

Abstract: The board of directors of each state member bank must designate a security officer to assume the responsibility for the development and administration of a written security program within 180 days of opening for business. Each state member bank must develop and implement a written security program for the bank's main office and branches and maintain it in the bank's records. The designated security officer must report at least annually to the bank's board of directors on the implementation, administration, and effectiveness of the written security program. There is no formal reporting form and the information is not submitted to the Federal Reserve.

Legal authorization and confidentiality: This recordkeeping requirement is mandatory pursuant to section 3 of the Bank Protection Act (12 U.S.C. 1882(a)) and Regulation H (12 CFR 208.61). Because written security programs are maintained at state member banks, no issue of confidentiality under the Freedom of Information Act (FOIA) normally arises. However, copies of such documents included in examination work papers would, in such form, be confidential pursuant to exemption 8 of FOIA (5 U.S.C. 552(b)(8)). In addition, the records may also be exempt from disclosure under exemption 4 of FOIA (5 U.S.C. 552(b)(4)).

Current Actions: On February 23, 2016, the Board published a notice in the **Federal Register** (81 FR 8958) requesting public comment for 60 days on the proposal to extend the FR 4004 for three years without revision. The comment period for the notice expired on April 25, 2016. The Federal Reserve did not receive any comments, and the information collection will be extended as proposed.

2. Report title: Risk-Based Capital Guidelines: Market Risk.

Agency form number: FR 4201.

OMB control number: 7100-0314.

Frequency: Varied—some requirements are done at least quarterly and some at least annually.

Reporters: State member banks, bank holding companies, and certain savings and loan holding companies.

Number of respondents: 28.

Estimated burden per respondent: 1,964 hours.

Total estimated annual burden: 54,992 hours.

Abstract: The market risk rule is an important component of the Board's regulatory capital framework (12 CFR 217) that requires banking organizations to measure and hold capital to cover their exposure to market risk. On July 2, 2013, the Federal Reserve adopted a

revised regulatory capital framework, including the market risk rule, which was expanded to include certain savings and loan holding companies. The information-collection requirements in the market risk rule provide the most current statistical data available to identify areas of market risk on which to focus for onsite and offsite examinations and allow the Federal Reserve to assess and monitor the levels and components of each reporting institution's risk-based capital requirements for market risk and the adequacy of the institution's capital under the market risk rule. The reporting, recordkeeping, and disclosure requirements are found in sections 12 CFR 217.203–217.210, and 217.212. These requirements enhance risk sensitivity and introduce requirements for public disclosure of certain qualitative and quantitative information about a financial institution's market risk. There are no required reporting forms associated with this information collection.

Legal authorization and confidentiality: The FR 4201 is authorized under 12 U.S.C. 324, 1844(c), and 1467a(b)(2)(A). Information collected pursuant to the reporting requirements of the FR 4201 (specifically, information related to seeking regulatory approval for the use of certain incremental and comprehensive risk models and methodologies under sections 217.208 and 217.209) is exempt from disclosure pursuant to exemption (b)(8) of FOIA (5 U.S.C. 552(b)(8)), and exemption (b)(4) of FOIA (5 U.S.C. 552(b)(4)). Exemption (b)(8) applies because the reported information is contained in or related to examination reports. Exemption (b)(4) applies because the information provided to obtain regulatory approval of the incremental or comprehensive risk models is confidential business information the release of which could cause substantial competitive harm to the reporting company. The recordkeeping requirements of the FR 4201 require banking organizations to maintain documentation regarding certain policies and procedures, trading and hedging strategies, and internal models. These documents would remain on the premises of the banking organizations and accordingly would not generally be subject to a FOIA request. To the extent these documents are provided to the regulators, they would be exempt under exemption (b)(8), and may be exempt under exemption (b)(4). Exemption (b)(4) protects from disclosure "trade secrets and commercial or financial information

obtained from a person and privileged or confidential." The disclosure requirements of the FR 4201 do not raise any confidentiality issues because they require banking organizations to make certain information public.

Current Actions: On February 23, 2016, the Board published a notice in the **Federal Register** (81 FR 8958) requesting public comment for 60 days on the proposal to extend the FR 4201 for three years without revision. The comment period for the notice expired on April 25, 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 23, 2016.

Robert deV. Frierson,
Secretary of the Board.

[FR Doc. 2016-12470 Filed 5-25-16; 8:45 am]

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GENERAL SERVICES ADMINISTRATION

[OMB Control No. 3090-0302; Docket 2016-0001; Sequence 2]

General Services Administration Acquisition Regulation; Submission for OMB Review; Modifications 552.238-81

AGENCY: Office of Acquisition Policy, General Services Administration (GSA).

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 information collection requirement regarding the Modifications clause.

DATES: Submit comments on or before: June 27, 2016.

FOR FURTHER INFORMATION CONTACT: Ms. Dana Munson, Procurement Analyst, General Services Acquisition Policy Division, GSA, 202-357-9652 or email dana.munson@gsa.gov.

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 inputting “Information Collection 3090–0302, Modifications,” under the heading “Enter Keyword or ID” and selecting “Search”. Select the link “Submit a Comment” that corresponds with “Information Collection 3090–0302, Modifications.” Follow the instructions provided at the “Submit a Comment” screen. Please include your name, company name (if any), and “Information Collection 3090–0302, Modifications,” on your attached document.

- **Mail:** General Services Administration, Regulatory Secretariat (MVCB), 1800 F Street NW., Washington, DC 20405. ATTN: Ms. Flowers/IC 3090–0302, Modifications.

Instructions: Please submit comments only and cite Information Collection 3090–0302, Modifications, 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).

SUPPLEMENTARY INFORMATION:

A. Purpose

The General Services Administration Acquisition Regulation (GSAR) clause 552.238–81 Modifications requires vendors to request a contract modification by submitting a request to the Contracting Officer for approval, except for electronic File updates. At a minimum, every request shall describe the proposed change(s) and provide the rationale for the requested change(s). A notice was published in the **Federal Register** at 81 FR 12731 on March 10, 2016. No comments were received.

B. Annual Reporting Burden

Respondents: 19,500.
Responses per Respondent: 2.
Total Responses: 39,000.
Hours per Response: 5.
Total Burden Hours: 195,000.

C. Public Comments

Public comments are particularly invited on: Whether this collection of information is necessary 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.

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. 3090–0302, “Modifications” in all correspondence.

Jeffrey A. Koses,

Director, Office of Acquisition Policy, Office of Government-wide Policy.

[FR Doc. 2016–12370 Filed 5–25–16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–855(A, B, I)]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

ACTION: Notice.

SUMMARY: The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS’ intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (PRA), federal agencies are required to publish a notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, and to allow a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency’s functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments on the collection(s) of information must be received by the OMB desk officer by June 27, 2016.

ADDRESSES: When commenting on the proposed information collections, please reference the document identifier or OMB control number. To be assured consideration, comments and

recommendations must be received by the OMB desk officer via one of the following transmissions: OMB, Office of Information and Regulatory Affairs, Attention: CMS Desk Officer, Fax Number: (202) 395–5806 *OR* Email: OIRA_submission@omb.eop.gov.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS’ Web site address at <http://www.cms.hhs.gov/PaperworkReductionActof1995>.
2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov.
3. Call the Reports Clearance Office at (410) 786–1326.

FOR FURTHER INFORMATION CONTACT: Reports Clearance Office at (410) 786–1326.

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice that summarizes the following proposed collection(s) of information for public comment:

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Medicare Enrollment Application; *Use:* The primary function of the CMS–855 Medicare enrollment application is to gather information from a provider or supplier that tells us who it is, whether it meets certain qualifications to be a health care provider or supplier, where it practices or renders its services, the identity of the owners of the enrolling entity, and other information necessary to establish correct claims payments. No comments were received during the 60-day comment period (April 1, 2016 (81 FR 18855)). *Form Number:* CMS–855(A,