**FEDERAL ELECTION COMMISSION**

**Sunshine Act Meeting**

**AGENCY:** Federal Election Commission.

**DATE AND TIME:** Tuesday, March 7, 2017 at 10:00 a.m. and its Continuation at the Conclusion of the open meeting on March 9, 2017.

**PLACE:** 999 E Street NW., Washington, DC.

**STATUS:** This meeting will be closed to the public.

**ITEMS TO BE DISCUSSED:** Compliance matters pursuant to 52 U.S.C. 30109.

Investigatory records compiled for law enforcement purposes and production would disclose investigative techniques.

Information the premature disclosure of which would be likely to have a considerable adverse effect on the implementation of a proposed Commission action.

Matters concerning participation in civil actions or proceedings or arbitration.

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**PERSON TO CONTACT FOR INFORMATION:**

Judith Ingram, Press Officer, Telephone: (202) 694–1220.

Dayna C. Brown, Secretary and Clerk of the Commission.

[FR Doc. 2017–04196 Filed 3–1–17; 11:15 am]

**BILLING CODE 6715–01–P**

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**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 March 27, 2017.

**A. Federal Reserve Bank of Cleveland**

(Nadine Wallman, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to

Comments.applications@clev.frb.org:


**B. 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. Southern Bancorp, Indiana, Arkadelphia, Arkansas; to acquire 100 percent of the voting shares of Farmers Bank, Hamburg, Arkansas.

Board of Governors of the Federal Reserve System, February 27, 2017.

**FOR FURTHER INFORMATION CONTACT:**

Katherine White, Attorney, Division of Privacy and Identity Protection, Bureau of Consumer Protection, (202) 326–2878, 600 Pennsylvania Ave. NW., Room CC–8232, Washington D.C. 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.

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**FEDERAL TRADE COMMISSION**

**Agency Information Collection Activities; Proposed Collection; Comment Request; Extension**

**AGENCY:** Federal Trade Commission (“FTC” or “Commission”).

**ACTION:** Notice.

**SUMMARY:** The FTC intends to ask the Office of Management and Budget (“OMB”) to extend for an additional three years the current Paperwork Reduction Act (“PRA”) clearance for the FTC’s enforcement of the information collection requirements in its “Fair Credit Reporting Risk-Based Pricing Regulations” (“RBPR Rule”), which applies to certain motor vehicle dealers, and its shared enforcement with the Consumer Financial Protection Bureau (“CFPB”) of the risk-based pricing provisions (subpart H) of the CFPB’s Regulation V regarding other entities. That clearance expires on July 1, 2017.

**DATES:** Comments must be filed by May 2, 2017.

**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 “RBPR Rule, PRA Comment, P145403,” on your comment and file your comment online at https://ftcpublic.commentworks.com/ftc/rbprulepra 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:**


**SUPPLEMENTARY INFORMATION:** On July 21, 2010, President Obama signed into law the Dodd-Frank Wall Street Reform and Consumer Protection Act (“Dodd-Frank Act”).

The Dodd-Frank Act substantially changed the federal legal framework for financial services providers. Among the changes, the Dodd-Frank Act transferred to the CFPB most of the FTC’s rulemaking authority for the risk-based pricing provisions of the Fair Credit Reporting Act (“FCRA”), on July 21, 2011.

The FTC retains rulemaking authority for the RBPR Rule solely for motor vehicle dealers described in section 1029(a) of the Dodd-Frank Act that are predominantly engaged in the sale and servicing of motor vehicles, the leasing and servicing of motor vehicles, or both.

In addition, the FTC retains its authority to enforce the risk-based pricing provisions of the FCRA and the FTC and CFPB rules issued under those provisions. Thus, the FTC and CFPB...
have overlapping enforcement authority for many entities subject to the CFPB rule and the FTC has sole enforcement authority for the motor vehicle dealers subject to the FTC rule.

As an analytical framework to estimate PRA burden in the "Burden Statement" below, the FTC estimates burden pertaining to respondents over which both agencies have shared enforcement authority, divides the resulting total by one-half to reflect the FTC’s shared jurisdiction, and adds to the resulting subtotal the incremental estimated burden regarding the motor vehicle dealers described above over which the FTC retains exclusive enforcement (and rulemaking) authority.

**Burden Statement**

Under the PRA, 44 U.S.C. 3501–3521, Federal agencies must get OMB approval for each collection of information they conduct or sponsor. “Collection of information” includes agency requests or requirements to submit reports, keep records, or provide information to a third party. 44 U.S.C. 3502(3); 5 CFR 1320.3(c). The FTC is seeking clearance for its assumed share of the estimated PRA burden regarding the disclosure requirements under the FTC and CFPB Rules.

Pursuant to Section 3506(c)(2)(A) of the PRA, the FTC invites comments on: (1) Whether the disclosure requirements are necessary, including whether the information will be practically useful; (2) the accuracy of our burden estimates, including whether the methodology and assumptions used are valid; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of providing the required information to consumers. All comments should be filed as prescribed in the section above, and must be received on or before May 2, 2017.

Under §§ 640.3–640.4 of the FTC’s RBP Rule and §§ 1022.72–1022.73 of the CFPB Rule, a creditor must provide a risk-based pricing notice to a consumer when the creditor uses a consumer report to grant or extend credit to the consumer on material terms that are materially less favorable than the most favorable terms available to a substantial proportion of consumers from or through that creditor.

Additionally, these provisions require disclosure of credit scores and information relating to credit scores in risk-based pricing notices if a credit score of the consumer is used in setting the material terms of credit.

The FTC’s currently cleared burden totals, post-adjustment for the effects of the Dodd-Frank Act, are 9,652,500 hours based on an estimated population of 160,875 entities apportioned to FTC enforcement and/or rulemaking authority.7

Using the currently cleared estimates (post-adjustment for the effects of the Dodd-Frank Act) for the number of applicable motor vehicle dealers and their assumed recurring disclosure burdens, in addition to the estimated number of and burden for other entities over which the FTC shares enforcement burden with the CFPB, the FTC proposes the following updated estimates:

**A. Estimated Number of Respondents:** 160,250.

**B. Burden Hours:** 9,615,000.

Yearly recurring burden of 60 hours per respondent 8 to modify and distribute notices × 160,250 respondents = 9,615,000 hours, cumulatively.

**C. Labor Costs:** $167,974,050.

Labor costs are derived by applying appropriate estimated hourly cost figures to the burden hours described above. The FTC assumes that respondents will use correspondence

7 OMB Control No. 3084-0145.
8 This estimate derives in part from an analysis of the figures obtained from the North American Industry Classification System (NAICS) Association’s database of U.S. businesses. See [link]. Commission staff identified categories of entities under its jurisdiction that also directly provide credit to consumers. Those categories include retail, vehicle dealers, consumer lenders, and utilities. The estimate also included credit union entities, which are subject to the Commission’s jurisdiction. See 15 U.S.C. 1681s. For the latter category, Commission staff relied on estimates from the Credit Union National Association for the number of non-federal credit unions. See [link].

9,615,000.

The FTC estimates that $17.47,10 to modify and distribute notices to consumers, for a cumulative labor cost total of $167,974,050.

**D. Capital/Non-Labor Costs:** $0.

The FTC believes that the FTC and CFPB 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) for the information collections discussed above.

**Request for Comment**

You can file a comment online or on paper. For the Commission to consider your comment, we must receive it on or before May 2, 2017. Write “RBP Rule, PRA Comment, P145403,” 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 Web site, at [link]. As a matter of discretion, the Commission tries to remove individuals’ home contact information from comments before placing them on the Commission Web site.

Because your comment will be made public, you are solely responsible for making sure that your comment does not include any sensitive personal information, like anyone’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 debit card number. You are also solely responsible for making sure that your comment does not include any sensitive health information, like medical records or other individually identifiable health information. In addition, do not include any “[t]rade secret or any commercial or financial information which is . . . privileged or confidential” as provided in Section 6(f) of the FTC Act 15 U.S.C. 46(f), and FTC Rule 4.10(a)(2), 16CFR 4.10(a)(2). In particular, do not include competitively sensitive information such as costs, sales statistics, inventories, formulas, patterns devices, manufacturing processes, or customer names.

If you want the Commission to give your comment confidential treatment, you must file it in paper form, with a request for confidential treatment, and you have to follow the procedure

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6 16 CFR 640.3–640.4.
7 12 CFR 1022.72–1022.73.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Request for Nominations of Candidates To Serve on the Clinical Laboratory Improvement Advisory Committee (CLIAC)

The Centers for Disease Control and Prevention (CDC) is soliciting nominations for membership on CLIAC. CLIAC provides scientific and technical advice and guidance to the Secretary, Department of Health and Human Services (HHS); the Assistant Secretary for Health, HHS; the Director, Centers for Disease Control and Prevention (CDC); the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare & Medicaid Services (CMS). The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory medicine. In addition, the Committee provides advice and guidance on specific questions related to possible revision of the CLIA standards. Examples include providing guidance on studies designed to improve safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods, the electronic transmission of laboratory information, and mechanisms to improve the integration of public health and clinical laboratory practices.

CLIAC consists of 20 members including the Chair, represents a diverse membership across laboratory specialties, professional roles (laboratory management, technical specialists, physicians, nurses) and practice settings (academic, clinical, public health), and includes a consumer representative. In addition, the Committee includes three ex officio members (or designees), including the Director, CDC; the Administrator, CMS; and the Commissioner, FDA. A nonvoting representative from the Advanced Medical Technology Association (AdvaMed) serves as the industry liaison. The Designated Federal Official (DFO) or their designee and the Executive Secretary are present at all meetings to ensure meetings are within applicable statutory, regulatory and HHS General Administration manual directives.

Request for Candidates: Nominations are being sought for individuals who have expertise and qualifications necessary to contribute to accomplishing CLIAC’s objectives. Nominees will be selected by the HHS Secretary or designee from authorities knowledgeable across the fields of microbiology (including bacteriology, mycobacteriology, mycology, parasitology, and virology), immunology (including histocompatibility), chemistry, hematology, pathology (including histopathology and cytology), or genetic testing (including cytogenetics); representatives from the fields of medical technology, public health, and clinical practice; and consumer representatives. Members may be invited to serve for terms of up to four years.

The U.S. Department of Health and Human Services policy stipulates that Committee membership be balanced in terms of points of view represented, and the committee’s function. Appointments shall be made without discrimination on the basis of age, race, ethnicity, gender, sexual orientation, gender identity, HIV status, disability, and cultural, religious, or socioeconomic status. Nominees must be U.S. citizens, and cannot be full-time employees of the U.S. Government. Current participation on federal workgroups or prior experience serving on a federal advisory committee does not disqualify a candidate; however, HHS policy is to avoid excessive individual service on advisory committees and multiple committee memberships. Committee members are Special Government Employees, requiring the filing of financial disclosure reports at the beginning and annually during their terms. CDC reviews potential candidates for CLIAC membership each year, and provides a slate of nominees for consideration to the Secretary of HHS for final selection. HHS notifies selected candidates of their appointment near the start of the term in July, or as soon as the HHS selection process is completed. Note that the need for different expertise and individuals to maintain the appropriate demographic balance varies from year to year and a candidate who is not selected in one year may be reconsidered in a subsequent year.

Candidates should submit the following items to be considered for nomination. The deadline for receipt of materials for the 2018 term is May 1, 2017:

- Current curriculum vitae, including complete contact information (name,