FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than September 27, 2016.

A. Federal Reserve Bank of Cleveland (Nadine Wallman, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to *Comments.applications@clev.frb.org:*

1. *Eloise Hamilton Campbell,* Danville, Kentucky; to acquire additional shares of Middlefork Financial Group, Hyden, Kentucky, and thereby indirectly acquire control of Farmers & Traders Bank of Campton, Campton, Kentucky, Hyden Citizens Bank, Hyden, Kentucky and Farmers State Bank, Booneville, Kentucky.

Board of Governors of the Federal Reserve System, September 7, 2016.

Michele Taylor Fennell,

Assistant Secretary of the Board.

[FR Doc. 2016–21856 Filed 9–9–16; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[Docket 2016–0053; Sequence 16; OMB Control No. 9000–0095]

Submission for OMB Review; Commerce Patent Regulations

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 Department of Commerce patent regulations. A notice was published in the **Federal Register** at 81 FR 24103 on April 25, 2016. No comments were received.

DATES: Submit comments on or before October 12, 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 9000–0095, Commerce Patent Regulations''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0095, Commerce Patent Regulations'' on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0095, Commerce Patent Regulations.

Instructions: Please submit comments only and cite Information Collection 9000–0095, Commerce Patent Regulations, 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: Ms. Zenaida Delgado, Procurement Analyst, Office of Governmentwide Acquisition Policy, GSA, 202–969–7207 or email *zenaida.delgado@gsa.gov.*

SUPPLEMENTARY INFORMATION:

A. Purpose

FAR subpart 27.3, Patents Rights under Government Contracts, implements the Department of Commerce regulation (37 CFR 401) based on chapter 18 of title 35 U.S.C., Presidential Memorandum on Government Patent Policy to the Heads of Executive Departments and Agencies, dated February 18, 1983, and Executive Order 12591, Facilitating Access to Science and Technology, dated April 10, 1987. Under the subpart, a contracting officer may insert clauses 52.227–11, Patent Rights-Ownership by the Contractor, or 52.227–13, Patent Rights-Ownership by the Government, in solicitations and contracts pertaining to inventions made in the performance of experimental, developmental, or research work.

In accordance with the clauses, a Government contractor must report all subject inventions to the contracting officer, submit a disclosure of the invention, and identify any publication, or sale, or public use of the invention (52.227–11(c), 52.227–13(e)(1)). The contracting officer may modify 52.227–11(e) or otherwise supplement the clause to require contractors to submit periodic or interim and final reports listing subject inventions (27.303(b)(2)(i) and (ii)). In order to ensure that subject inventions are reported, the contractor is required to establish and maintain effective procedures for identifying and disclosing subject inventions (52.227–11, Alternate IV; 52.227–13(e)(1)). In addition, the contractor must require his employees, by written agreements, to disclose subject inventions (52.227–11(e)(2); 52.227–13(e)(4)). The contractor also has an obligation to utilize the subject invention, and agree to report, upon request, the utilization or efforts to utilize the subject invention (27.302(e); 52.227–11(f)).

B. Annual Reporting Burden

Respondents: 3759.
Responses per Respondent: 3.8.
Total Responses: 14,338.
Hours per Response: 4.0.
Total Burden Hours: 57,352.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary for the proper performance of functions of the 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 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 (GSA), Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0095, Commerce Patent Regulations, in all correspondence.

Dated: September 6, 2016.

Lorin S. Curit,

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

[FR Doc. 2016–21790 Filed 9–9–16; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–0852]

Agency Forms Undergoing Paperwork Reduction Act Review

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

ACTION: Notice with comment period; withdrawal.

SUMMARY: The Centers for Disease Control and Prevention (CDC) in the Department of Health and Human Services (HHS) announces the withdrawal of the notice published under the same title on August 25, 2016 for public comment.

DATES: Effective September 12, 2016.

FOR FURTHER INFORMATION CONTACT: Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: On August 25, 2016 CDC published a notice in the **Federal Register** titled ''Agency Forms Undergoing Paperwork Reduction Act Review'' (Vol. 81, No. 165 FR Doc. 2016–20366, Pages 58513–58514). This notice was published prematurely and inadvertently. The notice is being withdrawn immediately for public comment. A new notice will be published at a later date for public comment.

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2016–21884 Filed 9–9–16; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–16–16ARH]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and clarity of the information to be collected; (d) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses; and (e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to *omb@cdc.gov.* Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, Washington, DC 20503 or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

Proposed Project

Poison Center Collaborations for Public Health Emergencies—NEW—National Center for Environmental Health (NCEH), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Centers for Disease Control and Prevention (CDC) is requesting a three-year approval for a new generic information collection request (Generic ICR) plan titled ''Poison Center Collaborations for Public Health Emergencies.''

CDC's key partner, the American Association of Poison Control Centers (AAPCC), is a national network of 55 poison centers working to prevent and treat poison exposures. The goal for this new Generic ICR is to create a timely mechanism to allow poison centers, in collaboration with CDC, to obtain critical exposure and health information during public health emergencies. This information is not captured during initial poison center calls about triage and treatment of potential poison exposures. Additional data collections are needed quickly to further characterize exposures, risk factors, and illnesses.

When a public health emergency of interest to CDC and AAPCC occurs, the CDC and AAPCC hold a meeting to mutually decide whether the incident needs further investigation. For a public health emergency to be selected for call-back, adverse health effects must have occurred and a response is needed to prevent further morbidity and mortality. The event must meet the criteria below:

(1) The event is a public health emergency causing adverse health effects.

(2) Timely data are urgently needed to inform rapid public health action to prevent or reduce injury, disease, or death.

(3) The event is characterized by a natural or man-made disaster, contaminated food or water, a new or existing consumer product, or an emerging public health threat.

(4) The event has resulted in calls to a poison center, and the poison center agrees to conduct the call-back data collection.

(5) The event is domestic.

(6) Data collection will be completed in 60 days or less.

Trained poison center staff will conduct the call-back telephone survey, after administering consent.