animal and animal product trade. To facilitate the export of U.S. animals and animal products, the Animal and Plant Health Inspection Service (APHIS) of the U.S. Department of Agriculture maintains information regarding the import health requirements of other countries for animals and animal products, including aquaculture animals, exported from the United States.

While APHIS does not currently require the approval or certification of laboratories that conduct disease tests for the export of aquaculture animals, some countries that import these animals from the United States require them to be tested for certain diseases and the test results recorded on the export certificates. In addition, the test results must originate from a laboratory approved by the competent authority of the exporting country, which is APHIS in this case. State, university, and private laboratories can voluntarily seek APHIS approval of individual diagnostic methods. Though APHIS does not have regulations for the approval or certification of laboratories that conduct tests for the export of aquaculture animals, APHIS provides this approval as a service to U.S. exporters who export aquaculture animals to countries that require this certification.

APHIS evaluates diagnostic methods for detecting aquatic animal pathogens listed by the World Organization for Animal Health (OIE) in the OIE diagnostic manual and other supporting scientific literature. APHIS lists the laboratories approved to conduct diagnostic testing in support of export health certification of aquatic species at [https://www.aphis.usda.gov/animal_health/lab_info_services/downloads/ApprovedLabs_Aquaculture.pdf](https://www.aphis.usda.gov/animal_health/lab_info_services/downloads/ApprovedLabs_Aquaculture.pdf). Once approved, the laboratories are inspected by APHIS every 2 years to maintain their approval.

The approval of laboratories to conduct tests for the export of aquaculture animals requires the use of certain information collection activities including notification of intent to request approval, application for APHIS approval, protocol statement, submission and recordkeeping of sample copies of diagnostic reports, quality assurance/control plans and their recordkeeping, notification of proposed changes to assay protocols, recordkeeping of supporting assay documentation, and request for removal of approved status.

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 burden for this collection of information is estimated to average 11.4 hours per response.

Respondents: State, university, and private laboratory personnel.

Estimated annual number of respondents: 8.

Estimated annual number of responses per respondent: 70.

Estimated annual number of responses: 560.

Estimated total annual burden on respondents: 6,382 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 4th day of January 2021.

Mark Davidson,
Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2021–00664 Filed 1–7–21; 8:45 am]

BILLING CODE 3410–34–P
SUPPLEMENTARY INFORMATION:

Background

The Animal and Plant Health Inspection Service (APHIS) is proposing to issue permits for the release of an insect, Lilioceris egena, into the continental United States for use as a biological control agent to reduce the severity of air potato (Dioscorea bulbifera) infestations.

Air potato is an herbaceous, twining vine that can grow 65 feet long or more, capable of climbing and out-competing native vegetation. Since its introduction to Florida in 1905, air potato has aggressively spread throughout the State; this species is reportedly naturalized in Georgia, Alabama, Mississippi, Louisiana, Texas, and Hawaii. In 1999, the Florida Department of Agricultural and Consumer Services added air potato to its list of noxious weeds in an attempt to protect the State’s native plant species from being displaced or hybridized. Presently, the air potato is well established in Florida and probably throughout the Gulf States where it has the potential to severely disrupt entire ecosystems.

Existing air potato management options, which include chemical and mechanical control methods, are ineffective, expensive, temporary, or have non-target impacts. Thus, a permit application has been submitted to APHIS for the purpose of releasing an insect, L. egena, into the continental United States for use as a biological control agent to reduce the severity of air potato infestations.

APHIS’ review and analysis of the proposed action are documented in detail in an environmental assessment (EA) titled “Field Release of the Beetle Lilioceris egena (Coleoptera: Chrysomelidae) for Classical Biological Control of Air Potato, Dioscorea bulbifera (Dioscoreaceae), in the Continental United States” (October 2019). We are making the EA available to the public for review and comment. We will consider all comments that we receive on or before the date listed under the heading DATES at the beginning of this notice.

The EA may be viewed on the Regulations.gov website or in our reading room (see ADDRESSES above for instructions for accessing Regulations.gov and information on the location and hours of the reading room). You may request paper copies of the EA by calling or writing to the person listed under FOR FURTHER INFORMATION CONTACT. Please refer to the title of the EA when requesting copies.

The EA has been prepared in accordance with: (1) The National Environmental Policy Act of 1969 (NEPA), as amended (42 U.S.C. 4321 et seq.), (2) regulations of the Council on Environmental Quality for implementing the procedural provisions of NEPA (40 CFR parts 1500–1508), (3) USDA regulations implementing NEPA (7 CFR part 1b), and (4) APHIS NEPA Implementing Procedures (7 CFR part 372).

Done in Washington, DC, this 4th day of January 2021.

Michael Watson,
Acting Administrator, Animal and Plant Health Inspection Service.

[FR Doc. 2021–00063 Filed 1–7–21; 8:45 am]

BILLING CODE 3410–34–P

DEPARTMENT OF COMMERCE

Bureau of Economic Analysis

Agency Information Collection Activities; Submission to the Office of Management and Budget (OMB) for Review and Approval; Comment Request; Direct Investment Surveys: BE–605, Quarterly Survey of Foreign Direct Investment in the United States—Transactions of U.S. Affiliate With Foreign Parent

The Department of Commerce will submit the following information collection request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995, on or after the date of publication of this notice. We invite the general public and other Federal agencies to comment on proposed, and continuing information collections, which helps us assess the impact of our information collection requirements and minimize the public’s reporting burden. Public comments were previously requested via the Federal Register on October 23, 2020 during a 60-day comment period. This notice allows for an additional 30 days for public comments.

Agency: Bureau of Economic Analysis (BEA), Commerce.


OMB Control Number: 0608–0009.

Form Number: BE–605.

Type of Request: Regular submission.

Number of Responses: 17,800 annually.

Average Hours per Response: One hour is the average but may vary considerably among respondents because of differences in company size and complexity.

Estimated Total Annual Burden Hours: 17,800.

Needs and Uses: The Quarterly Survey of Foreign Direct Investment in the United States—Transactions of U.S. Affiliate with Foreign Parent (Form BE–605) obtains quarterly data on transactions and positions between foreign-owned U.S. business enterprises and their “affiliated foreign groups” (i.e., their foreign parents and foreign affiliates of their foreign parents). The survey is a sample survey that covers all U.S. affiliates above a size-exemption level. The sample data are used to derive universe estimates of direct investment transactions, positions, and income in non-benchmark years from similar data reported in the BE–12, Benchmark Survey of Foreign Direct Investment in the United States, which is conducted every five years and will next be conducted for the fiscal year ending in 2022. The data collected through the BE–605 survey are essential for the preparation of the U.S. international transactions, national income and product, and input-output accounts and the international investment position of the United States. The data are needed to measure the size and economic significance of foreign direct investment in the United States, measure changes in such investment, and assess its impact on the U.S. economy.

Affected Public: Businesses or other-for-profit organizations.

Frequency: Quarterly.

Respondent’s Obligation: Mandatory.


This information collection request may be viewed at www.reginfo.gov. Follow the instructions to view Department of Commerce collections currently under review by OMB.

Written comments and recommendations for the proposed information collection should be submitted within 30 days of the publication of this notice on the following website http://www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function and entering either the title of