degradeable'' without qualification on the front of the package. Tender's website and promotional materials also made the claim. According to the FTC complaint, respondent represented that Fresh Bath Wipes and Fresh Bath Travel Wipes and their packages will completely break down and return to nature, *i.e.*, decompose into elements found in nature, within a reasonably short period of time after customary disposal. The complaint alleges respondent's biodegradable claim is false because a substantial majority of total household waste is disposed of either in landfills, incinerators, or recycling facilities and these customary disposal methods do not present conditions that would allow for the wipes and their packaging to completely break down and return to nature, *i.e.*, decompose into elements found in nature, within a reasonably short period of time. The complaint further alleges that respondent failed to have substantiation for the biodegradable claim. The proposed consent order contains provisions designed to prevent respondent from engaging in similar acts and practices in the future.

Part I.A of the proposed order prohibits respondent from making a representation that certain of its products are degradable unless the representation is true, not misleading, and substantiated by competent and reliable scientific evidence. Part I.B prohibits respondent from making any other environmental benefit claim about such products, unless at the time the representation is made, it is truthful and not misleading, and substantiated by competent and reliable evidence, which when appropriate must be competent and reliable scientific evidence.

Part II of the proposed order requires respondent to specify whether its degradability claim applies to the product, package, or components of either.

Parts III through VI require respondent to keep copies of relevant advertisements and materials substantiating claims made in the advertisements; to provide copies of the order to certain of their personnel; to notify the Commission of changes in corporate structure that might affect compliance obligations under the order; and to file compliance reports with the Commission and respond to other requests from FTC staff. Part VII provides that the order will terminate after twenty (20) years under certain circumstances.

The purpose of this analysis is to facilitate public comment on the proposed order. It is not intended to constitute an official interpretation of the agreement and proposed order or to modify in any way their terms.

By direction of the Commission.

**Donald S. Clark**

*Secretary.*

[FR Doc. E9–14107 Filed 6–15–09: 8:45 am]

**BILLING CODE: 6750–01–S**

## DEPARTMENT OF DEFENSE

## GENERAL SERVICES ADMINISTRATION

## NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

**[OMB Control No. 9000–0053]**

### Federal Acquisition Regulation; Information Collection; Permits, Authorities, or Franchises Certification

**AGENCIES:** Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

**ACTION:** Notice of request for comments regarding the reinstatement of a previously existing OMB clearance.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal Acquisition Regulation (FAR), Regulatory Secretariat will be submitting to the Office of Management and Budget (OMB) a request to review and approve an extension of a currently approved information collection requirement concerning Permits, Authorities, or Franchises Certification.

Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

**DATES:** Submit comments on or before August 17, 2009.

**ADDRESSES:** Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: General Services Administration, Regulatory Secretariat (VPR), 1800 F Street NW., Room 4041, Washington, DC 20405. Please cite OMB Control No. 9000–0053, Permits, Authorities, or Franchises Certification, in all correspondence.

**FOR FURTHER INFORMATION CONTACT:** Ms. Jeritta Parnell, Procurement Analyst, Contract Policy Division, GSA, (202) 501–4082.

**A. Purpose**

This certification and copies of authorizations are needed to determine that the offeror has obtained all authorizations, permits, etc., required in connection with transporting the material involved. The contracting officer reviews the certification and any documents requested to ensure that the offeror has complied with all regulatory requirements and has obtained any permits, licenses, etc., that are needed.

**B. Annual Reporting Burden**

Respondents: *1,106.*
Responses Per Respondent: *3.*
Annual Responses: *3,318.*
Hours Per Response: *.094.*
Total Burden Hours: *312.*

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VPR), 1800 F Street NW., Room 4041, Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 9000–0053, Permits, Authorities, or Franchises Certification, in all correspondence.

Dated: June 9, 2009.

**Al Matera,**

*Director, Office of Acquisition Policy.*

[FR Doc. E9–14104 Filed 6–15–09; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF DEFENSE

## GENERAL SERVICES ADMINISTRATION

## NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

**[OMB Control No. 9000–0082]**

### Federal Acquisition Regulation; Information Collection; Economic Purchase Quantity—Supplies

**AGENCIES:** Department of Defense (DOD), General Services Administration (GSA), and National Aeronautics and Space Administration (NASA).

**ACTION:** Notice of reinstatement request for an information collection requirement regarding an existing OMB clearance.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35), the Federal

Acquisition Regulation, Regulatory Secretariat (VPR) will be submitting to the Office of Management and Budget (OMB) a request to reinstate a previously approved information collection requirement concerning Economic Purchase Quantity—Supplies.

Public comments are particularly invited on: Whether this collection of information is necessary; whether it will have practical utility; whether our estimate of the public burden of this collection of information is accurate, and based on valid assumptions and methodology; ways to enhance the quality, utility, and clarity of the information to be collected; and ways in which we can minimize the burden of the collection of information on those who are to respond, through the use of appropriate technological collection techniques or other forms of information technology.

**DATES:** Submit comments on or before August 17, 2009.

**ADDRESSES:** Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden, to: General Services Administration, Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC 20405. Please cite OMB Control No. 9000–0082, Economic Purchase Quantity—Supplies, in all correspondence.

**FOR FURTHER INFORMATION CONTACT:** Ms. Lori Sakalos, Procurement Analyst, Contract Policy Division, GSA, (202) 208–0498.

**SUPPLEMENTARY INFORMATION:**

**A. Purpose**

The provision at 52.207–4, Economic Purchase Quantity—Supplies, invites offerors to state an opinion on whether the quantity of supplies on which bids, proposals, or quotes are requested in solicitations is economically advantageous to the Government. Each offeror who believes that acquisitions in different quantities would be more advantageous is invited to (1) recommend an economic purchase quantity, showing a recommended unit and total price, and (2) identify the different quantity points where significant price breaks occur. This information is required by Public Law 98–577 and Public Law 98–525.

**B. Annual Reporting Burden**

*Respondents:* 1,524.
*Responses per Respondent:* 25.
*Annual Responses:* 38,100.
*Hours per Response:* .83.
*Total Burden Hours:* 31,623.

*Obtaining Copies of Proposals:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VPR), 1800 F Street, NW., Room 4041, Washington, DC, 20405, telephone (202) 501–4755. Please cite OMB Control No. 9000–0082, Economic Purchase Quantity—Supplies, in all correspondence.

Dated: June 9, 2009.

**Al Matera,**

*Director, Office of Acquisition Policy.*

[FR Doc. E9–14105 Filed 6–15–09; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2009–N–0261]**

### Agency Emergency Processing Under Office of Management and Budget Review; Reporting and Recordkeeping Requirements for Reportable Food Registry

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for emergency processing under the Paperwork Reduction Act of 1995 (the PRA). This notice solicits comments on the proposed collection of information associated with the draft guidance document entitled ''Questions and Answers Regarding the Reportable Food Registry as Established by the Food and Drug Administration Amendments Act of 2007.'' The draft guidance, when finalized, will assist the industry in complying with the Reportable Food Registry requirements prescribed by the Food and Drug Administration Amendments Act of 2007 (FDAAA).

**DATES:** Fax written comments on the collection of information by July 16, 2009. FDA is requesting approval of this emergency processing by August 17, 2009.

**ADDRESSES:** To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974. All comments should be identified with the docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794.

**SUPPLEMENTARY INFORMATION:** FDA has requested emergency processing of this proposed collection of information under section 3507(j) of the PRA (44 U.S.C. 3507(j)) and 5 CFR 1320.13. On September 27, 2007, the President signed FDAAA into law (Public Law 110–85). Section 1005 of FDAAA amends the Federal Food, Drug, and Cosmetic Act (the act) by creating a new section 417 (21 U.S.C. 350f), among other things. Section 417 of the act requires the Secretary of Health and Human Services to establish, within FDA, a Reportable Food Registry (the Registry); the Registry is to be established not later than 1 year after the date of enactment (i.e., by September 27, 2008).

To further the development of the Registry, section 417 of the act requires FDA to establish, also within 1 year after the date of enactment (i.e., by September 27, 2008), an electronic portal (the Reportable Food electronic portal) by which instances of reportable food must be submitted to FDA by responsible parties and may be submitted by public health officials.

FDA made the decision that the most efficient and cost effective means to implement the requirements of section 417 of the act relating to the Registry was to utilize the business enterprise system currently under development within the agency: The MedWatch$^{Plus}$ Portal. This would permit the agency to establish an electronic portal through which instances of reportable food may be submitted to the agency. However, FDA recognized that the MedWatch$^{Plus}$ Portal would not be implemented in time to meet the September 27, 2008, deadline for establishing the Reportable Food electronic portal and therefore announced that it was delaying its implementation until spring 2009 (73 FR 30405; May 27, 2008).

The agency now expects the system to be operational on September 8, 2009.

Section 1005(f) of FDAAA required FDA to issue guidance to industry about submitting reports through the electronic portal of instances of reportable food and providing notifications to other persons in the supply chain of such article of food. In a notice published in the **Federal Register** of June 11, 2009, FDA announced the availability of the draft guidance document entitled ''Questions