requirements of CFPB's Regulation B is mandatory. Because the recordkeeping and disclosure requirements of the CFPB's Regulation B require creditors to retain their own records and to make certain disclosures to customers, the Freedom of Information Act (FOIA) would only be implicated if the Board's examiners retained a copy of this information as part of an examination of a bank. Records obtained as a part of an examination or supervision of a bank are exempt from disclosure under FOIA exemption (b)(8), for examination material (5 U.S.C. 552(b)(8)). In addition, the records may also be exempt under FOIA exemption (b)(4) or (b)(6). Records would be exempt under (b)(4) if the records contained ''trade secrets and commercial or financial information obtained from a person [that is] privileged or confidential'' and the disclosure of the information is likely to cause substantial harm to the competitive position of the respondents (5 U.S.C. 552(b)(4)). Records would be exempt under (b)(6) if the records contained personal information, the disclosure of which would ''constitute a clearly unwarranted invasion of personal privacy'' (5 U.S.C. 552(b)(6)).

Current actions: On April 13, 2018, the Board published a notice in the **Federal Register** (83 FR 16098) requesting public comment for 60 days on the extension, without revision, of the FR B. The comment period for this notice expired on June 12, 2018. The Board received one comment letter that addressed matter outside the scope of this proposal.

Board of Governors of the Federal Reserve System, June 28, 2018.

Michele Taylor Fennell,

Assistant Secretary of the Board.

[FR Doc. 2018–14305 Filed 7–2–18; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[CDC–2018–0006; Docket Number NIOSH–306]

Final National Occupational Research Agenda for Services

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 availability.

SUMMARY: NIOSH announces the availability of the final *National Occupational Research Agenda for Services.*

DATES: The final document was published on June 26, 2018.

ADDRESSES: The document may be obtained at the following link: *https://www.cdc.gov/niosh/nora/sectors/serv/agenda.html.*

FOR FURTHER INFORMATION CONTACT: Emily Novicki, M.A., M.P.H, (*NORACoordinator@cdc.gov*), National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention, Mailstop E–20, 1600 Clifton Road NE, Atlanta, GA 30329, phone (404) 498–2581 (not a toll free number).

SUPPLEMENTARY INFORMATION: On January 29, 2018, NIOSH published a request for public review in the **Federal Register** [83 FR 4058] of the draft version of the *National Occupational Research Agenda for Services.* All comments received were reviewed and addressed where appropriate.

John J. Howard,

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

[FR Doc. 2018–14227 Filed 7–2–18; 8:45 am]

BILLING CODE 4163–19–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10673]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services.

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 (the PRA), federal agencies are required to publish 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 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

DATES: Comments must be received by September 4, 2018.

ADDRESSES: When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to *http://www.regulations.gov.* Follow the instructions for ''Comment or Submission'' or ''More Search Options'' to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number ___, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

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' website address at *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html.*

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: William Parham at (410) 786–4669.

SUPPLEMENTARY INFORMATION:

Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement and associated materials (see **ADDRESSES**).

CMS–10673 Medicare Advantage Qualifying Payment Arrangement Incentive (MAQI) Demonstration

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management