Notices

Federal Register

Vol. 81, No. 176

Monday, September 12, 2016

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. APHIS-2016-0063]

Notice of Request for Revision to and Extension of Approval of an Information Collection; Communicable Diseases in Horses

AGENCY: Animal and Plant Health Inspection Service, USDA.

ACTION: Revision to 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 a revision to and extension of approval of an information collection associated with the regulations for the interstate movement of horses that have tested positive for equine infectious anemia.

DATES: We will consider all comments that we receive on or before November 14, 2016.

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

- Federal eRulemaking Portal: Go to http://www.regulations.gov/#!docketDetail;D=APHIS-2016-0063.
- Postal Mail/Commercial Delivery: Send your comment to Docket No. APHIS-2016-0063, 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-2016-0063 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) 799–7039 before coming.

FOR FURTHER INFORMATION CONTACT: For information on the regulations for the interstate movement of horses that have tested positive for equine infectious anemia, contact Dr. Rory Carolan, National Equine Programs, Surveillance, Preparedness and Response Services, VS, APHIS, 4700 River Road Unit 46, Riverdale, MD 20737; (301) 851–3558. For copies of more detailed information on the information collection, contact Ms. Kimberly Hardy, APHIS' Information Collection Coordinator, at (301) 851–2727.

SUPPLEMENTARY INFORMATION:

Title: Communicable Diseases in Horses.

OMB Control Number: 0579–0127. Type of Request: Revision to and extension of approval of an information collection.

Abstract: Under the authority of the Animal Health Protection Act (7 U.S.C. 8301 et seq.), the Animal and Plant Health Inspection Service (APHIS) of the U.S. Department of Agriculture (USDA) regulates the importation and interstate movement of animals and animal products, and conducts various other activities to protect the health of U.S. livestock and poultry.

Equine infectious anemia (EIA) is an infectious and potentially fatal viral disease of equines. There is no vaccine or treatment for the disease. It is often difficult to differentiate from other fever-producing diseases, including anthrax, influenza, and equine encephalitis.

The regulations in 9 CFR 75.4 govern the interstate movement of equines that have tested positive to an official test for EIA (EIA reactors) and provide for the approval of laboratories, diagnostic facilities, and research facilities. Ensuring the safe movement of these horses requires the use of information collection activities, including an EIA laboratory test form, a certificate or permit for the interstate movement of an EIA reactor, a supplemental investigation form if a horse tests positive for EIA, agreements, request for hearing, and written notification of withdrawal of approval.

The regulations also require laboratories conducting an official EIA test to be approved by the APHIS

Administrator in consultation with the appropriate State animal health officials of the State. Approval of a laboratory requires the collection of information, such as the name of the director, location, facilities, appropriate resources, and training and proficiency of employees. This information helps us determine a laboratory's capacity to conduct accurate and reliable testing and to meet the requirements in the regulations. In addition, a laboratory must enter an agreement with APHIS and undergo regular inspections to receive and maintain approval. We are adding these activities to this collection.

We are asking the Office of Management and Budget (OMB) to approve our use of these information collection activities, as described, 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 0.083 hours per response.

Respondents: Producers, veterinarians, State veterinarians, and laboratory directors.

Estimated annual number of respondents: 235,005.

Estimated annual number of responses per respondent: 6. Estimated annual number of

responses: 1,416,075.

Éstimated total annual burden on respondents: 118,010 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 6th day of September 2016.

Jere L. Dick,

Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2016–21840 Filed 9–9–16; 8:45 am]

BILLING CODE 3410-34-P

DEPARTMENT OF AGRICULTURE

Commodity Credit Corporation

Farm Service Agency

Information Collection Request; Application for Payment of Amounts Due Persons Who Have Died, Disappeared, or Have Been Declared Incompetent

AGENCY: Commodity Credit Corporation and Farm Service Agency, USDA. **ACTION:** Notice; request for comments.

SUMMARY: In accordance with the Paperwork Reduction Act of 1995, the Commodity Credit Corporation (CCC) and the Farm Service Agency (FSA) are requesting comments from all interested individuals and organizations on an extension of a currently approved information collection. CCC and FSA use the information to determine whether representatives or survivors of a producer are entitled to receive payments earned by a producer who dies, disappears, or is declared incompetent before receiving payments or other disbursements.

DATES: We will consider comments that we receive by November 14, 2016.

ADDRESSES: We invite you to submit comments on this notice. In your comments, include date, volume, and page number of this issue of the Federal Register. You may submit comments by any of the following methods:

• Federal eRulemaking Portal: Go to http://regulations.gov. Follow the online instructions for submitting comments.

• Mail: Joe Lewis Jr., Agricultural Program Specialist, USDA, FSA STOP 0572, 1400 Independence Avenue SW., Washington, DC 20250–0572.

You may also send comments to the Desk Officer for Agriculture, Office of Information and Regulatory Affairs, Office of Management and Budget, Washington, DC 20503. Copies of the information collection may be requested by contacting Joe Lewis Jr. at the above address.

FOR FURTHER INFORMATION CONTACT: Joe Lewis Jr., (202) 720–0795.

SUPPLEMENTARY INFORMATION:

Title: Application for Payment of Amounts Due Persons Who Have Died, Disappeared, or Have Been Declared Incompetent.

OMB Control Number: 0560–0026. Expiration Date: December 31, 2016. Type of Request: Extension.

Abstract: Persons desiring to claim payments earned, but not yet paid to a person who has died, disappeared, or has been declared incompetent must complete form FSA-325, Application for Payment of Amounts Due Persons Who Have Died, Disappeared, or Have Been Declared Incompetent. This information required by form FSA-325 is used by FSA county office employees to document the relationship of heirs, beneficiaries, or others who claim payment that was earned, but not vet paid to the person who died, disappeared, or who has been declared incompetent, and to determine the share and order of precedence for disbursing payments to such persons.

Information is obtained only when a person claims that they are due a payment that was earned, but not paid to a producer that has died, disappeared, or has been declared incompetent, and documentation is needed to determine if any individuals are entitled to receive such payments or disbursements.

The formula used to calculate the total burden hours is the estimated average time per response times total annual responses.

Estimate of Burden: Public reporting burden for this collection of information is estimated to average 1.5 hours per response. The average travel time, which is included in the total annual burden, is estimated to be 1 hour per respondent.

Respondents: Producers.

Estimated Number of Respondents: 2,000.

Estimated Annual Number of Responses per Respondent: 1. Estimated Total Annual Responses:

2,000.

Estimated Average Time per Responses: 1.5 hours.

Estimated Total Annual Burden Hours on Respondents: 3,000.

We are requesting comments on all aspects of this information collection to help us to:

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

(2) Evaluate the accuracy of FSA's estimate of burden including the

validity of the methodology and assumptions used;

- (3) Enhance the quality, utility, and clarity of the information to be collected:
- (4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

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

Val Dolcini,

Executive Vice President, Commodity Credit Corporation, and Administrator, Farm Service Agency.

[FR Doc. 2016-21654 Filed 9-9-16; 8:45 am]

BILLING CODE 3410-05-P

DEPARTMENT OF AGRICULTURE

Food Safety and Inspection Service [Docket No. FSIS-2016-0029]

Codex Alimentarius Commission: Meeting of the Codex Committee on Food Hygiene

AGENCY: Office of the Deputy Under Secretary for Food Safety, USDA.

ACTION: Notice of public meeting and request for comments.

SUMMARY: The Office of the Deputy Under Secretary for Food Safety, U.S. Department of Agriculture (USDA), and the Food and Drug Administration (FDA), U.S. Department of Health and Human Services (HHS), are sponsoring a public meeting on October 11, 2016. The objective of the public meeting is to provide information and receive public comments on agenda items and draft United States (U.S.) positions to be discussed at the 48th Session of the Codex Committee on Food Hygiene (CCFH) of the Codex Alimentarius Commission (Codex), taking place in Los Angeles, CA, November 7-11, 2016. The Deputy Under Secretary for Food Safety and the FDA recognize the importance of providing interested parties the opportunity to obtain background information on the 48th Session of the CCFH and to address items on the agenda.

DATES: The public meeting is scheduled for Tuesday, October 11, 2016, from 1:00 p.m.—4:00 p.m.

ADDRESSES: The public meeting will take place at the USDA, Jamie L.