request to review the documents to ensure that the contractor has complied with all regulatory requirements.

**C. Annual Reporting Burden**

*Respondents:* 8,256.
*Total Annual Responses:* 8,256.
*Total Burden Hours:* 4,128.

**D. Public Comment**

A 60 day notice was published in the **Federal Register** at 84 FR 13921, on April 9, 2019. No comments were received.

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

Dated: July 16, 2019.

**Janet Fry,**

*Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.*

[FR Doc. 2019–15451 Filed 7–19–19; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF DEFENSE

## GENERAL SERVICES ADMINISTRATION

## NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

**[OMB Control No. 9000–0082; Docket No. 2019–0003; Sequence No. 2]**

### Submission for OMB Review; Economic Purchase Quantity—Supplies

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

**ACTION:** Notice.

**SUMMARY:** Under the provisions of the Paperwork Reduction Act, the Regulatory Secretariat Division will be submitting to the Office of Management and Budget (OMB) a request to review and approve a revision and extension of a previously approved information collection requirement concerning economic purchase quantity—supplies.

**DATES:** Submit comments on or before August 21, 2019.

**ADDRESSES:** Submit comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to: Office of Information and Regulatory Affairs of OMB, Attention: Desk Officer for GSA, Room 10236, NEOB, Washington, DC 20503. Additionally submit a copy to GSA by any of the following methods:

- *Federal eRulemaking Portal:* This website provides the ability to type short comments directly into the comment field or attach a file for lengthier comments. Go to *http://www.regulations.gov* and follow the instructions on the site.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW, Washington, DC 20405. ATTN: Ms. Mandell/IC 9000–0082, Economic Purchase Quantity—Supplies.

*Instructions:* All items submitted must cite Information Collection 9000–0082, Economic Purchase Quantity—Supplies. Comments received generally will be posted without change to *http://www.regulations.gov,* including any personal and/or business confidential information provided. To confirm receipt of your comment(s), please check *www.regulations.gov,* approximately two-to-three days after submission to verify posting (except allow 30 days for posting of comments submitted by mail).

**FOR FURTHER INFORMATION CONTACT:** Mr. Michael O. Jackson, Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA, 202–208–4949, or email at *michaelo.jackson@gsa.gov.*

**SUPPLEMENTARY INFORMATION:**

**A. OMB Number, Title, and Any Associated Form(s)**

9000–0082, Economic Purchase Quantity—Supplies.

**B. Needs and Uses**

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.

**C. Annual Reporting Burden**

*Respondents:* 3,000.
*Total Annual Responses:* 75,000.
*Total Burden Hours:* 75,000.
*Affected Public:* Business or other for-profit entities.
*Respondent's Obligation:* Voluntary.
*Type of Request:* Extension of a currently approved collection.
*Reporting Frequency:* On occasion.

**D. Public Comment**

A 60 day notice was published in the **Federal Register** at 84 FR 10828 on March 22, 2019. No comments were received.

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

Dated: July 16, 2019.

**Janet Fry,**

*Director, Federal Acquisition Policy Division, Office of Governmentwide Acquisition Policy, Office of Acquisition Policy, Office of Governmentwide Policy.*

[FR Doc. 2019–15452 Filed 7–19–19; 8:45 am]

**BILLING CODE 6820–EP–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2012–N–0873]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Bar Code Label Requirement for Human Drug and Biological Products

**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 review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Fax written comments on the collection of information by August 21, 2019.

**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: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All

comments should be identified with the OMB control number 0910–0537. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

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

**Bar Code Label Requirement for Human Drug and Biological Products**

*OMB Control Number 0910–0537—Extension*

In the **Federal Register** of February 26, 2004 (69 FR 9120), FDA issued a final rule that requires human drug product and biological product labels to have bar codes. Specifically, the final rule requires bar codes on most human prescription drug products and on over-the-counter (OTC) drug products that are dispensed under an order and commonly used in healthcare facilities. It also requires machine-readable information on blood and blood components. For human prescription drug products and OTC drug products that are dispensed under an order and commonly used in healthcare facilities, the bar code must contain the national drug code number for the product. For blood and blood components, the final rule specifies the minimum contents of the label in a format that is machine readable and approved for use by the Director, Center for Biologics Evaluation and Research. We believe that the final rule helps reduce the number of medication errors in hospitals and other healthcare settings by allowing healthcare professionals to use bar code scanning equipment to verify that the right drug (in the right dose and right route of administration) is being given to the right patient at the right time.

Although most of the information collections created by the final rule have now been incorporated in OMB approved information collections supporting the applicable regulations, respondents to the collection may continue to seek an exemption from the bar code label requirement under § 201.25(d) (21 CFR 201.25(d)). Section 201.25(d) requires submission of a written request for an exemption and describes the information that must be included in such a request. Based on the number of exemption requests we have received previously, we estimate that approximately two exemption requests will be submitted annually and each exemption request will require 24 hours to complete. This results in an annual reporting burden of 48 hours, as reflected in table 1.

In the **Federal Register** of November 1, 2018 (83 FR 54930), we published a 60-day notice requesting public comment on the proposed collection of information. No comments were received.

We estimate the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN [1]

| 21 CFR section | Number of respondents | Number of responses per respondent | Total annual responses | Average burden per response | Total hours |
|---|---|---|---|---|---|
| 21 CFR 201.25(d) | 2 | 1 | 2 | 24 | 48 |

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

Based on a review of the information collection since our last request for OMB approval, we have made no adjustments to our burden estimate.

Dated: July 16, 2019.

**Lowell J. Schiller,**

*Principal Associate Commissioner for Policy.*

[FR Doc. 2019–15488 Filed 7–19–19; 8:45 am]

**BILLING CODE 4164–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. FDA–2019–N–1265]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Labeling: Nutrition Facts Label and Supplement Facts Label

**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 review and clearance under the Paperwork Reduction Act of 1995.

**DATES:** Fax written comments on the collection of information by August 21, 2019.

**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: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0813. Also include the FDA docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Domini Bean, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–5733, *PRAStaff@fda.hhs.gov.*

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

**Food Labeling: The Nutrition Facts Label and Supplement Facts Label—21 CFR 101.9**

*OMB Control Number 0910–0813—Extension*

This information collection supports requirements for the Nutrition Facts and Supplemental Facts labels. Section 403(q) of the Federal Food, Drug, and Cosmetic Act (FD&C Act) (21 U.S.C. 343(q)) specifies certain nutrients to be declared in nutrition labeling and authorizes the Secretary of Health and Human Services (Secretary) to require other nutrients to be declared if the Secretary determines that a nutrient will provide information regarding the nutritional value of such food that will assist consumers in maintaining healthy dietary practices. The Secretary also has