

*Legal authorization and confidentiality:* The Board is authorized to collect the information in the FR 3075 from bank holding companies (and their subsidiaries) under section 5(c) of the Bank Holding Company Act (12 U.S.C. 1844(c)); from savings and loan holding companies under section 10(b)(2) of the Home Owners Loan Act (12 U.S.C. 1467a(b)(2)); from non-BHC/SLHC systemically important financial institutions under section 161(a) of the Dodd-Frank Act (12 U.S.C. 5361(a)); from the combined domestic operations of certain foreign banking organizations under section 8(a) of the International Banking Act of 1978 (12 U.S.C. 3106(a)) and section 5(c) of the Bank Holding Company Act (12 U.S.C. 1844(c)); from state member banks under section 9 of the Federal Reserve Act (12 U.S.C. 324); from Edge and agreement corporations under sections 25 and 25A of the Federal Reserve Act (12 U.S.C. 602 and 625) and from U.S. branches and agencies of foreign banks under section 7(c)(2) of the International Banking Act of 1978 (12 U.S.C. 3105(c)(2)) and under section 7(a) of the Federal Deposit Insurance Act (12 U.S.C. 1817(a)).

These surveys would be conducted on a voluntary basis. The confidentiality of information provided by respondents to the FR 3075 surveys will be determined on a case-by-case basis depending on the type of information provided for a particular survey. Depending upon the survey questions, confidential treatment may be warranted under exemptions 4, 6, and 8 of the Freedom of Information Act (5 U.S.C. 552(b)(4), (6), and (8)).

*Consultation outside the agency:* Surveys conducted under the FR 3075 may include data collections sponsored by bodies such as the BCBS and the FSB.

Board of Governors of the Federal Reserve System, February 12, 2018.

**Ann E. Misback,**  
*Secretary of the Board.*

[FR Doc. 2018-03209 Filed 2-15-18; 8:45 am]

**BILLING CODE 6210-01-P**

## FEDERAL RESERVE SYSTEM

### Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the notices are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The notices are available for immediate inspection at the Federal Reserve Bank indicated. The notices also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than March 7, 2018.

*A. Federal Reserve Bank of Kansas City* (Dennis Denney, Assistant Vice President) 1 Memorial Drive, Kansas City, Missouri 64198-0001:

1. *Robert McMahan, John McMahan, and the McMahan Family Trust, Edythe McMahan, trustee, all of Altus, Oklahoma;* to retain voting shares of First Altus Bancorp, Inc., Altus, Oklahoma, and for approval to join members of the McMahan family group, which owns shares of First Altus Bancorp, Altus, Oklahoma, and owns shares of Frazer Bank, Altus, Oklahoma.

2. *Tanner Johnson, Courtland, Kansas; Travis Johnson, Wichita, Kansas; and Tara Renze, Leawood, Kansas, individually and as members of the Johnson Family Group;* to retain voting shares of Swedish-American Bancshares, Inc., and thereby indirectly retain shares of Swedish-American State Bank, both of Courtland, Kansas.

Board of Governors of the Federal Reserve System, February 13, 2018.

**Ann Misback,**  
*Secretary of the Board.*

[FR Doc. 2018-03262 Filed 2-15-18; 8:45 am]

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## FEDERAL TRADE COMMISSION

### Agency Information Collection Activities; Submission for OMB Review; Comment Request; Extension

**AGENCY:** Federal Trade Commission ("FTC" or "Commission").

**ACTION:** Notice.

**SUMMARY:** The information collection requirements described below will be submitted to the Office of Management and Budget ("OMB") for review, as required by the Paperwork Reduction Act ("PRA"). The FTC intends to ask OMB to extend for an additional three years the current PRA clearance for the FTC's enforcement of the information collection requirements in its Fair Packaging and Labeling Act regulations ("FPLA Rules"). That clearance expires on April 30, 2018.

**DATES:** Comments must be filed by March 19, 2018.

**ADDRESSES:** Interested parties may file a comment online or on paper, by

following the instructions in the Request for Comment part of the **SUPPLEMENTARY INFORMATION** section below. Write "FPLA Rules, PRA Comment, P074200" on your comment and file your comment online at <https://ftcpublic.commentworks.com/ftc/fplaregspra2> by following the instructions on the web-based form. If you prefer to file your comment on paper, mail your comment to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC-5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024.

**FOR FURTHER INFORMATION CONTACT:** Megan Gray, Attorney, Division of Enforcement, Bureau of Consumer Protection, (202) 326-3408, [mgray@ftc.gov](mailto:mgray@ftc.gov), 600 Pennsylvania Ave. NW, Room CC-9541, Washington, DC 20580.

**SUPPLEMENTARY INFORMATION:** On December 6, 2017, the FTC sought public comment on the information collection requirements associated with the FPLA Rules (December 6, 2017 Notice <sup>1</sup>), 16 CFR parts 500-503 (OMB Control Number 3084-0110).<sup>2</sup> No relevant comments were received. Pursuant to the OMB regulations, 5 CFR part 1320, that implement the PRA, 44 U.S.C. 3501 *et seq.*, the FTC is providing this second opportunity for public comment while seeking OMB approval to renew the pre-existing clearance for the Rules.

### Burden Statement

As detailed in the December 6, 2017 Notice, the FTC estimates cumulative annual burden on affected entities to be 8,084,250 hours and \$199,680,975 in labor costs. Commission staff believes that the FPLA Rules impose negligible capital or other non-labor costs, as the affected entities are likely to have the necessary supplies and/or equipment already (*e.g.*, offices and computers) to implement the packaging and labeling disclosure requirements under the FPLA Rules.

<sup>1</sup> 82 FR 57599.

<sup>2</sup> Section 4 of the FPLA specifically requires packages or labels to be marked with: (1) A statement of identity; (2) a net quantity of contents disclosure; and (3) the name and place of business of the company responsible for the product. The FPLA Rules, 16 CFR parts 500-503, specify how manufacturers, packagers, and distributors of "consumer commodities" must do this.

### Request for Comment

You can file a comment online or on paper. For the FTC to consider your comment, we must receive it on or before March 19, 2018. Write “FPLA Rules, PRA Comment, P074200” on your comment. Your comment—including your name and your state—will be placed on the public record of this proceeding, including, to the extent practicable, on the public Commission website, at <http://www.ftc.gov/os/publiccomments.shtm>.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online, or to send them to the Commission by courier or overnight service. To make sure that the Commission considers your online comment, you must file it at <https://ftcpublic.commentworks.com/ftc/fplaregspra2>, by following the instructions on the web-based form. When this Notice appears at <http://www.regulations.gov/#/home>, you also may file a comment through that website.

If you file your comment on paper, write “FPLA Rules, PRA Comment, P074200” on your comment and on the envelope, and mail it to the following address: Federal Trade Commission, Office of the Secretary, 600 Pennsylvania Avenue NW, Suite CC–5610 (Annex J), Washington, DC 20580, or deliver your comment to the following address: Federal Trade Commission, Office of the Secretary, Constitution Center, 400 7th Street SW, 5th Floor, Suite 5610 (Annex J), Washington, DC 20024. If possible, submit your paper comment to the Commission by courier or overnight service.

Because your comment will be placed on the publicly accessible FTC website at <https://www.ftc.gov/>, you are solely responsible for making sure that your comment does not include any sensitive or confidential information. In particular, your comment should not include any sensitive personal information, such as your or anyone else’s Social Security number; date of birth; driver’s license number or other state identification number, or foreign country equivalent; passport number; financial account number; or credit or debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, such as medical records or other individually identifiable health information. In addition, your comment should not include any “trade secret or any

commercial or financial information which \* \* \* is privileged or confidential”—as provided by Section 6(f) of the FTC Act, 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16 CFR 4.10(a)(2)—including in particular competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns, devices, manufacturing processes, or customer names.

Comments containing material for which confidential treatment is requested must be filed in paper form, must be clearly labeled “Confidential,” and must comply with FTC Rule 4.9(c). In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c). Your comment will be kept confidential only if the General Counsel grants your request in accordance with the law and the public interest. Once your comment has been posted on the public FTC website—as legally required by FTC Rule 4.9(b)—we cannot redact or remove your comment from the FTC website, unless you submit a confidentiality request that meets the requirements for such treatment under FTC Rule 4.9(c), and the General Counsel grants that request.

The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before March 19, 2018. For information on the Commission’s privacy policy, including routine uses permitted by the Privacy Act, see <https://www.ftc.gov/site-information/privacy-policy>. For supporting documentation and other information underlying the PRA discussion in this Notice, see <http://www.reginfo.gov/public/jsp/PRA/pradashboard.jsp>.

Comments on the information collection requirements subject to review under the PRA should additionally be submitted to OMB. If sent by U.S. mail, they should be addressed to Office of Information and Regulatory Affairs, Office of Management and Budget, Attention: Desk Officer for the Federal Trade Commission, New Executive Office Building, Docket Library, Room 10102, 725 17th Street NW, Washington, DC 20503. Comments sent to OMB by U.S. postal mail, however, are subject to delays due to heightened security precautions. Thus, comments instead

can also be sent by email to [wliberante@omb.eop.gov](mailto:wliberante@omb.eop.gov).

David C. Shonka,  
Acting General Counsel.

[FR Doc. 2018–03289 Filed 2–15–18; 8:45 am]

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## 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:** Through this Request for Information (RFI), the Agency for Healthcare Research and Quality (AHRQ) is seeking information from the public, hospitals and other health care organizations, clinicians, quality improvement experts, researchers, and quality measure developers about current use of the AHRQ Quality Indicators (AHRQ QIs) for quality improvement efforts. AHRQ recognizes that the AHRQ QIs have been adopted for other uses, but for the purpose of this RFI, the Agency is specifically seeking information about quality improvement initiatives such as those that seek to: Improve clinical practice (e.g., adherence to guidelines, coordination of care); improve patient safety or reduce harm; address disparities in health or care; improve prevention practices; and collaborate with community groups to improve health or care. AHRQ is also seeking information about the ways in which the Agency can increase use of the AHRQ QI measures for quality improvement, for example by refining measures, summarizing best practices, creating training materials, developing standardized metrics, and/or convening learning networks. To learn more about the AHRQ QIs, visit <https://www.qualityindicators.ahrq.gov/>.

**DATES:** Comments on this notice must be received by the deadline on or before March 8, 2018.

**ADDRESSES:** Written comments should be submitted to: Maushami DeSoto, Ph.D., MHA, Health Scientist Administrator, Center for Delivery Organization and Markets, Agency for Healthcare Research and Quality, 5600 Fishers Lane, Rockville, MD 20857, or by email at [Maushami.Desoto@ahrq.hhs.gov](mailto:Maushami.Desoto@ahrq.hhs.gov).