

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 27, 2018.

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. *Summit Bancshares, Inc., Chesterfield, Missouri*; to become a bank holding company by acquiring 100 percent of the voting shares of The Bank of Houston, Houston, Missouri.

Board of Governors of the Federal Reserve System, August 28, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–18997 Filed 8–30–18; 8:45 am]

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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 September 19, 2018.

A. Federal Reserve Bank of Minneapolis (Mark A. Rauzi, Vice President), 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291:

1. *Brian Solsrud, individually and as trustee of the Descendant's Separate Trust fbo Brian K. Solsrud under the Glenn A. Solsrud Augusta Irrevocable Trust dated December 28, 2012 and the Descendant's Separate Trust fbo Brian K. Solsrud under the Ardath K. Solsrud Augusta Irrevocable Trust dated December 28, 2012, all of North Oaks, Minnesota; and Rachel Goodell, Augusta, Wisconsin; Corinne Solsrud, Mosinee, Wisconsin; and Gregory Solsrud, Dunwoody, Georgia, each individually*; to acquire voting shares of Augusta Financial Corporation and thereby indirectly acquire shares of Unity Bank, both of Augusta, Wisconsin.

2. *Brian Solsrud, individually and as trustee of the Descendant's Separate Trust fbo Brian K. Solsrud under the Glenn A. Solsrud Caprice Irrevocable Trust dated December 28, 2012 and the Descendant's Separate Trust fbo Brian K. Solsrud under the Ardath K. Solsrud Caprice Irrevocable Trust dated December 28, 2012, all of North Oaks, Minnesota; and Rachel Goodell, Augusta, Wisconsin; Corinne Solsrud, Mosinee, Wisconsin; and Gregory Solsrud, Dunwoody, Georgia, each individually*; to acquire voting shares of Caprice Corporation, Augusta, Wisconsin, and thereby indirectly acquire shares of Unity Bank North, Red Lake Falls, Minnesota.

B. Federal Reserve Bank of San Francisco (Gerald C. Tsai, Director, Applications and Enforcement) 101 Market Street, San Francisco, California 94105–1579:

1. *Kevin Stacy Garn, Layton, Utah, Courtney Allphin, Layton, Utah, Gabe Garn, Syracuse, Utah, Jake Garn, Layton, Utah, Jordan Garn, Farmington, Utah, Talmage Garn, Salt Lake City, Utah, and Taylee Goff, Farmington, Utah*; to retain voting shares of FNB Bancorp, and thereby indirectly retain voting shares of First National Bank of Layton, both of Layton, Utah.

Board of Governors of the Federal Reserve System, August 28, 2018.

Ann Misback,

Secretary of the Board.

[FR Doc. 2018–18996 Filed 8–30–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30Day–18–0743]

Agency Forms Undergoing Paperwork Reduction Act Review

In accordance with the Paperwork Reduction Act of 1995, the Centers for Disease Control and Prevention (CDC) has submitted the information collection request titled Monitoring Breastfeeding-Related Maternity Care—U.S. hospitals to the Office of Management and Budget (OMB) for review and approval. CDC previously published a “Proposed Data Collection Submitted for Public Comment and Recommendations” notice on November 22, 2017, to obtain comments from the public and affected agencies. CDC received 12 comments related to the previous notice. This notice serves to allow an additional 30 days for public and affected agency comments.

CDC will accept all comments for this proposed information collection project. The Office of Management and Budget is particularly interested in comments that:

(a) 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;

(b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(c) Enhance the quality, utility, and clarity of the information to be collected;

(d) 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 submission of responses; and

(e) Assess information collection costs.

To request additional information on the proposed project or to obtain a copy of the information collection plan and instruments, call (404) 639–7570 or send an email to omb@cdc.gov. Direct written comments and/or suggestions regarding the items contained in this notice to the Attention: CDC Desk Officer, Office of Management and Budget, 725 17th Street NW, Washington, DC 20503 or by fax to (202)

395–5806. Provide written comments within 30 days of notice publication.

Proposed Project

Monitoring Breastfeeding-Related Maternity Care—U.S. Hospitals (OMB Control No. 0920–0743, Exp. 9/30/2016)—Reinstatement with Change—Division of Nutrition, Physical Activity, and Obesity, National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Substantial evidence demonstrates the social, economic, and health benefits of breastfeeding for both the mother and infant as well as for society in general. Breastfeeding mothers have lower risks of breast and ovarian cancers and type 2 diabetes, and breastfeeding better protects infants against infections, chronic diseases like diabetes and obesity, and even childhood leukemia and sudden infant death syndrome (SIDS). However, the groups that are at higher risk for diabetes, obesity, and poor health overall persistently have the lowest breastfeeding rates.

Health professionals recommend at least 12 months of breastfeeding, and Healthy People 2020 establishes specific national breastfeeding goals. In addition to increasing overall rates, a significant public health priority in the United States is to reduce variation in breastfeeding rates across population subgroups. Although CDC surveillance data indicate that breastfeeding initiation rates in the United States are climbing, rates for duration and exclusivity continue to lag, and significant disparities persist between African American and white women in breastfeeding rates.

The health care system is one of the most important and effective settings to improve breastfeeding. Recognition of the hospital stay as a crucial influence in later breastfeeding outcomes led to the addition of two objectives in Healthy People 2020 to allow national monitoring of improvements in support for breastfeeding during this time. In 2007, CDC conducted the first national survey of Maternity Practices in Infant

Nutrition and Care (known as the mPINC Survey) in health care facilities (hospitals and free-standing childbirth centers). This survey was designed to provide baseline information and to be repeated every two years. The survey was conducted again in 2009, 2011, 2013, and 2015. The survey inquired about patient education and support for breastfeeding throughout the maternity stay as well as staff training and maternity care policies.

Prior to the fielding of the 2009 iteration, CDC was requested to provide a report to OMB on the results of the 2007 collection. In this report, CDC provided survey results by geographic and demographic characteristics and a summary of activities that resulted from the survey. A summary of mPINC findings was also the anchor of all activities related to the CDC August 2011 Vital Signs activity, marking the first time that CDC highlighted improving hospital maternity practices as the CDC-wide public health priority. A summary of mPINC findings provided the basis of the CDC October 2015 Vital Signs report, which updated the 2011 Vital Signs report and concluded that although maternity care policies and practices supportive of breastfeeding are improving nationally; more work is needed to ensure all women receive optimal breastfeeding support during the birth hospitalization.

The planned methodology for the 2018 and 2020 national survey of Maternity Practices in Infant Nutrition and Care (mPINC) will closely match that of the previously administered mPINC surveys in 2007, 2009, 2011, 2013, and 2015. Changes described in this Reinstatement with change include: (1) Deployment of 2018 and 2020 Surveys; (2) data collection via web-survey only (no paper surveys); (3) surveying hospitals only (not birth centers); (4) requesting contact information for two individuals per facility (previously only one); (5) an updated American Hospital Association (AHA) database will be acquired to identify hospitals not currently on the list for recruitment in the 2018 survey. This process will not occur for the 2020

survey, but additional hospitals identified from the new database for 2018 will be included in the 2020 survey; (6) 2018 and 2020 survey content has been updated.

A major strength of the mPINC survey is its structure as an ongoing national census, which does not employ sampling methods. Facilities are identified by using the American Hospital Association (AHA) Annual Survey of Hospitals. Facilities that will be invited to participate in the survey include hospitals that participated in previous iterations and those that were invited but did not participate in the previous iterations, as well as those that have become eligible since the most recent mPINC survey. All hospitals with ≥1 registered maternity bed will be screened via a brief phone call to assess their eligibility, identify additional satellite locations, and identify the appropriate point of contact. The high response rates to the previous iterations of the mPINC survey (82–83% in 2007, 2009, 2011, 2013, and 2015) indicate that the methodology is appropriate and also reflects high interest among the study population.

As with the initial surveys, a major goal of the 2018 and 2020 follow-up surveys is to be fully responsive to hospitals’ needs for information and technical assistance. CDC will provide direct feedback to hospital respondents in a customized benchmark report of their results. CDC will use information from the mPINC surveys to identify, document, and share information related to incremental changes in practices and care processes over time at the hospital, state, and national levels. Data are also used by researchers to better understand the relationships between hospital characteristics, maternity-care practices, state level factors, and breastfeeding initiation and continuation rates. Participation in the survey is voluntary, and responses may be submitted through a Web-based system. The total estimated annual Burden Hours are 855. There are no costs to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Maternity Hospital	Screening Call Script Part A	1,952	1	1/60
Maternity Hospital	Screening Call Script Part B	1,672	1	4/60
Maternity Hospital	mPINC Facility Survey	1,421	1	30/60

Jeffrey M. Zirger,

Acting Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2018–19012 Filed 8–30–18; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60 Day–18–16JO; Docket No. CDC–2018–0077]

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 proposed information collection project titled Pregnancy Risk Assessment Monitoring System (PRAMS). PRAMS provides an important supplement to vital records data by providing state-specific information not available through birth certificate data on maternal behaviors and experiences before, during and after pregnancy on health conditions, prenatal care, postpartum care, access to care, and health insurance status.

DATES: CDC must receive written comments on or before October 30, 2018.

ADDRESSES: You may submit comments, identified by Docket No. CDC–201x–xxxx 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 (regulations.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 Jeffrey 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

The Pregnancy Risk Assessment Monitoring System (PRAMS)—Existing Collection in Use without an OMB Control Number—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) seeks OMB approval to collect information through the Pregnancy Risk Assessment Monitoring System (PRAMS) for three years as a generic clearance. OMB approval for new modules will be submitted through the part of generic clearance mechanism.

PRAMS supplements vital records data by providing state-specific information on maternal behaviors and experiences before, during and after pregnancy. Every month, in each participating state, a sample of women who have recently given birth to a live born or stillborn infant is selected from birth certificates or fetal death files. The sample is stratified based on the state's population of interest to ensure high-risk populations are represented in the data. PRAMS is a state customized mail and telephone survey conducted in 51 sites and covers 83% of all live births in the United States. Information is collected by self-administered mail survey with telephone follow-up for non-responders. Because PRAMS uses standardized data collection methods, it allows data to be compared among states.

The PRAMS survey instrument is based on a core set of questions common across all states. Core questions request information that is not available from vital records; information about health conditions, prenatal care, postpartum care, access to care, or health insurance status; information about contraception, health habits or risk behaviors; and information about other topics such as breastfeeding. In addition, CDC provides participating states with standard questions from optional modules that states may use to customize survey content for their specific needs at the beginning of each Phase of data collection. In addition, on occasion, states may be funded to address emerging topics of interest to collect supplemental data on optional modules of interest. These questions can be used to address state-specific priorities and special topics such as, for example, substance use, including prescription and illicit opioid use, disease epidemics, or other topics related to healthy pregnancy; these supplements can be administered to women identified in the usual manner or via hospital records. States not intending to implement the survey on an ongoing basis, can instead employ a point-in-time survey. Because PRAMS infrastructure was developed to access a specific and vulnerable subpopulation, the PRAMS infrastructure can be