

of the Board of Governors. Comments must be received not later than June 26, 2012.

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

1. *The Gause Family, consisting of Bryce and Sheila Gause, Lynnville, Iowa; HW and Nancy Barnhouse, Vero Beach, Florida; Lester and Kay Gause, Newton, Iowa; Charles Gause, Providence, North Carolina; Gary and Joan Ales, Lakewood Ranch, Florida; Rebecca Barnhouse, Youngstown, Ohio; Richard Buls, New Market, Maryland; Kristy Crawford, Frederick, Maryland; Curtis Gause, Pleasant Hill, Iowa; Peggy Gause, Roanoke, Virginia; Russell Gause, Pasadena, Texas; Carrie Holub, Davenport, Iowa; Connie Kopacek, Urbandale, Iowa; Cynthia Smith, Newton, Iowa; and Kimberly Soulen, Meyersville, Maryland*, all acting in concert, to retain control of First State Bank Holding Company, and thereby indirectly retain control of First State Bank, both in Lynnville, Iowa.

Board of Governors of the Federal Reserve System, June 6, 2012.

Michael J. Lewandowski,

Assistant Secretary of the Board.

[FR Doc. 2012-14086 Filed 6-8-12; 8:45 am]

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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 July 5, 2012.

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

1. *DFW Capital Holdings, Inc.*, and DFW Capital Holdings Merger Corporation, both in Dallas, Texas; to become bank holding companies by acquiring Schwertner State Bank, Schwertner, Texas.

Board of Governors of the Federal Reserve System, June 5, 2012.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 2012-13980 Filed 6-8-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Implementation of Federal Financial Report—Upcoming Mandatory Use of the Federal Financial Report System in the eRA Commons

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

ACTION: Notice.

Purpose

Beginning October 1, 2012, CDC will implement the expenditure data portion of the Federal Financial Report (FFR) in the Electronic Research Administration (eRA) Commons. The transition to use the FFR for reporting expenditure data includes new reporting dates for annual FFRs, and reporting of cumulative data only.

Background

The Office of Management and Budget has consolidated the Financial Status Report (FSR or SF-269/SF-269A) and the Federal Cash Transaction Report (FCTR or SF-272/SF-272A) into a single form known as the Federal Financial Report (FFR or SF-425/SF-425A). Since January 1, 2010, CDC grantees have been required to report cash transaction data via the Payment Management System (PMS) using the FFR cash transaction data elements. The FSR/FFR module allows grantees to

electronically submit a statement of expenditures associated with their grant to the sponsor of the grant via eRA Commons. The new eRA Commons system was piloted with 5 Federal Demonstration Partnership (FDP) institutions that assisted CDC with feedback and testing during the Third Quarter of Fiscal Year 2011.

The Electronic Research Administration (eRA) was established by the National Institutes of Health (NIH) in response to the Government Paperwork Elimination Act requiring federal agencies to pursue electronic means of production. The intent of eRA is to provide for secure receipt, review and administration of electronic grants. The eRA Commons offers a meeting place for grantees tracking research grants administration information and applications and will now be used by grantees to submit their FFRs.

DATES: The effective date is October 1, 2012.

FOR FURTHER INFORMATION CONTACT:

Technical Information Management Section (TIMS), Procurement and Grants Office, Centers for Disease Control and Prevention, 2920 Brandywine Rd, NE., Atlanta, GA 30341; telephone (770) 488-2700; email @ PGOTIM@CDC.GOV.

Implementation

All CDC Financial Expenditure data due on/after October 1, 2012 must be submitted using the FFR via the eFSR/FFR system in the eRA Commons. All Federal Reporting in the Payment Management System is unchanged. All new submissions should be prepared and submitted as FFRs.

CDC's implementation of the FFR retains a financial reporting period that coincides with the budget period of a particular project. However, the due date for annual FFRs will be 90 days after the end of the calendar quarter in which the budget period ends. Note that this is a change in due dates of annual FFRs and may provide up to 60 additional days to report, depending upon when the budget period end date falls within a calendar quarter. For example, if the budget period ends 1/30/2012, the annual FFR is due 6/30/2012 (90 days after the end of the calendar quarter of 3/31/2012).

Due dates of final reports will remain unchanged. The due date for final FFRs will continue to be 90 days after the project period end date.

Grantees must submit closeout reports in a timely manner. Unless the Grants Management Officer (GMO) of the awarding Institute or Center approves an extension, grantees must submit a final FFR, final progress report, and

Final Invention Statement and Certification within 90 days of the end of grant period. Failure to submit timely and accurate final reports may affect future funding to the organization or awards under the direction of the same Project Director/Principal Investigator.

FFR (SF 425) instructions for CDC Grantees are now available at <http://grants.nih.gov/grants/forms.htm>. For further information, contact GrantsInfo@nih.gov. Additional resources concerning the eFSR/FFR system, including a User Guide and an on-line demonstration, can be found on the eRA Commons Support Page: <http://www.cdc.gov/od/pgo/funding/grants/eramain.shtm>.

Getting Started—eRA Commons Registration

The submission of FFRs to CDC will require organizations to register with eRA Commons (Commons) (<https://commons.era.nih.gov/commons/>). CDC recommends that this one time registration process be completed at least 2 weeks prior to the submittal date of a FFR submission.

Organizations may verify their current registration status by running the “List of Commons Registered Organizations” query found at: <http://era.nih.gov/commons/>. Organizations not yet registered can go to <https://commons.era.nih.gov/commons/registration/registrationInstructions.jsp> for instructions. It generally takes several days to complete this registration process. This registration is independent of Grants.gov and may be done at any time.

The individual designated as the Principle Investigator (PI) on the application must also be registered in the Commons. The PI must hold a PI account *and* be affiliated with the applicant organization. This registration must be done by an organizational official or their delegate who is already registered in the Commons. To register PIs in the Commons, refer to the eRA Commons User Guide found at: <http://era.nih.gov/commons/index.cfm>.

Inquiries

General questions concerning using the eRA Commons should be directed to the eRA Commons Helpdesk at: eRA Commons Help Desk Web: <http://ithelpdesk.nih.gov/eRA/>. (Preferred method of contact).

Toll-free: 1-866-504-9552
 Phone: 301-402-7469
 TTY: 301-451-5939
 Hours: Mon-Fri, 7 a.m. to 8 p.m. Eastern Standard Time.
 Email: commons@od.nih.gov.

Dated: June 5, 2012.

Alan A. Kotch,
 Director, Procurement and Grants Office,
 Centers for Disease Control and Prevention.
 [FR Doc. 2012-14049 Filed 6-8-12; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Health and Nutrition Examination Survey (NHANES) DNA Samples

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

ACTION: Notice.

SUMMARY: The National Health and Nutrition Examination Survey (NHANES) will not be receiving DNA proposals in the near future. NHANES is changing its plan for making DNA available for genetic research and its proposal guidelines. NHANES will announce when it will reopen its repository for use of DNA specimens for research protocols once it has developed its new plan of operation.

DATES: Effective date is date of publication in the **Federal Register**.

ADDRESSES: Geraldine McQuillan, Ph.D., Division of Health and Nutrition Examination Surveys, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 4204, Hyattsville, MD

20782, Phone: 301-458-4371, Fax: 301-458-4028, EMail: NHANESgenetics@cdc.gov.

FOR FURTHER INFORMATION CONTACT: Dr. Geraldine McQuillan, Division of Health and Nutrition Examination Surveys, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 4204, Hyattsville, MD 20782, Phone: 301-458-4371, Fax: 301-458-4028, E-Mail: NHANESgenetics@cdc.gov.

Juliana Cyril,
 Deputy Director, Office of Science Quality,
 Office of the Associate Director for Science,
 Centers for Disease Control and Prevention.
 [FR Doc. 2012-14056 Filed 6-8-12; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:
Title: Extension to HS Transportation Requirement.

OMB No.: 0970-0260.

Description: The Office of Head Start is proposing to renew authority to collect information regarding the Head Start transportation requirement without changes. The transportation requirement provides the requirement that each child be seated in a child restraint system while the vehicle is in motion, and the requirement that each bus have at least one bus monitor on board at all times. Waivers would be granted when the Head Start or Early Head Start grantee demonstrates that compliance with the requirement(s) for which the waiver is being sought will result in a significant disruption to the Head Start program or the Early Head Start program and that waiving the requirement(s) is in the best interest of the children involved.

Respondents: Head Start and Early Head Start program grants recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per respondent	Total burden hours
Form	275	1	1	275

Estimated Total Annual Burden Hours: 275

In compliance with the requirements of Section 506(c)(2)(A) of the Paperwork Reduction Act of 1995, the

Administration for Children and Families is soliciting public comment on the specific aspects of the