General description of report: The notices are authorized under the Securities Exchange Act of 1934, as amended 1 (the Act), which requires a financial institution that is a broker or dealer of government securities dealer to notify the appropriate regulatory agency (ARA) that it is a government securities broker or a government securities dealer (Form G-FIN notice), or that it has ceased to act as such (Form G-FINW notice). In addition, 15 U.S.C. 780-5(b)(1) directs the Treasury to adopt rules requiring every government securities broker and government securities dealer to collect information and to provide reports to the applicable ARA. The Board is an ARA. 15 U.S.C. 78c(a)(34)(G)(ii). Further support for the creation and collection of these notices by the Board is found in Treasury regulations, authorized by 15 U.S.C. 780-5(b)(l), instructing that any amendments or corrections to a financial institution's status as a government securities broker or dealer also be filed with the ARA on the Form GFIN notice. 17 CFR 400.5(b).

Under the Act, the Secretary of the Treasury is authorized to exempt any government securities broker or dealer, or class thereof, from the notice requirement of section 78o-5(a)(1)(B). See 15 U.S.C. 780-5(a)(5). Thus, the obligation to file the notices with the Board is mandatory for those financial institutions for which the Board serves as the ARA, unless the financial institution is exempted from the notice filing requirement by Treasury regulations (17 CFR part 401). If an exemption no longer applies, the institution must immediately file a notice. The filing of these notices is event generated.

Respondents file two copies of the notices directly with the Board. Under the statute, the Board forwards one copy to the Securities and Exchange Commission (SEC), and the notices are then made public by the SEC. 15 U.S.C. 780–5(a)(l)(B)(iii). While the statute only requires the SEC to produce the notices to the public, the notices are also available to the public upon request made to the Board. Accordingly, the Board does not consider these data to be confidential.

Abstract: The Act requires financial institutions to notify their ARA of their intent to engage in government securities broker or dealer activity, to amend information submitted previously, and to record their termination of such activity. The Federal Reserve is the ARA for state member banks, foreign banks, uninsured

state branches or state agencies of foreign banks, commercial lending companies owned or controlled by foreign banks, and Edge corporations. The Federal Reserve uses the information in its supervisory capacity to measure compliance with the Act.

Board of Governors of the Federal Reserve System, February 24, 2016.

#### Robert deV. Frierson,

Secretary of the Board.

[FR Doc. 2016-04282 Filed 2-26-16; 8:45 am]

BILLING CODE 6210-01-P

### **DEPARTMENT OF DEFENSE**

# GENERAL SERVICES ADMINISTRATION

# NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000-0070; Docket 2015-0055; Sequence 26]

# Submission for OMB Review; Payments

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

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

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 an extension of a previously approved information collection requirement concerning Payments. A 60-day notice was published in the Federal Register at 80 FR 76492 on December 9, 2015. No comments were received.

**DATES:** Submit comments on or before March 30, 2016.

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:

• Regulations.gov: http:// www.regulations.gov. Submit comments via the Federal eRulemaking portal by searching the OMB control number. Select the link "Submit a Comment" that corresponds with "Information Collection 9000–0070, Payments". Follow the instructions provided at the

- "Submit a Comment" screen. Please include your name, company name (if any), and "Information Collection 9000–0070, Payments" on your attached document.
- Mail: General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0070, Payments.

Instructions: Please submit comments only and cite Information Collection 9000–0070, Payments, in all correspondence related to this collection. 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:

Kathlyn Hopkins, Procurement Analyst, Office of Acquisition Policy, GSA at 202–969–7226 or email at kathlyn.hopkins@gsa.gov.

## SUPPLEMENTARY INFORMATION:

## A. Purpose

Firms performing under Federal contracts must provide adequate documentation to support requests for payment under these contracts. The documentation may range from a simple invoice to detailed cost data. The information is usually submitted once, at the end of the contract period or upon delivery of the supplies or services, but could be submitted more often depending on the payment schedule established under the contract (see FAR 52.232-1 through 52.232-4, and FAR 52.232-6 through 52.232-11). The information is used to determine the proper amounts to be paid to Federal contractors.

## **B.** Annual Reporting Burden

Respondents: 80,000. Responses per Respondent: 120. Total Responses: 9,600,000. Hours per Response: .25. Total Burden Hours: 2,400,000.

# C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the Federal Acquisition Regulations (FAR), and 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

<sup>&</sup>lt;sup>1</sup> 15 U.S.C. 780-5(a)(l)(B)(i).

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.

Obtaining Copies of Proposals: 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. 9000–0070, Payments, in all correspondence.

Dated: February 23, 2016.

#### Lorin S. Curit,

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

[FR Doc. 2016–04280 Filed 2–26–16; 8:45 am]

BILLING CODE 6820-EP-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC-2013-0025; Docket Number NIOSH-266]

### **Issuance of Final Publication**

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

**ACTION:** Notice of issuance of final publication.

**SUMMARY:** The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), announces the availability of the following publication: *NIOSH Criteria for a Recommended Standard:* Occupational Exposure to Heat and Hot Environments [2016–106].

**ADDRESSES:** This document may be obtained at the following link: http://www.cdc.gov/niosh/docs/2016-106/.

FOR FURTHER INFORMATION CONTACT: Brenda Jacklitsch, NIOSH Education and Information Division, 1090 Tusculum Ave, Mail Stop C–32, Cincinnati, OH 45226, email address: gwe6@cdc.gov.

Dated: February 19, 2016.

### John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2016–04297 Filed 2–26–16; 8:45 am]

BILLING CODE 4163-19-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Food and Drug Administration**

[Docket No. FDA-2011-D-0147]

Agency Information Collection
Activities; Announcement of Office of
Management and Budget Approval;
Guidance for Industry and Food and
Drug Administration Staff; Section
905(j) Reports: Demonstrating
Substantial Equivalence for Tobacco
Products and Demonstrating the
Substantial Equivalence of a New
Tobacco Product: Responses to
Frequently Asked Questions

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

**ACTION:** Notice.

SUMMARY: The Food and Drug
Administration (FDA) is announcing
that a collection of information entitled
"Guidance for Industry and Food and
Drug Administration Staff; Section
905(j) Reports: Demonstrating
Substantial Equivalence for Tobacco
Products and Demonstrating the
Substantial Equivalence of a New
Tobacco Product: Responses to
Frequently Asked Questions" has been
approved by the Office of Management
and Budget (OMB) under the Paperwork
Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, PRAStaff@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: On July 16, 2015, the Agency submitted a proposed collection of information entitled "Guidance for Industry and Food and Drug Administration Staff; Section 905(j) Reports: Demonstrating Substantial Equivalence for Tobacco Products and Demonstrating the Substantial Equivalence of a New Tobacco Product: Responses to Frequently Asked Questions" to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

OMB has now approved the information collection and has assigned OMB control number 0910–0673. The approval expires on January 31, 2019. A copy of the supporting statement for this information collection is available on the Internet at http://www.reginfo.gov/public/do/PRAMain.

Dated: February 23, 2016.

#### Leslie Kux,

 $Associate\ Commissioner\ for\ Policy.$  [FR Doc. 2016–04222 Filed 2–26–16; 8:45 am]

BILLING CODE 4164-01-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Food and Drug Administration**

[Docket No. FDA-2012-N-1093]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Additive Petitions and Investigational Food Additive Exemptions

**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 March 30, 2016.

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 0910–0546. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, 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.