SUPPLEMENTARY INFORMATION:

Title: Importation of Baby Squash and Baby Courgettes From Zambia.

OMB Number: 0579–0347.

Type of Request: Extension of an 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 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 319.56–1 through 319.56–50).

Under these regulations, baby squash and baby courgettes from Zambia are subject to certain conditions before entering the United States to ensure that exotic plant pests are not introduced into the United States. Allowing baby squash and baby courgettes to be imported necessitates the use of certain information collection activities, including completing phytosanitary inspection certificates, maintaining inspection records, and labeling cartons.

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 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 1 hour per response.

Respondents: National plant protection organization of Zambia.

Estimated annual number of respondents: 2.

Estimated annual number of responses per respondent: 2.

Estimated annual number of responses: 4.

Estimated total annual burden on respondents: 4 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 20th day of June 2011.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2011–15867 Filed 6–23–11; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2011–0056]

Notice of Revision and Request for Extension of Approval of an Information Collection; Pale Cyst Nematode

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Revision and 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 regulations for the interstate movement of regulated articles to prevent the spread of the pale cyst nematode to noninfested areas of the United States.

DATES: We will consider all comments that we receive on or before August 23, 2011.

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

- *Federal eRulemaking Portal:* Go to *http://www.regulations.gov/#!documentDetail;D=APHIS-2011-0056-0001.*
- *Postal Mail/Commercial Delivery:* Send your comment to Docket No. APHIS–2011–0056, Regulatory Analysis and Development, PPD, APHIS, Station 3A–03.8, 4700 River Road, Unit 118, Riverdale, MD 20737–1238.

Supporting documents and any comments we receive on this docket may be viewed at *http://www.regulations.gov/#!docketDetail;D=APHIS-2011-0056* or in our reading room, which 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) 6902817 before coming.

FOR FURTHER INFORMATION CONTACT: For information on the interstate movement of regulated articles to prevent the spread of pale cyst nematode, contact Mr. Jonathan Jones, Program Manager, Emergency and Domestic Programs, PPQ, APHIS, 4700 River Road, Unit 160, Riverdale, MD 20737; (301) 734–5038. For copies of more detailed information on the information collection, contact Mrs. Celeste Sickles, APHIS' Information Collection Coordinator, at (301) 851–2908.

SUPPLEMENTARY INFORMATION:

Title: Pale Cyst Nematode.

OMB Number: 0579–0322.

Type of Request: Revision and extension of approval of an information collection.

Abstract: As authorized by the Plant Protection Act (7 U.S.C. 7701 *et seq.*) (PPA), the Secretary of Agriculture, either independently or in cooperation with States, may carry out operations or measures to detect, eradicate, suppress, control, prevent, or retard the spread of plant pests that are new to or not widely distributed within the United States. This authority has been delegated to the Animal and Plant Health Inspection Service (APHIS) of the U.S. Department of Agriculture, which administers regulations to implement the PPA.

In accordance with the regulations in ''Subpart—Pale Cyst Nematode'' (7 CFR 301.86 through 301.86–9), APHIS restricts the interstate movement of certain articles to help prevent the artificial spread of pale cyst nematode, a major pest of potato crops in cool-temperature areas, to noninfested areas of the United States. The regulations contain requirements for the interstate movement of regulated articles and involve information collection activities, including certificates, permits, and compliance agreements. We are revising the title of the collection to ''Pale Cyst Nematode'' because the title of the original subpart, ''Potato Cyst Nematode,'' was changed in a 2009 final rule.

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 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 0.2057 hours per response.

Respondents: U.S. potato producers, packers, processors, and handlers of potatoes.

Estimated Annual Number of Respondents: 152.

Estimated Annual Number of Responses per respondent: 10.934.

Estimated Annual Number of Responses: 1,662.

Estimated total annual burden on respondents: 342 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 20th day of June 2011.

Kevin Shea,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2011–15871 Filed 6–23–11; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF AGRICULTURE

Animal and Plant Health Inspection Service

[Docket No. APHIS–2011–0061]

Notice of Request for Extension of Approval of an Information Collection; Virus-Serum-Toxin Act and Regulations

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 Virus-Serum-Toxin Act and regulations.

DATES: We will consider all comments that we receive on or before August 23, 2011.

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

- *Federal eRulemaking Portal:* Go to *http://www.regulations.gov/#!documentDetail;D=APHIS-2011-0061-0001.*
- *Postal Mail/Commercial Delivery:* Send your comment to Docket No. APHIS–2011–0061, Regulatory Analysis and Development, PPD, APHIS, Station 3A–03.8, 4700 River Road Unit 118, Riverdale, MD 20737–1238.

Supporting documents and any comments we receive on this docket may be viewed at *http://www.regulations.gov/#!docketDetail;D=APHIS-2011-0061* or in our reading room, which 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.

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) 734–8725. For copies of more detailed information on the information collection, contact Mrs. Celeste Sickles, APHIS' Information Collection Coordinator, at (301) 851–2908.

SUPPLEMENTARY INFORMATION:

Title: Virus-Serum-Toxin Act and Regulations.

OMB Number: 0579–0013.

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

Abstract: Under the Virus-Serum-Toxin Act (21 U.S.C. 151–159), the Animal and Plant Health Inspection Service is authorized to promulgate regulations designed to prevent the importation, preparation, sale, or shipment of harmful veterinary biological products. These regulations are contained in title 9, Code of Federal Regulations, subchapter E, parts 102 to 124.

Veterinary biological products include viruses, serums, toxins, and analogous products of natural or synthetic origin, such as vaccines, antitoxins, or the immunizing components of microorganisms intended for the diagnosis, treatment, or prevention of diseases in domestic animals.

APHIS issues licenses to qualified establishments that produce veterinary 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.

To help ensure that veterinary biological products used in the United States are pure, safe, potent, and effective, APHIS requires certain information collection activities, including establishment license applications, product license applications, product import permit applications, product and test report forms, and field study summaries.

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 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 3.42359 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