U.S.C. 611a, the Federal Reserve may ''issue rules and regulations'' governing such entities ''consistent with and in furtherance of the purposes'' of that subchapter.

Because the information collection is called for in guidance and not in a statute or regulation, it is considered voluntary.

Because the information collected by the Proposed Guidance is maintained at the institutions, issues of confidentiality would not normally arise. Should the information be obtained by the Board in the course of an examination, it would be exempt from disclosure under exemption 8 of Freedom of Information Act (FOIA), 5 U.S.C. 552(b)(8). In addition, some or all of the information may be confidential commercial or financial information protected from disclosure under exemption 4 of FOIA, under the standards set forth in *National Parks & Conservation Ass'n* v. *Morton,* 498 F.2d 765 (D.C. Cir. 1974).

Abstract: The interagency guidance outlines high-level principles related to safe and sound leveraged lending activities, including underwriting considerations, assessing and documenting enterprise value, risk management expectations for credits awaiting distribution, stress testing expectations and portfolio management, and risk management expectations. This guidance applies to all financial institutions substantively engaged in leveraged lending activities supervised by the Federal Reserve, FDIC, and OCC (the Agencies).

The Agencies identified certain aspects of the proposed guidance that may constitute a collection of information. In particular, these aspects are the provisions that state a banking organization should (a) have underwriting policies for leveraged lending, including stress testing procedures for leveraged credits; (b) have risk management policies, including stress testing procedures for pipeline exposures; and (c) have policies and procedures for incorporating the results of leveraged credit and pipeline stress tests into the firm's overall stress testing framework.

Although the guidance is applicable to all institutions that originate or participate in leverage lending, due to the large exposures created by these types of loans, these credits are most likely originated primarily by larger institutions.

Current Actions: On March 15, 2016, the Board published a notice in the **Federal Register** (81 FR 13791) requesting public comment for 60 days on the proposal to extend the FR 4203 for three years without revision. The comment period for the notice expired on May 16, 2016. The Federal Reserve did not receive any comments, and the information collection will be extended as proposed.

3. Report title: Reporting, Recordkeeping, and Disclosure Requirements Associated with Regulation NN.

Agency form number: Reg NN.

OMB control number: 7100–0353.

Frequency: On occasion.

Reporters: Banking organizations seeking to engage in off-exchange transactions in foreign currency with retail customers.

Estimated annual reporting hours: 1,972 hours.

Estimated average hours per response: Reporting, 16 hours; Recordkeeping, 183 hours; Disclosure, 787 hours.

Number of respondents: 2.

General description of report: This information collection is required by the Commodity Exchange Act (7 U.S.C. Section 2(c)(2)(E)), the Federal Reserve Act (12 U.S.C. Sections 248 and 321–338), the Federal Deposit Insurance Act (12 U.S.C. Section 1818), the International Banking Act (12 U.S.C. Section 3108), and Regulation NN (12 CFR part 240). The information collection is mandatory. The reported data are regarded as confidential under the Freedom of Information Act (5 U.S.C. Section 552(b)(4)).

Abstract: The reporting requirements associated with Regulation NN are found in section 240.4; the recordkeeping requirements are found in sections 240.7, 240.9, and 240.13(a); and the disclosure requirements are found in sections 240.5, 240.6, 240.10, 240.13b–d, 240.15, and 240.16. These requirements permit banking organizations under the Federal Reserve's supervision to engage in off-exchange transactions in foreign currency with retail customers and to describe various requirements with which banking organizations must comply to conduct such transactions.

Current Actions: On March 17, 2016, the Board published a notice in the **Federal Register** (81 FR 14444) requesting public comment for 60 days on the proposal to extend the FR 4203 for three years without revision. The comment period for the notice expired on May 16, 2016. The Federal Reserve did not receive any comments, and the information collection will be extended as proposed.

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

Robert deV. Frierson,

Secretary of the Board.

[FR Doc. 2016–12604 Filed 5–26–16; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF DEFENSE

GENERAL SERVICES ADMINISTRATION

NATIONAL AERONAUTICS AND SPACE ADMINISTRATION

[OMB Control No. 9000–0173; Docket 2016–0053; Sequence 28]

Information Collection; Limitations on Pass-Through Charges

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 a previously approved information collection requirement regarding Limitations on Pass-Through Charges.

DATES: Submit comments on or before July 26, 2016.

ADDRESSES: Submit comments identified by Information Collection 9000–0173, Limitations on Pass-Through Charges 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 9000–0173, Limitations on Pass-Through Charges''. Follow the instructions provided at the ''Submit a Comment'' screen. Please include your name, company name (if any), and ''Information Collection 9000–0173, Limitations on Pass-Through Charges'' on your attached document.
- *Mail:* General Services Administration, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 9000–0173, Limitations on Pass-Through Charges.

Instructions: Please submit comments only and cite Information Collection 9000–0173, Limitations on Pass-Through Charges, 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: Mr. Michael O. Jackson, Procurement Analyst, Office of Acquisition Policy, at telephone 202–208–4949 or via email to *michaelo.jackson@gsa.gov*.

SUPPLEMENTARY INFORMATION:

A. Purpose

To enable contracting officers to verify that pass-through charges are not excessive, the provision at 52.215–22 requires offerors submitting a proposal for a contract, task order, or delivery order to provide the following information with its proposal: (1) The percent of effort the offeror intends to perform and the percent expected to be performed by each subcontractor. (2) If the offeror intends to subcontract more than 70 percent of the total cost of work to be performed—(i) The amount of the offeror's indirect costs and profit/fee applicable to the work to be performed by the subcontractor(s); and (ii) A description of the value added by the offeror as related to the work to be performed by the subcontractor(s). (3) If any subcontractor intends to subcontract to a lower-tier subcontractor more than 70 percent of the total cost of work to be performed under its subcontract—(i) The amount of the subcontractor's indirect costs and profit/fee applicable to the work to be performed by the lower-tier subcontractor(s); and (ii) A description of the value added by the subcontractor as related to the work to be performed by the lower-tier subcontractor(s).

B. Annual Reporting Burden

Respondents: 4,638.
Responses per Respondent: 8.7.
Total Responses: 40,347.
Hours per Response: 2.
Total Burden Hours: 80,694.
Frequency of Collection: On Occasion.
Affected Public: Businesses or other for-profit and not-for-profit institutions.

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, Regulatory Secretariat Division (MVCB), 1800 F Street NW., Washington, DC 20405, telephone 202–501–4755. Please cite OMB Control No. 9000–0173, Limitations on Pass-Through Charges, in all correspondence.

Dated: May 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. 2016–12554 Filed 5–26–16; 8:45 am]

BILLING CODE 6820–EP–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Agency for Healthcare Research and Quality, HHS.

ACTION: Notice.

SUMMARY: This notice announces the intention of the Agency for Healthcare Research and Quality (AHRQ) to request that the Office of Management and Budget (OMB) approve the proposed information collection project: ''*Eisenberg Center Voluntary Customer Survey Generic Clearance.*'' In accordance with the Paperwork Reduction Act, 44 U.S.C. 3501–3521, AHRQ invites the public to comment on this proposed information collection.

DATES: Comments on this notice must be received by July 26, 2016.

ADDRESSES: Written comments should be submitted to: Doris Lefkowitz, Reports Clearance Officer, AHRQ, by email at *doris.lefkowitz@AHRQ.hhs.gov.*

Copies of the proposed collection plans, data collection instruments, and specific details on the estimated burden can be obtained from the AHRQ Reports Clearance Officer.

FOR FURTHER INFORMATION CONTACT: Doris Lefkowitz, AHRQ Reports Clearance Officer, (301) 427–1477, or by email at *doris.lefkowitz@AHRQ.hhs.gov.*

SUPPLEMENTARY INFORMATION:

Proposed Project

Eisenberg Center Voluntary Customer Survey Generic Clearance

The Agency for Healthcare Research and Quality (AHRQ) requests that the Office of Management and Budget (OMB) renew under the Paperwork Reduction Act of 1995 AHRQ's Generic Clearance to collect information from users of work products and services produced by AHRQ's John M. Eisenberg Center for Clinical Decisions and Communications Science (Eisenberg Center). The Eisenberg Center is an innovative effort aimed at improving communication of findings to a variety of audiences (''customers''), including consumers, clinicians, and health care policymakers. The Eisenberg Center compiles research results into a variety of useful formats for stakeholders.

This effort has the following goals:

(1) Conduct research into effective communication of research findings in order to improve the usability and rapid incorporation of findings into medical practice and decision making.

(2) Conduct research into effective strategies for disseminating evidence-based products, tools, and resources to consumers, clinicians, and other health care professionals, and policymakers.

(3) Evaluate outcomes reported by clinicians and other health care professionals resulting from participation in continuing medical education (CME) initiatives and activities.

(4) Conduct research into factors associated with successful collaboration between AHRQ and partnering institutions and organizations in synthesizing, translating, and disseminating evidence-based research.

Clearance is being requested to cover a three-year period in which differing numbers of products and research activities may be conducted during each year. The collections proposed include activities to assist in the development of materials to be disseminated through the Eisenberg Center and to provide feedback to AHRQ on the extent to which these products meet customer needs. These materials include summary documents that summarize and translate the findings of research reports for various decision-making audiences, such as consumers, clinicians, and policymakers. The summaries are designed to help these decision makers use research evidence to maximize the benefits of health care, minimize harm, and optimize the use of health care resources. In addition, each year, a unique research project will be undertaken to study successful approaches to disseminating AHRQ