

Supporting Statement
Special Needs Request Under the
Plant Protection Act
OMB NO. 0579-0291

A. Justification

1. Explain the circumstances that make the collection of information necessary. Identify any legal or administrative requirements that necessitate the collection.

The United States Department of Agriculture (USDA) is responsible for preventing plant diseases or insect pests from entering the United States, preventing the spread of pests and noxious weeds not widely distributed in the United States, and eradicating those imported pests when eradication is feasible. The Plant Protection Act authorizes the Department to carry out this mission.

The Plant Protection Act (PPA, 7 U.S.C. 7701 et seq.) gives authority to the Secretary of Agriculture to prohibit or restrict the importation, entry, exportation, or movement in interstate commerce of any plant, plant product, biological control organism, noxious weed, article, or means of conveyance if the Secretary determines that the prohibition or restriction is necessary to prevent the introduction of plant pests or noxious weeds into the United States, or the dissemination of plant pests or noxious weeds within the United States. The Secretary has delegated this authority to the Administrator of the Animal and Plant Health Inspection Service (APHIS).

Under section 436 of the PPA (7 U.S.C. 7756), no State or political subdivision of a State may regulate the movement in interstate commerce of any article, means of conveyance, plant, biological control organism, plant pest, noxious weed, or plant product in order to: (1) control a plant pest or noxious weed; (2) eradicate a plant pest or noxious weed; or (3) prevent the introduction or dissemination of a biological control organism, plant pest, or noxious weed if the Secretary has issued a regulation or order to prevent the dissemination of the biological control organism, plant pest, or noxious weed within the United States.

APHIS amended its domestic quarantine regulations to establish a process by which a State or political subdivision of a State could request approval to impose prohibitions or restrictions on the movement in interstate commerce of specific articles that are in addition to the prohibitions and restrictions imposed by APHIS. In Title 7, Code of Federal Regulations (CFR), Part 301, "Subpart-Special Need Request" (7 CFR 301.1 through 301.1-3), APHIS sets out procedures for the criteria, action, and submission of a special need request.

APHIS is asking OMB to approve, for three additional years, its use of this information collection activity associated with APHIS' efforts to establish a process for special needs requests.

2. Indicate how, by whom, and for what purpose the information is used. Except for a new collection, indicate the actual use the agency has made of the information received from the current collection.

APHIS believes that this specific information, which would be considered along with more general information available to APHIS, would be necessary for the Administrator to be able to determine whether to grant or deny a request for a special need exemption. The administrator's determination would be based upon his or her review of the information submitted by the State or political subdivision in support of its request and would take into account any comments received. The Administrator's finding that the State or political subdivision has demonstrated, based on sound scientific data or a thorough risk assessment, that there is a special need for additional prohibitions or restrictions would mean that the State or political subdivision would be authorized to impose specific prohibitions or restrictions that go beyond those identified in the regulations or orders issued by APHIS.

3. Describe whether, and to what extent, the collection of information involves the use of automated, electronic, mechanical, or other forms of information technology, e.g., permitting electronic submission of responses, and the basis for the decision for adopting this means of collection. Also describe any consideration of using information technology to reduce burden.

Automation of the data detection survey and other documents are not possible for GPEA. Each of the above criteria to be met in a request for a special needs exemption will require a unique site, host-and/or pest-specific analysis. These analyzes could include a review of existing data and the collection of new data to support the request. In addition, in order for the requester to produce a complete defensible analysis, he or she will need to review available scientific literature, and consult with scientific and other experts. Whoever prepares the request for the special needs exemption will also require a full understanding of the unique circumstances that could justify the request and he or she will need to articulate these in the written request.

4. Describe efforts to identify duplication. Show specifically why any similar information already available cannot be used or modified for use for the purpose described in item 2 above.

The information requested is specific to each plant pest, host, and geographic location and would not be available from any other source.

5. If the collection of information impacts small businesses or other small entities, describe any methods used to minimize burden.

There are no small entities involved with this information collection.

6. Describe the consequences to Federal program or policy activities if the collection is not conducted or is conducted less frequently, as well as any technical or legal obstacles to reducing burden.

Failing to collect this information would leave a State or political subdivision of a State with unique or special needs less than adequately protected from the dissemination of destructive biological control organisms, plant pests, and noxious weeds within the United States. Such gaps could cripple APHIS' efforts to prevent the introduction and establishment of plant pests.

7. Explain any special circumstances that require the collection to be conducted in a manner inconsistent with the general information collection guidelines in 5 CFR 1320.5.

This information collection is conducted in a manner consistent with the guidelines established in 5 CFR 1320.5.

8. Describe efforts to consult with persons outside the agency to obtain their views on the availability of data, frequency of collection, the clarity of instructions and recordkeeping, disclosure, or reporting form, and on the data elements to be recorded, disclosed, or reported. If applicable, provide a copy and identify the date and page number of publication in the Federal Register of the agency's notice, soliciting comments on the information collection prior to submission to OMB.

APHIS did not consult with any outside parties concerning the burden of producing the needed paperwork. However, several states did request exemptions to APHIS' Federal order and regulations.

On Monday, December 8, 2008, pages 74452-74453, APHIS published in the Federal Register a 60-day notice seeking public comments on its plans to request a 3-year renewal of this information collection. No comments from the public were received.

9. Explain any decision to provide any payment or gift to respondents, other than reenumeration of contractors or grantees.

This information collection activity involves no payments or gifts to respondents.

10. Describe any assurance of confidentiality provided to respondents and the basis for the assurance in statute, regulation, or agency policy.

No additional assurance of confidentiality is provided with this information collection. However, the confidentiality of information is protected under 5 U.S.C. 552a.

11. Provide additional justification for any questions of a sensitive nature, such as sexual behavior and attitudes, religious beliefs, and others that are considered private. This justification should include the reasons why the agency considers the questions necessary, the specific uses to be made of the information, the explanation to be given to persons from whom the information is requested, and any steps to be taken to obtain their consent.

This information collection activity asks no questions of personal or sensitive nature.

12. Provide estimates of the hour burden of the collection of information. Indicate the number of respondents, frequency of response, annual hour burden, and an explanation of how the burden was estimated.

. Indicate the number of respondents, frequency of response, annual hour burden, and an explanation of how the burden was estimated. If this request for approval covers more than one form, provide separate hour burden estimates for each form and aggregate the hour burdens in Item 13 of OMB Form 83-I.

See APHIS Form 71 for hour burden estimates.

. Provide estimates of annualized cost to respondents for the hour burdens for collections of information, identifying and using appropriate wage rate categories.

1,600 hours X \$26.32 = \$ 42,112.00.

13. Provide estimates of the total annual cost burden to respondents or recordkeepers resulting from the collection of information, (do not include the cost of any hour burden shown in items 12 and 14). The cost estimates should be split into two components: (a) a total capital and start-up cost component annualized over its expected useful life; and (b) a total operation and maintenance and purchase of services component.

There is zero annual cost burden associated with capital and start-up, operation and maintenance, and purchase of services in connection with this program.

14. Provide estimates of annualized cost to the Federal government. Provide a description of the method used to estimate cost and any other expense that would not have been incurred without this collection of information.

The estimated cost for the Federal Government is \$ 11,770.43 (see APHIS 79 attached).

15. Explain the reasons for any program changes or adjustments reported in Items 13 or 14 of the OMB Form 83-I.

This is a continuation of a previously approved program. There are no changes in burden.

16. For collections of information whose results are planned to be published, outline plans for tabulation and publication.

APHIS has no plans to publish information collected in connection with this program.

17. If seeking approval to not display the expiration date for OMB approval of the information collection, explain the reasons that display would be inappropriate.

There are no USDA forms associated with this collection.

18. Explain each exception to the certification statement identified in the "Certification for Paperwork Reduction Act."

APHIS is able to certify compliance with all the provisions under the Act.

B. Collections of Information Employing Statistical Methods

Statistical methods are not used in this information collection.