immune system or immune response. The term ''biological products'' includes, but is not limited to, vaccines, bacterins, allergens, antibodies, antitoxins, toxoids, immunostimulants, certain cytokines, antigenic or immunizing components of live organisms, and diagnostic components that are of natural or synthetic origin or that are derived from synthesizing or altering various substances or components of substances, such as microorganisms, genes or genetic sequences, carbohydrates, proteins, antigens, allergens, or antibodies.

To accomplish its mission, APHIS issues licenses to qualified establishments that produce biological products and issues permits to importers of such products. We also enforce requirements concerning production, packaging, labeling, and shipping of these products and set standards for the testing of these products.

Fulfilling this responsibility requires us to use certain information collection activities such as establishment license applications, product license applications, product import permit applications, product and test report forms, and field study summaries. This information helps us to ensure that biological products used in the United States are pure, safe, potent, and effective. If we did not collect this information, we would be unable to carry out this mission.

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 these information collection activities. 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 4.337559606 hours per response.

Respondents: U.S. importers, exporters, and shippers of veterinary biological products; State veterinary authorities; and operators of establishments that produce or test veterinary biological products or that engage in product research and development.

Estimated Annual Number of Respondents: 500.

Estimated Annual Number of Responses per Respondent: 43.452.

Estimated Annual Number of Responses: 21,726.

Estimated Total Annual Burden on Respondents: 94,237.82 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 27th day of August 2007.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. E7–14988 Filed 8–1–07; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2007–0076]

Notice of Request for Extension of Approval of an Information Collection; Importation of Tomatoes From Spain, Chile, France, Morocco, and Western Sahara

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 our regulations governing the importation of tomatoes from Spain, Chile, France, Morocco, and Western Sahara.

DATES: We will consider all comments that we receive on or before October 1, 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–2007–0076 to submit or view public comments and to view supporting and related materials available electronically. Information on using *http://www.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–2007–0076, 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–2007–0076.

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 regarding foreign quarantine regulations, contact Ms. Sharon Porsche, Import Specialist, Commodity Import Analysis and Operations, PPQ, APHIS, 4700 River Road Unit 133, Riverdale, MD 20737–1236; (301) 734–5281. 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 Tomatoes from Spain, Chile, France, Morocco, and Western Sahara.

OMB Number: 0579–0131.

Type of Request: Extension of Approval of an Information Collection.

Abstract: The Plant Protection Act (PPA, 7 U.S.C. 7701 *et seq.*) authorizes the Secretary of Agriculture to restrict the importation, entry, or interstate movement of plants, plant products, and other articles to prevent the introduction of plant pests, including fruit flies, into the United States or their dissemination within the United States. Regulations authorized by the PPA concerning the importation of fruits and vegetables into the United States from certain parts of the world are contained in ''Subpart—Fruits and Vegetables'' (7 CFR 319.56 through 319.56–46).

The regulations in 319.56–28 allow tomatoes from Spain, Chile, France, Morocco, and Western Sahara to be imported into the United States subject to certain conditions designed to protect the tomatoes from infestation by the Mediterranean fruit fly (Medfly). Allowing tomatoes to be imported necessitates the use of certain information collection activities, including completing phytosanitary inspection certificates and maintaining records regarding trap placement and Medfly captures. The information we collect serves as the supporting documentation needed to confirm that the tomatoes meet the conditions set forth in the regulations.

We are asking the Office of Management and Budget (OMB) to approve our use of this information collection activity 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 collection of information, 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 collection of information 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 0.6960 hour per response.

Respondents: Importers, foreign officials, shippers.

Estimated Annual Number of Respondents: 34.

Estimated Annual Number of Responses Per Respondent: 72.

Estimated Annual Number of Responses: 2,448.

Estimated Total Annual Burden on Respondents: 1,704 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 27th day of July 2007.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. E7–15008 Filed 8–1–07; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2007–0077]

Notice of Request for Extension of Approval of an Information Collection; Black Stem Rust; Identification Requirements for Addition of Rust-Resistant Varieties

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 black stem rust quarantine and regulations.

DATES: We will consider all comments that we receive on or before October 1, 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–2007–0077 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–2007–0077, 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–2007–0077.

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,