

must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than November 30, 2015.

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

1. *Town and Country Financial Corporation*, Springfield, Illinois; to merge with West Plains Investors, Inc., and thereby indirectly acquire Premier Bank of Jacksonville, both in Jacksonville, Illinois.

B. Federal Reserve Bank of St. Louis (Yvonne Sparks, Community Development Officer) P.O. Box 442, St. Louis, Missouri 63166-2034:

1. *First Breckinridge Bancshares, Inc.*, Irvington, Kentucky; to acquire 100 percent of the voting shares of American Bank & Trust Company, Inc., Bowling Green, Kentucky.

Board of Governors of the Federal Reserve System, November 2, 2015.

Michael J. Lewandowski,

Associate Secretary of the Board.

[FR Doc. 2015-28262 Filed 11-4-15; 8:45 am]

BILLING CODE 6210-01-P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Savings and Loan Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Home Owners' Loan Act (12 U.S.C. 1461 *et seq.*) (HOLA), Regulation LL (12 CFR part 238), and Regulation MM (12 CFR part 239), and all other applicable statutes and regulations to become a savings and loan holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a savings association and nonbanking companies owned by the savings and loan 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 HOLA (12 U.S.C. 1467a(e)). 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 10(c)(4)(B) of the HOLA (12 U.S.C. 1467a(c)(4)(B)). 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 November 30, 2015.

A. Federal Reserve Bank of Atlanta (Chapelle Davis, Assistant Vice President) 1000 Peachtree Street NE., Atlanta, Georgia 30309:

1. *Oculina Banc Corp*, Vero Beach, Florida; proposes to merge with its parent company, Colonial Banc Corp, Vero Beach, Florida. Oculina Banc Corp will survive the merger. Colonial Banc Corp and Oculina Banc Corp control Oculina Bank, Vero Beach, Florida.

Board of Governors of the Federal Reserve System, November 2, 2015.

Michael J. Lewandowski,

Associate Secretary of the Board.

[FR Doc. 2015-28261 Filed 11-4-15; 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 November 30, 2015.

A. Federal Reserve Bank of Richmond (Adam M. Drimer, Assistant Vice President) 701 East Byrd Street, Richmond, Virginia 23261-4528:

1. *Southern BancShares (N.C.), Inc.*, Mount Olive, North Carolina; to acquire voting shares of Heritage Bankshares Inc., and thereby indirectly acquire Heritage Bank, both in Norfolk, Virginia.

B. Federal Reserve Bank of Minneapolis (Jacquelyn K. Brunmeier, Assistant Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480-0291:

1. *Darwin Bancshares, Inc.*, Darwin, Minnesota; to merge with Winthrop Bancshares, Inc., and thereby indirectly acquire Winthrop State Bank, both in Winthrop, Minnesota.

Board of Governors of the Federal Reserve System, October 30, 2015.

Margaret McCloskey Shanks,

Deputy Secretary of the Board.

[FR Doc. 2015-28128 Filed 11-4-15; 8:45 am]

BILLING CODE 6210-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-16-16CB; Docket No. CDC-2015-0094]

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 efforts to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. This notice invites comment on the evaluation of the progress of CDC partners that receive awards distributed via contracts, grants and cooperative agreements, from the Procurements and Grants Office (PGO). PGO is responsible for the stewardship of these funds while providing excellent, professional services to our partners and stakeholders. Data will be collected for the purpose of evaluating the progress of programmatic activities.

DATES: Written comments must be received on or before January 4, 2016.

ADDRESSES: You may submit comments, identified by Docket No. CDC-2015-0094 by any of the following methods:

- *Federal eRulemaking Portal:*

Regulations.gov. Follow the instructions for submitting comments.

- *Mail:* Leroy A. Richardson, 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. All relevant comments received will be posted without change to Regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to Regulations.gov.

Please note: All public comment should be submitted 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 the 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 OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information 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. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information.

Proposed Project

Performance Progress and Evaluation Report (PPER)—Existing Collection in use without an OMB Control Number—Procurements and Grants Office (PGO), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Each year, approximately 80% of the Centers for Disease Control and Prevention's (CDC) budget is distributed via contracts, grants and cooperative agreements, from the Procurements and Grants Office (PGO) to partners throughout the world to promote health, prevent disease, injury and disability and prepare for new health threats. PGO

is responsible for the stewardship of these funds while providing excellent, professional services to our partners and stakeholders.

Currently, CDC uses SF-PPR (a progress report form for Non-Research awards) to collect information semi-annually from Awardees regarding the progress made over specified time periods on CDC funded projects. The SF-PPR (OMB Control Number: 0970-0406, Expiration Date: 10/31/2015) is owned by the Administration for Children and Families (ACF) within the Department of Health and Human Services (HHS). This New ICR is being developed by CDC to create a CDC-wide collection tool called the Progress Performance and Evaluation Report (PPER) that will be used to collect data on the progress of CDC Awardees for the purposes of evaluation and to bring the Awardee reporting procedure into compliance with the Paperwork Reduction Act (PRA).

The information collected will enable the accurate, reliable, uniform and timely submission to CDC of each Awardee's work plans and progress reports, including strategies, activities and performance measures. The information collected by the PPER is designed to align with, and support the goals outlined for each of the CDC Awardees. Collection and reporting of the information will occur in an efficient, standardized, and user-friendly manner that will generate a variety of routine and customizable reports. The PPER will allow each Awardee to summarize activities and progress towards meeting performance measures and goals over a specified time period specific to each award. CDC will also have the capacity to generate reports that describe activities across multiple Awardees. In addition, CDC will use the information collection to respond to inquiries from HHS, Congress and other stakeholder inquiries about program activities and their impact.

The total estimated burden is 16,000 hours. There is no cost to respondents other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
CDC Non-Research contract, grant, and cooperative agreement Awardees.	Performance Progress and Evaluation Report (PPER).	8,000	1	120/60	16,000
Total	16,000

Leroy A. Richardson,

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. 2015–28155 Filed 11–4–15; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–16–0850; Docket No. CDC–2015–
0093]

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 efforts to reduce public
burden and maximize the utility of
government information, invites the
general public and other Federal
agencies to take this opportunity to
comment on proposed and/or
continuing information collections, as
required by the Paperwork Reduction
Act of 1995. This notice invites
comment on the proposed extension of
the Laboratory Response Network
information collection.

DATES: Written comments must be
received on or before January 4, 2016.

ADDRESSES: You may submit comments,
identified by Docket No. CDC–2015–
0093 by any of the following methods:

- *Federal eRulemaking Portal:*
Regulation.gov. Follow the instructions
for submitting comments.

- *Mail:* Leroy A. Richardson,
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. All relevant comments
received will be posted without change
to Regulations.gov, including any
personal information provided. For
access to the docket to read background
documents or comments received, go to
Regulations.gov.

Please note: All public comment should be
submitted 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 the 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 OMB for approval. To
comply with this requirement, we are
publishing this notice of a proposed
data collection as described below.

Comments are invited on: (a) Whether
the proposed collection of information
is necessary for the proper performance
of the functions of the agency, including
whether the information shall have
practical utility; (b) the accuracy of the
agency's estimate of the burden of the
proposed collection of information; (c)
ways to enhance the quality, utility, and
clarity of the information to be
collected; (d) ways to minimize the
burden of the collection of information
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. Burden means
the total time, effort, or financial
resources expended by persons to
generate, maintain, retain, disclose or
provide information to or for a Federal
agency. This includes the time needed
to review instructions; to develop,
acquire, install and utilize technology
and systems for the purpose of
collecting, validating and verifying
information, processing and
maintaining information, and disclosing
and providing information; to train
personnel and to be able to respond to
a collection of information, to search
data sources, to complete and review
the collection of information; and to
transmit or otherwise disclose the
information.

Proposed Project

Laboratory Response Network—
Extension—(OMB Control No. 0920–
0850, expires April 30, 2016), National
Center for Emerging and Zoonotic
Infectious Diseases (NCEZID), Centers
for Disease Control and Prevention
(CDC).

Background and Brief Description

The Laboratory Response Network
(LRN) was established by the
Department of Health and Human
Services (HHS), Centers for Disease
Control and Prevention (CDC) in
accordance with Presidential Decision
Directive 39, which outlined national
anti-terrorism policies and assigned
specific missions to Federal
departments and agencies. The LRN's
mission is to maintain an integrated
national and international network of
laboratories that can respond to
suspected acts of biological, chemical,
or radiological threats and other public
health emergencies.

When Federal, State and local public
health laboratories voluntarily join the
LRN, they assume specific
responsibilities and are required to
provide information to the LRN Program
Office at CDC. Each laboratory must
submit and maintain complete
information regarding the testing
capabilities of the laboratory.
Biennially, laboratories are required to
review, verify and update their testing
capability information. Complete testing
capability information is required in
order for the LRN Program Office to
determine the ability of the Network to
respond to a biological or chemical
threat event. The sensitivity of all
information associated with the LRN
requires the LRN Program Office to
obtain personal information about all
individuals accessing the LRN Web site.
In addition, the LRN Program Office
must be able to contact all laboratory
personnel during an event so each
laboratory staff member that obtains
access to the restricted LRN Web site
must provide his or her contact
information to the LRN Program Office.

As a requirement of membership, LRN
Laboratories must report all biological
and chemical testing results to the LRN
Program at CDC using a CDC developed
software tool called the LRN Results
Messenger. This information is essential
for surveillance of anomalies, to support
response to an event that may involve
multiple agencies and to manage limited
resources. LRN Laboratories must also
participate in and report results for
Proficiency Testing Challenges or
Validation Studies. LRN Laboratories
participate in multiple Proficiency