Current actions: On January 22, 2018 the Board published a notice in the **Federal Register** (83 FR 2983) requesting public comment for 60 days on the extension, with revision, of the Reporting Requirements Associated with Resolution Plans (Regulation QQ). The revision to the clearance is burden increase due to a reassessment of the burden hours associated with responding to the informational requirements of Regulation QQ and to guidance, feedback, and additional requests for information by the agencies as part of the iterative resolution planning process. The increase in burden is mitigated by the postponement of the July 2018 submission date for the resolution plans of the complex domestic filers, which account for the largest percentage of overall burden hours. The comment period for this notice expired on March 23, 2018. The Board received one comment on the proposal. The commenter recommended a number of potential changes to Regulation QQ intended to enhance the quality of the information collected pursuant to the regulation and reduce the burden of the information collection requirements.[7]

The Board is not adopting any of the recommended changes at this time. Either a revision to the Board's Regulation QQ or joint action with the FDIC would be necessary to implement each of the recommended changes. Most of the recommendations would require changes to the Board's Regulation QQ, which could only be accomplished pursuant to a rulemaking. In addition, the Board could not unilaterally take the actions requested by these comments, even those that would not require a rulemaking, as they fall under the purview of a rule that the Board proposed jointly with the FDIC and a process that is jointly administered by the two agencies.[8] However, the Board will consider the recommended changes in due course as it determines, in consultation with the FDIC, whether to conduct a joint rulemaking. The revisions will be implemented as proposed.

Board of Governors of the Federal Reserve System, August 15, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–17964 Filed 8–20–18; 8:45 am]

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[7] These recommended changes include:

(i) Extending the annual resolution plan filing cycle to a two-year cycle;

(ii) providing additional clarity on filing deadlines;

(iii) requiring that any agency guidance be provided more than 12 months in advance of each filing deadline;

(iv) allowing firms to satisfy some of their Regulation QQ requirements by incorporating their IDI plans by reference;

(v) providing for further tailoring based on the systemic risk posed by each firm,

(vi) further reducing the need for duplicative reporting;

(vii) adjusting the forecasting expected from the firms;

(viii) providing greater guidance regarding regulatory expectations related to the resolution of financial market utilities;

(ix) eliminating the strategic analysis section from tailored plans;

(x) providing an opportunity for notice and comment on any new information requirements, the framework used for assessing resolution plans, and the procedures related to remediation;

(xi) requiring the agencies to provide feedback on plans within six months of plan submission;

(xii) refraining from making feedback provided to the firms public or providing firms more time to consider the feedback before it is made public; and

(xiii) reconsidering the procedures the Board and FDIC undertake to engage with firms.

[8] See 12 U.S.C. 5365(d)(8) (requiring the Board and FDIC to issue joint rules implementing the Dodd-Frank Act's resolution planning requirements), 12 CFR. Part 243 (the Board's resolution planning rule), and 12 CFR. Part 381 (the FDIC's resolution planning rule). Aspects of the statute and regulations require joint actions or determinations by the Board and FDIC and therefore the agencies have jointly developed a coordinated resolution plan review process.

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 10 of the Home Owners' Loan Act (12 U.S.C. 1467a) (HOLA) and Regulation LL, (12 CFR part 238) 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 § 238.53 of Regulation LL (12 CFR 225.53). 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 10(c)(4)(B) of the HOLA 12 U.S.C. 1467a(c)(4)(B).

Unless otherwise noted, comments regarding the notices must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than September 4, 2018.

A. Federal Reserve Bank of Chicago (Colette A. Fried, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690–1414:

1. *McHenry Bancorp, Inc., McHenry, Illinois;* to engage de novo in purchasing and servicing loans, and holding and managing improved real estate, pursuant to sections 238.53(b)(1) and (8) of Regulation LL.

Board of Governors of the Federal Reserve System, August 16, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–17975 Filed 8–20–18; 8:45 am]

BILLING CODE 6210–01–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 September 18, 2018.

A. Federal Reserve Bank of Dallas (Robert L. Triplett III, Senior Vice President) 2200 North Pearl Street, Dallas, Texas 75201–2272:

1. *Woodforest Financial Group Employee Stock Ownership Plan, The Woodlands, Texas; and Woodforest Financial Group Employee Stock Ownership Trust, Spring, Texas*; to acquire up to an additional 28 percent

of Woodforest Financial Group, Inc., The Woodlands, Texas, and thereby indirectly acquire Woodforest National Bank, Houston, Texas.

Board of Governors of the Federal Reserve System, August 16, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–17974 Filed 8–20–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60-Day–18–0210; Docket No. CDC–2018–0069]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed and/or continuing information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a continuing information collection project titled List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products.

DATES: CDC must receive written comments on or before October 22, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2018–0069 by any of the following methods:

- *Federal eRulemaking Portal: Regulations.gov*. Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS–D74, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *Regulations.gov*.

Please note: Submit all comments through the Federal eRulemaking portal (egulations.gov) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffery M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road, NE, MS–D74, Atlanta, Georgia 30329; phone: 404–639–7570; Email: *omb@cdc.gov*.

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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected; and

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses.

5. Assess information collection costs.

Proposed Project

List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products—Extension (OMB# 0920–0210 Exp.Date 12/31/2018)—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Cigarette smoking is the leading preventable cause of premature death and disability in our Nation. Each year more than 480,000 deaths occur as the result of cigarette smoking-related diseases.

The CDC's Office on Smoking and Health (OSH) has the primary responsibility for the HHS smoking and health program. Since 1986, as required by the Comprehensive Smoking Education Act (CSEA) of 1984, which amended the Federal Cigarette Labeling and Advertising Act, 15 U.S.C. 1335a, CDC has collected information about the ingredients used in cigarette products. HHS has delegated responsibility for implementing the required information collection to CDC's OSH. Respondents are commercial cigarette manufacturers, packagers, or importers (or their representatives), who are required by the CSEA to submit ingredient reports to HHS on an annual basis.

Respondents are not required to submit specific forms; however, they are required to submit a list of all ingredients used in their products. CDC requires the ingredient report to be submitted by chemical name and Chemical Abstract Service (CAS) Registration Number, consistent with accepted reporting practices for other companies currently required to report ingredients added to other consumer products. The information collected is subject to strict confidentiality provisions.

Ingredient reports are due annually on March 31. Information is submitted to CDC by mailing or faxing a written report on the respondent's letterhead. All faxed lists should be followed up with a mailed original. Data may also be submitted to CDC by CD, three-inch floppy disk, or thumb drive. Electronic mail submissions are not accepted. Mail Annual Ingredient Submissions to Attention: FCLAA Program Manager, Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion, Centers for Disease Control and Prevention, 4770 Buford Highway, NE, MS S107–7, Atlanta, GA 30341–3717

Upon receipt and verification of the annual ingredient report, CDC issues a Certificate of Compliance to the respondent. As deemed appropriate by the Secretary of HHS, HHS is authorized to use the information to report to Congress the health effects of ingredients, research activities related to the health effects of ingredients, and other information that the Secretary determines to be of public interest. There are no costs to respondents other