To receive additional written information and to request an application kit, call 1–888–GRANTS4 (1–888–472–6874). You will be asked to leave your name and address and will be instructed to identify the Announcement number of interest. Please refer to Program Announcement 99113 when you request information. After reviewing the Program Announcement, for business management assistance, contact: Joanne Wojick, Grants Management Specialist Grants Management Branch, Procurement and Grants Office, Announcement 99113, Centers for Disease Control and Prevention (CDC), 2920 Brandywine Road, Suite 3000, Atlanta, GA 30341–4146, Telephone (404) 488–2717, Email address jcw6@cdc.gov.

For program technical assistance, contact: Paul Burlack, Centers for Disease Control and Prevention, National Center for Injury Prevention and Control, 4770 Buford Highway N.E., Mailstop F41, Atlanta, GA 30341–3724, Telephone (770) 488–4031, pab5@cdc.gov.

Dated: May 3, 1999.

John L. Williams,

Director, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–11484 Filed 5–6–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Collection of Fees at United States Ports Designated To Conduct Rodent Infestation Inspections and Issue Deratting and Deratting Exemption Certificates

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).

ACTION: Notice.

SUMMARY: The Centers for Disease Control and Prevention (CDC) is adopting a requirement for collection of user fees for conducting rodent infestation inspection of ships, and issuing Deratting and Deratting Exemption Certificates. While the United States does not require these certificates for ships to enter its seaports, the United States conducts inspections and issues certificates in accordance with 42 CFR 71.46 and Article 17 of the International Health Regulations.

DATES: Effective date is June 6, 1999.

FOR FURTHER INFORMATION CONTACT: James E. Barrow, Chief, Program Operations Branch, Division of Quarantine, National Center for Infectious Diseases, CDC, Mailstop E–03, Atlanta, Georgia 30333, telephone: (404) 639–8107, fax (404) 639–2599, e-mail: jeb1@cdc.gov.

Authority: 42 U.S.C. 264–271, 42 CFR 71.46, IHR Articles 17 and 53.

SUPPLEMENTARY INFORMATION: A proposal to charge fees for rodent infestation inspections of ships, and issuance of Deratting and Deratting Exemption Certificates, where these services are provided directly by employees or vendors of the CDC was published in the **Federal Register** on November 24, 1998 (63 FR 64967).

Comments Received

Interested parties were afforded an opportunity to comment on the proposal. One media inquiry and no comments were received during the comment period.

Conclusion

CDC has determined that in the interest of defraying the cost of inspection and certificate issuance, user fees will be implemented for rodent infestation of ships, and issuance of Deratting and Deratting Exemption Certificates. Rodent infestation inspections for ships will be conducted at 11 major ports upon request, including: Baltimore, Maryland; Honolulu, Hawaii; Houston, Texas; Jacksonville, Florida; Los Angeles, California; Miami, Florida; New Orleans, Louisiana; New York, New York; San Francisco, California; Savannah, Georgia; and Seattle, Washington.

Cost Impact

The United States does not require a Rodent Infestation Inspection, or a Deratting or Deratting Exemption Certificate, for ships to enter its seaports. Article 17 of the International Health Regulations, published by the World Health Organization, Geneva, Switzerland, requires that each Health Administration provide these services, and Article 82 outlines the criteria for charging fees. 42 CFR 71.46 authorizes the performance of these services by the Public Health Service as carried out by CDC. CDC has for many years offered these services at no cost to the owners or agents of ships requesting them. These user fees will, in a manner consistent with most other countries, pass the cost of conducting these services along as a charge to those receiving and benefitting from the inspections and certificates.

Regulatory Impact

The requirements adopted herein will not have substantial direct effects on the States, on the relationship between the national government and the States, or on the distribution of power and responsibilities among the various levels of government. Therefore, in accordance with Executive Order 12612, it is determined that this action does not have sufficient federalism implications to warrant the preparation of a Federalism Assessment.

This action (1) is not a ''significant regulatory action'' under Executive Order 12866; (2) will not have a significant economic impact, positive or negative, on a substantial number of small entities under the criteria of the Regulatory Flexibility Act; and (3) does not impose additional costs upon any State or local government as a result of a mandate imposed upon them as a government agency, as described in the Unfunded Mandates Reform Act.

Collection of Information

This final rule contains no new collection-of-information requirements under the Paperwork Reduction Act (44 U.S.C. 3501 *et seq.*).

User Fee Administration

1. Effective June 6, 1999, user fees will be collected for all rodent infestation inspections of ships, and the associated issuance of Deratting and Deratting Exemption Certificates, by CDC and its vendors.

2. Rodent infestation inspections for ships will be conducted at 11 ports upon request, including: Baltimore, Maryland; Honolulu, Hawaii; Houston, Texas; Jacksonville, Florida; Los Angeles, California; Miami, Florida; New Orleans, Louisiana; New York, New York; San Francisco, California; Savannah, Georgia; and Seattle, Washington.

3. Costs are determined by taking into consideration such items as salaries, benefits, vendor services, printing, supplies, and agency overhead. The charge for the first full year during which fees for rodent infestation inspections and issuance of Deratting and Deratting Exemption Certificates are assessed is $150 for each inspection conducted. Shipping companies will be provided by mail instructions for submitting fees. The fees will be due at the address specified in the bill, not later than 30 days following the inspection. Arrangements may also be made to prepay user fees and draw against those prepayments.

Dated: May 3, 1999.

Joseph R. Carter,

Acting Associate Director for Management and Operation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 99–11485 Filed 5–6–99; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Quarterly Performance Report, ORR–6.

OMB No.: 0970–0036.

Description: Data gathered from the Quarterly Performance Report (Form ORR–6) are used by ORR to estimate the number of months of Refugee Cash Assistance (RCA) and Refugee Medical Assistance (RMA) that ORR can provide based on appropriations; to determine priorities; and standards, budget requests, and assistance policies; to analyze data on service caseloads and program outcomes in order to monitor performance; and to compute refuge medical assistance (RMA) utilization rates.

Respondents: State, Local or Tribal Governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Program Estimates (CMA)	48	4	3.875	744

Estimated Total Annual Burden Hours: 744.

Additional Information: Copies of the proposed collection may be writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW, Washington, DC 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 to 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW, Washington, DC 20503, Attn: Ms. Lori Schack.

Dated: May 3, 1999.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 99–11536 Filed 5–6–99; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–0123]

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Food Labeling; Notification Procedures for Statements on Dietary Supplements

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Submit written comments on the collection of information by June 7, 1999.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attn: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Peggy Schlosburg, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1223.

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has submitted the following proposed collection of information to OMB for review and clearance.

Food Labeling; Notification Procedures for Statements on Dietary Supplements—21 CFR 101.93 (OMB Control Number 0910–0331—Extension)

Section 403(r)(6) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 343(r)(6)) requires that the agency be notified by manufacturers, packers, and distributors of dietary supplements that they are marketing a dietary supplement product that bears on its label or in its labeling a statement provided for in section 403(r)(6) of the act. Section 403(r)(6) of the act requires that the agency be notified, with a submission about such statements, no later than 30 days after the first marketing of the dietary supplement. Information that is required in the submission includes: (1) The name and address of the manufacturer, packer, or distributor of the dietary supplement product; (2) the text of the statement that is being made; (3) the name of the dietary ingredient or supplement that is the subject of the statement; (4) the name of the dietary supplement (including the brand name); and (5) a signature of a responsible individual who can certify the accuracy of the information presented.

The agency established § 101.93 (21 CFR 101.93) as the procedural regulation for this program. Section 101.93 provides details of the procedures associated with the submission and identifies the information that must be included in order to meet the requirements of section 403 of the act.

Description of Respondents: Businesses or other for-profit organizations.

In the **Federal Register** of February 4, 1999 (64 FR 5664), the agency requested comments on the proposed collections of information. No comments were received.

FDA estimates the burden of this collection of information as follows: