

Frank Wall Street Reform and Consumer Protection Act (Dodd-Frank Act), which permits nonbank companies that own at least one registered securities broker or dealer, and that are required by a foreign regulator or provision of foreign law to be subject to comprehensive consolidated supervision, to register with the Board and subject themselves to supervision by the Board.

Proposal to approve under OMB delegated authority the extension for three years, with revision, of the following report

1. *Report title:* Request for Proposal and Request for Price Quotations.
Agency form number: RFP and RFPQ.
OMB control number: 7100–0180.
Frequency: On occasion.
Reporters: Vendors of goods and services.

Estimated annual reporting hours:
 RFP: 17,500 hours; RFPQ: 4,400 hours;
 Subcontractor report: 50 hours.

Estimated average hours per response:
 RFP: 50 hours; RFPQ: 2 hours;
 Subcontractor report: 20 minutes.

Number of respondents: RFP: 350;
 RFPQ: 2,200; Subcontractor report: 150.

General description of report: The RFP and RFPQ are required to obtain a benefit and are authorized by Sections 10(3), 10(4), and 11(1) of the Federal Reserve Act (12 U.S.C. 243, 244, and 248(l)). With regard to the Subcontracting Report, Section 342(c) of Dodd-Frank requires the Federal Reserve to develop and implement standards and procedures to assess the diversity policies and practices in all business and activities of the agency at all levels, including procurement, insurance, and all types of contracts. (12 U.S.C. 5452(c)(1)). “Such procedure shall include a written statement, in a form and with such content as the Director [of OMWI] shall prescribe . . . that a contractor shall ensure . . . the fair inclusion of women and minorities in the workforce of the contractor and, as applicable, subcontractors.” (12 U.S.C. 5452(c)(2)).

Proposals from vendors that are not accepted and incorporated into contracts with the Federal Reserve would be protected from Freedom of Information (FOIA) disclosure by 41 U.S.C. 4702, which expressly prohibits FOIA disclosure of these proposals. Moreover, during the solicitation process vendors are permitted to mark information contained in their proposals that is proprietary or confidential with the label RESTRICTED DATA. For information so marked, the Federal Reserve also may determine on a case-by-case basis whether FOIA exemption 4, which applies to “trade secrets and

commercial or financial information,” would protect information from disclosure pursuant to a FOIA request (5 U.S.C. 552(b)(4)).

Abstract: The Federal Reserve uses the RFP and the RFPQ as appropriate to obtain competitive proposals and contracts from approved vendors of goods and services. This information collection is required to collect data on prices, specifications of goods and services, and qualifications of prospective vendors.

Current Actions: In connection with the RFP and RFPQ process, the Federal Reserve proposes to require prime contractors to submit a Subcontracting Report that would collect information about their subcontractors’ commitments toward diversity and inclusion of minority-owned and women-owned vendors in the subcontractor’s activities.

Board of Governors of the Federal Reserve System, July 20, 2015.

Robert deV. Frierson,

Secretary of the Board.

[FR Doc. 2015–18059 Filed 7–22–15; 8:45 am]

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DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0152; Docket 2015–0055; Sequence 17]

Information Collection; Service Contracting

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

ACTION: Notice of request for public comments regarding an extension to an 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 currently approved information collection requirement concerning service contracting.

DATES: Submit comments on or before September 21, 2015.

ADDRESSES: Submit comments identified by Information Collection 9000–0152, Service Contracting, 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 9000–0152, Service Contracting”. Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 9000–0152, Service Contracting” on your attached document.

- Mail: General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0152, Service Contracting.

Instructions: Please submit comments only and cite Information Collection 9000–0152, Service Contracting, in all correspondence related to this collection. All comments received will be posted without change to <http://www.regulations.gov>, including any personal and/or business confidential information provided.

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

SUPPLEMENTARY INFORMATION:

A. Purpose

The policies implemented at FAR 37.115, Uncompensated Overtime, are based on Section 834 of Public Law 101–510 (10 U.S.C. 2331). The policies require insertion of FAR provision 52.237–10, Identification of Uncompensated Overtime, in all solicitations valued above the simplified acquisition threshold, for professional or technical services to be acquired on the basis of the number of hours to be provided.

The provision requires that offerors identify uncompensated overtime hours, in excess of 40 hours per week, and the uncompensated overtime rate for direct charge Fair Labor Standards Act-exempt personnel. This permits Government contracting officers to ascertain cost realism of proposed labor rates for professional employees and discourages the use of uncompensated overtime.

B. Annual Reporting Burden

The burden placed on offerors is the time required to identify and support any hours in excess of 40 hours per week included in their proposal or subcontractor’s proposal. It is estimated that there will be 17,500 service

contracts awarded annually at \$100,000 or more, of which 65 percent or 11,375 contracts will be competitively awarded. About 7 proposals will be received for each contract award. Of the total 79,625 (11,375 × 7) proposals received, only 25 percent or 19,906 proposals are expected to include uncompensated overtime hours. It is estimated that offerors will take about 30 minutes to identify and support any hours in excess of 40 hours per week included in their proposal or subcontractor's proposal.

Number of Respondents: 19,906.

Responses Per Respondent: 1.

Total Annual Responses: 19,906.

Average Burden Hours Per Response: .5.

Total Burden Hours: 9,953.

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 Regulation (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, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202-501-4755. Please cite OMB Control No. 9000-0152, Service Contracting, in all correspondence.

Dated: July 20, 2015.

Edward Loeb,

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

[FR Doc. 2015-18077 Filed 7-22-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee: Notice of Charter Amendment

This gives notice under the Federal Advisory Committee Act (Pub. L. 92-463) of October 6, 1972, that the Clinical Laboratory Improvement Advisory Committee, Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS), has amended their charter to reduce the number of annual meetings and to change the designation of CDC, FDA and CMS from voting to non-voting ex officio members. The amended filing date is July 9, 2015.

For information, contact Nancy Anderson, Chief, Laboratory Practice Standards Branch, Division of Laboratory Programs, Standards, and Services, Center for Surveillance, Epidemiology and Laboratory Services, Office of Public Health Scientific Services, CDC, 1600 Clifton Road, NE., Mailstop F-11, Atlanta, Georgia 30329-4018; telephone (404) 498-2741.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2015-18065 Filed 7-22-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Injury Prevention and Control, (BSC, NCIPC)

Correction: This notice was published in the **Federal Register** on June 16, 2015, Volume 80, Number 115, Page 34435. The Matters For Discussion and Contact Person For More Information should read as follows:

Matters For Discussion: The BSC, NCIPC will discuss, research strategies needed to guide the Center's focus, updates on the current research

portfolio review and the Pediatric mild-Traumatic Injury Workgroup. There will be 15 minutes allotted for public comments at the end of the open session.

On the second day, the BSC, NCIPC will meet to conduct a Secondary Peer Review of extramural research grant applications received in response to four (4) Funding Opportunity Announcements (FOAs): PHS 2014002 Omnibus Solicitation of the NIH, CDC, FDA and ACF for Small Business Innovation Research Grant Applications (Parent SBIR {R42/R44}); CE15-003, Evaluating Structural, Economic, Environmental, or Policy Primary Prevention Strategies for Intimate Partner Violence and Sexual Violence; CE15-004, Evaluating Innovative and Promising Strategies to Prevent Suicide among Middle-Aged Men; and CE15-005, Research to Evaluate the CDC Heads Up Initiative in Youth Sports. Applications will be assessed as they relate to the Center's mission and programmatic balance. Recommendations from the secondary review will be voted upon and the application will be forwarded to the Center Director for consideration for funding support.

Contact Person For More Information: Arlene Greenspan, DrPH, MPH, PT, Associate Director for Science, Acting Designated Federal Officer, NCIPC, CDC, 4770 Buford Highway NE., Mailstop F-63, Atlanta, GA 30341, Telephone (770) 488-1279.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

Director, Management Analysis and Services Office Centers for Disease Control and Prevention.

[FR Doc. 2015-18064 Filed 7-22-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2015-N-2523]

Intent To Review a Study Data Reviewer's Guide Template

AGENCY: Food and Drug Administration, HHS.