Type of burden	Number of respondents	Estimated time per response	Frequency of response	Total estimated annual burden (hours)
Recordkeeping	1	166	On Occasion	166
Disclosure	1	1,332	On Occasion	1,332
Total Estimated Annual Burden				1,514

General Description of Collection: This information collection implements section 742(c)(2) of the Dodd-Frank Act (7 U.S.C. 2(c)(2)(E) and FDIC regulations governing retail foreign exchange transactions as set forth at 12 CFR part 349, subpart B. The regulation allows banking organizations under FDIC supervision to engage in off-exchange transactions in foreign currency with retail customers provided they comply with various reporting, recordkeeping and third-party disclosure requirements specified in the rule. If an institution elects to conduct such transactions, compliance with the information collection is mandatory.

Reporting Requirements—Part 349, subpart B requires that, prior to initiating a retail foreign exchange business; a banking institution must provide the FDIC with a notice certifying that the institution has written policies and procedures, and risk measurement and management systems and controls in place to ensure that retail foreign exchange transactions are conducted in a safe and sound manner. The institution must also provide information about it intends to manage customer due diligence, new product approvals and haircuts applied to noncash margin.

Recordkeeping Requirements—Part 349 subpart B requires that institutions engaging in retail foreign exchange transactions keep full, complete and systematic records of account, financial ledger, transaction, memorandum orders and post execution allocations of bunched orders. In addition, institutions are required to maintain records regarding their ratio of profitable accounts, possible violations of law, records of noncash margin and monthly statements and confirmations issued.

Disclosure Requirements—The regulation requires that, before opening an account that will engage in retail foreign exchange transactions, a banking institution must obtain from each retail foreign exchange customer an acknowledgement of receipt and understanding of a written disclosure specified in the rule and of disclosures about the banking institution's fees and other charges and of its profitable accounts ratio. The institution must also provide monthly statements to each retail foreign exchange customer and must send confirmation statements following every transaction. The customer dispute resolution provisions of the regulation require certain endorsements, acknowledgements and signature language as well as the timely provision of a list of persons qualified to handle a customer's request for arbitration.

There is no change in the method or substance of the collection. At present no FDIC-supervised institution is engaging in activities that would make them subject to the information collection requirements. FDIC originally estimated that 3 institutions would be impacted by the rule. The agency is reducing the estimated number of respondents to one (1) as a placeholder in case an institution elects to engage in covered activities in the future. There has been no change in the frequency of response or in the estimated number of hours required to respond. Because of the reduction in the estimated number of respondents from three (3) to one (1), the estimated annual burden has decreased.

Request for Comment

Comments are invited on: (a) Whether the collections of information are necessary for the proper performance of the FDIC's functions, including whether the information has practical utility; (b) the accuracy of the estimates of the burden of the information collections, including the validity of the methodology and assumptions used; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collections of information on respondents, including through the use of automated collection techniques or other forms of information technology. All comments will become a matter of public record.

Dated at Washington, DC, this 21st day of July, 2017.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2017–15711 Filed 7–26–17; 8:45 am]

BILLING CODE 6714–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Bank Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Bank Holding Company Act of 1956 (12 U.S.C. 1841 *et seq.*) (BHC Act), Regulation Y (12 CFR part 225), and all other applicable statutes and regulations to become a bank holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a bank or bank holding company and all of the banks and nonbanking companies owned by the bank holding company, including the companies listed below.

The applications listed below, as well as other related filings required by the Board, are available for immediate inspection at the Federal Reserve Bank indicated. The applications will also be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the standards enumerated in the BHC Act (12 U.S.C. 1842(c)). If the proposal also involves the acquisition of a nonbanking company, the review also includes whether the acquisition of the nonbanking company complies with the standards in section 4 of the BHC Act (12 U.S.C. 1843). Unless otherwise noted, nonbanking activities will be conducted throughout the United States.

Unless otherwise noted, comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than August 23, 2017.

A. Federal Reserve Bank of St. Louis (David L. Hubbard, Senior Manager) P.O. Box 442, St. Louis, Missouri 63166–2034. Comments can also be sent electronically to *Comments.applications@stls.frb.org:*

1. *Ozarks Heritage Financial Group, Inc., Gainesville, Missouri;* and its top tier holding company Century Bancshares, Inc., Gainesville, Missouri, to acquire 100 percent of the voting shares of Financial Enterprises, Inc., Clinton, Missouri, and thereby indirectly acquire First National Bank of Clinton, Clinton, Missouri.

Board of Governors of the Federal Reserve System, July 24, 2017.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2017–15871 Filed 7–26–17; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Notice of Proposals To Engage in or To Acquire Companies Engaged in Permissible Nonbanking Activities

The companies listed in this notice have given notice under section 4 of the Bank Holding Company Act (12 U.S.C. 1843) (BHC Act) and Regulation Y, (12 CFR part 225) to engage *de novo,* or to acquire or control voting securities or assets of a company, including the companies listed below, that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) or that the Board has determined by Order to be closely related to banking and permissible for bank holding companies. Unless otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act.

Unless otherwise noted, comments regarding the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than August 11, 2017.

A. Federal Reserve Bank of New York (Ivan Hurwitz, Vice President) 33 Liberty Street, New York, New York 10045–0001. Comments can also be sent electronically to *Comments.applications@ny.frb.org:*

1. *China Merchants Group Limited, Hong Kong Special Administrative Region, the People's Republic of China;* to engage de novo in extending credit and servicing loans and the leasing of personal property through CIMC Leasing USA Inc., Oakbrook Terrace, Illinois, pursuant to sections 225.28(b)(1) and 225.28(b)(3) of Regulation Y.

Board of Governors of the Federal Reserve System, July 24, 2017.

Yao-Chin Chao,

Assistant Secretary of the Board.

[FR Doc. 2017–15872 Filed 7–26–17; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Supplemental Evidence and Data Request on Nonsurgical Treatments for Urinary Incontinence in Adult Women: A Systematic Review Update

AGENCY: Agency for Healthcare Research and Quality (AHRQ), HHS.

ACTION: Request for Supplemental Evidence and Data Submissions.

SUMMARY: The Agency for Healthcare Research and Quality (AHRQ) is seeking scientific information submissions from the public. Scientific information is being solicited to inform our review of *Nonsurgical Treatments for Urinary Incontinence in Adult Women: A Systematic Review Update,* which is currently being conducted by the AHRQ's Evidence-based Practice Centers (EPC) Program. Access to published and unpublished pertinent scientific information will improve the quality of this review.

DATES: Submission Deadline on or before August 28, 2017.

ADDRESSES:

Email submissions: SEADS@epc-src.org.

Print submissions: Mailing Address: Portland VA Research Foundation, Scientific Resource Center, ATTN: Scientific Information Packet Coordinator, P.O. Box 69539, Portland, OR 97239.

Shipping Address (FedEx, UPS, etc.): Portland VA Research Foundation, Scientific Resource Center, ATTN: Scientific Information Packet Coordinator, 3710 SW., U.S. Veterans Hospital Road, Mail Code: R&D 71, Portland, OR 97239.

FOR FURTHER INFORMATION CONTACT: Ryan McKenna, Telephone: 503–220–8262 ext. 51723 or Email: *SEADS@epc-src.org.*

SUPPLEMENTARY INFORMATION: The Agency for Healthcare Research and Quality has commissioned the Evidence-based Practice Centers (EPC) Program to complete a review of the evidence for *Nonsurgical Treatments for Urinary Incontinence in Adult Women: A Systematic Review Update.* AHRQ is conducting this systematic review pursuant to Section 902(a) of the Public Health Service Act, 42 U.S.C. 299a(a).

The EPC Program is dedicated to identifying as many studies as possible that are relevant to the questions for each of its reviews. In order to do so, we are supplementing the usual manual and electronic database searches of the literature by requesting information from the public (*e.g.,* details of studies conducted). We are looking for studies that report on *Nonsurgical Treatments for Urinary Incontinence in Adult Women: A Systematic Review Update,* including those that describe adverse events. The entire research protocol, including the key questions, is also available online at: *https://effectivehealthcare.ahrq.gov/index.cfm/search-for-guides-reviews-and-reports/?pageaction=displayproduct&productid=2479.*

This is to notify the public that the EPC Program would find the following information on *Nonsurgical Treatments for Urinary Incontinence in Adult Women: A Systematic Review Update* helpful:

- A list of completed studies that your organization has sponsored for this indication. In the list, please *indicate whether results are available on ClinicalTrials.gov along with the ClinicalTrials.gov trial number.*
- *For completed studies that do not have results on ClinicalTrials.gov,* please provide a summary, including the following elements: Study number, study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, primary and secondary outcomes, baseline characteristics, number of patients screened/eligible/enrolled/lost to follow-up/withdrawn/analyzed, effectiveness/efficacy, and safety results.
- A list of ongoing studies that your organization has sponsored for this indication. In the list, please provide the ClinicalTrials.gov trial number or, if the trial is not registered, the protocol for the study including a study number, the study period, design, methodology, indication and diagnosis, proper use instructions, inclusion and exclusion criteria, and primary and secondary outcomes.
- Description of whether the above studies constitute ALL Phase II and above clinical trials sponsored by your organization for this indication and an index outlining the relevant information in each submitted file.

Your contribution will be very beneficial to the EPC Program. Materials submitted must be publicly available or able to be made public. Materials that are considered confidential; marketing materials; study types not included in the review; or information on indications not included in the review cannot be used by the EPC Program. This is a voluntary request for information, and all costs for complying with this request must be borne by the submitter.