fair value option to financial or servicing assets and liabilities.

The agencies believe that the proposed information is necessary to more accurately assess the impact of fair value accounting and fair value measurements for safety and soundness purposes. The collection of the information on Schedule Q, as proposed, will facilitate and enhance the banking agencies' ability to monitor the extent of fair value accounting by branches, including the elective use of fair value accounting and the nature of the inputs used in the valuation process, pursuant to the disclosure requirements of ASC Topic 820. The information collected on Schedule Q is consistent with the disclosures required by ASC Topic 820 and consistent with industry practice for reporting fair value measurements and should, therefore, not impose significant incremental burden on branches.

## **Request for Comment**

Comments are invited on:

a. Whether the information collections are necessary for the proper performance of the agencies' functions, including whether the information has practical utility;

b. The accuracy of the agencies' 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;

d. Ways to minimize the burden of information collections on respondents, including through the use of automated collection techniques or other forms of information technology; and

e. Estimates of capital or start up costs and costs of operation, maintenance, and purchase of services to provide information.

Comments submitted in response to this notice will be shared among the agencies. All comments will become a matter of public record. Written comments should address the accuracy of the burden estimates and ways to minimize burden including the use of automated collection techniques or the use of other forms of information technology as well as other relevant aspects of the information collection request.

Board of Governors of the Federal Reserve System, January 5, 2011.

# Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 2011–270 Filed 1–10–11; 8:45 am] BILLING CODE 6210–01–P

# FEDERAL RESERVE SYSTEM

## Federal Open Market Committee; Domestic Policy Directive of December 14, 2010

In accordance with Section 271.25 of its rules regarding availability of information (12 CFR part 271), there is set forth below the domestic policy directive issued by the Federal Open Market Committee at its meeting held on December 14, 2010.<sup>1</sup>

The Federal Open Market Committee seeks monetary and financial conditions that will foster price stability and promote sustainable growth in output. To further its long-run objectives, the Committee seeks conditions in reserve markets consistent with federal funds trading in a range from 0 to  $\frac{1}{4}$  percent. The Committee directs the Desk to execute purchases of longer-term Treasury securities in order to increase the total face value of domestic securities held in the System Open Market Account to approximately \$2.6 trillion by the end of June 2011. The Committee also directs the Desk to reinvest principal payments from agency debt and agency mortgagebacked securities in longer-term Treasury securities. The System Open Market Account Manager and the Secretary will keep the Committee informed of ongoing developments regarding the System's balance sheet that could affect the attainment over time of the Committee's objectives of maximum employment and price stability.

By order of the Federal Open Market Committee, January 5, 2011.

# William B. English,

Secretary, Federal Open Market Committee. [FR Doc. 2011–348 Filed 1–10–11; 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 application also will 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 February 7, 2011.

A. Federal Reserve Bank of Dallas (E. Ann Worthy, Vice President)2200 North Pearl Street, Dallas, Texas 75201–2272:

1. Sulphur Springs Bancshares, Inc., Sulphur Springs, Texas, to acquire by merger 100 percent of First Mineola, Inc., and indirectly acquire The First National Bank of Mineola, both of Mineola, Texas.

Board of Governors of the Federal Reserve System, January 6, 2011.

#### Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. 2011–341 Filed 1–10–11; 8:45 am] BILLING CODE 6210–01–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS-0990-New; 60-Day Notice]

## Agency Information Collection Request; 60-Day Public Comment Request

**AGENCY:** Office of the Secretary, HHS. In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed information collection request for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information,

<sup>&</sup>lt;sup>1</sup>Copies of the Minutes of the Federal Open Market Committee at its meeting held on December 14, 2010, which includes the domestic policy directive issued at the meeting, are available upon request to the Board of Governors of the Federal Reserve System, Washington, DC 20551. The minutes are published in the Federal Reserve Bulletin and in the Board's Annual Report.

including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to

Sherette.funncoleman@hhs.gov, or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be directed to the OS Paperwork Clearance Officer at the above e-mail address within 60 days.

*Proposed Project:* Comparative Effectiveness Research Inventory—OMB No. 0990–New-Assistant Secretary for Planning and Evaluation (ASPE).

Abstract: The Office of the Assistance Secretary for Planning and Evaluation (ASPE) is requesting approval by OMB for the collection of information submitted by content users directly to a web-based inventory of comparative effectiveness research (CER). The CER Inventory will categorize and catalogue Federal and non-Federal CER outputs and activities across four main domains: Research, human & scientific capital (e.g., training/education, methods development), data infrastructure, and dissemination & translation. The CER inventory will serve as a valuable tool for researchers, providers, patients, policymakers, and other users.

The CER inventory will draw upon primary data sources, including PubMed, HSRProj, ClinicalTrials.gov, and NIH RePORTER. Working with these four major sources and using the Federal Coordinating Council for CER's definition of CER and strategic framework, selection criteria and tools to select and extract the appropriate subsets of these datasets for inclusion in the CER inventory will be identified. In addition, content owners wishing to submit CER records to the CER

inventory will be directed first to submit such records to one of these main primary source databases, as appropriate. This method will not only help to augment these existing databases, it will enable efficient and effective capture of CER information for the CER Inventory via CER search filters, etc., that have been developed for those respective source databases. If candidate CER records under consideration are not suitable for submission to one of these main databases, an alternative method that allows for direct submissions to the CER inventory will be made available to content users. Examples include reports and published articles or projects and programs that focus on areas of CER outside of primary research (e.g., training and education). The pilot inventory tool will provide a web form that may be used by content owners to submit CER records, subject to validation. This process for direct submission will draw from the experience with content owner submissions for such established databases as HSRProj and ClinicalTrials.gov.

#### ESTIMATED ANNUALIZED BURDEN TABLE

Form	Type of respondent	Number of respondents	Number of responses per respondent	Total responses	Average burden hours per response	Total burden hours
CER Inventory Direct Submission Form for Reports or Other Publications.	Researchers/ Research Assistants.	400	1	400	25/60	167
CER Inventory Direct Submission Form for Projects.	Researchers/ Research Assistants.	100	1	100	28/60	47
Total						214

#### Seleda M. Perryman,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer. [FR Doc. 2011–310 Filed 1–10–11; 8:45 am] BILLING CODE 4150–05–P

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# DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Centers for Disease Control and Prevention

[30 Day-11-0210]

## Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–5960 or send an email to *omb@cdc.gov*. Send written comments to CDC Desk Officer, Office of Management and Budget, Washington, DC or by fax to (202) 395–5806. Written comments should be received within 30 days of this notice.

# **Proposed Project**

List of Ingredients Added to Tobacco in the Manufacture of Cigarette Products (OMB# 0920–0210 Exp. 04/30/2011)— Extension—Office on Smoking and Health, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention, (CDC).

#### **Background and Brief Description**

The Centers for Disease Control and Prevention (CDC), Office on Smoking and Health (OSH) has the primary responsibility for the Department of Health and Human Services (HHS) smoking and health program. HHS's overall goal is to reduce death and disability resulting from cigarette smoking and other forms of tobacco use through programs of information, education and research.

Since 1986, as required by the Comprehensive Smoking Education Act of 1984 (CSEA, 15 U.S.C. 1336 or Pub. L. 98–474), CDC has collected information about the ingredients used in cigarette products. Respondents are commercial cigarette manufacturers, packagers, or importers (or their representatives), who are required by