

Norfolk, Virginia, and thereby acquire shares of Bank of the Commonwealth, Norfolk, Virginia.

B. Federal Reserve Bank of Kansas City (Todd Offerbacker, Assistant Vice President) 925 Grand Avenue, Kansas City, Missouri 64198-0001:

1. *Tri-Valley Bank Shares, Inc., Lincoln, Nebraska*; to become a bank holding company by acquiring 100 percent of the voting shares of Tri-Valley Bank, Talmage, Nebraska.

Board of Governors of the Federal Reserve System, July 10, 2009.

Robert deV. Frierson,
Deputy Secretary of the Board.
[FR Doc. E9–16739 Filed 7–14–09; 8:45 am]
BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 concerning opportunity for public comment on proposed collections of information, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the information

collection plans, call the SAMHSA Reports Clearance Officer on (240) 276–1243.

Comments are invited on: (a) Whether the proposed collections of information are 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; and (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.

Proposed Project Survey of Evidence-Based Practices in State Medicaid Plans: Coverage Structures, Access and Challenges—NEW

The Substance Abuse and Mental Health Services Administration (SAMHSA) is conducting a survey to gather information about current and planned State Medicaid activities and policies related to six mental health/substance abuse evidence-based practices (EBPs). This survey is part of a five-year project to increase attention to and understanding of Medicaid mental health and substance abuse service issues among State Medicaid and Mental Health/Substance Abuse Directors, as well as improve the effectiveness of State Medicaid mental health services.

The purpose of the survey is to determine the overall management and delivery of mental health and substance abuse services within Medicaid and the use of six specific evidence-based practices. The information provided through the survey will be vital to increasing awareness and understanding of Medicaid mental health/substance abuse evidence-based practice activities. This information will also be used to develop numerous products to help State Medicaid and Mental Health/Substance Abuse Directors adopt, deliver, and refine existing policies about mental health and substance abuse EBPs.

A survey will be sent to the director of each State Medicaid office in the 50 States and the District of Columbia, with responses expected over a four-week period. The survey contains a total of 116 questions on the overall management and delivery of mental health and substance abuse services within Medicaid and on the implementation of six EBPs within the State Medicaid program. However, respondents will complete part or all of the survey, depending on how many of the six EBPs are being implemented in their State. The survey will be sent electronically to State Medicaid Directors, and they may respond by e-mail or facsimile. To reduce burden, prior to administering the survey several survey questions will be pre-completed based on existing information, as available.

Below is the table of the estimated total burden hours:

Respondent	Number of respondents	Number of responses per respondent	Average burden hour	Total burden hours
State Medicaid Directors	51	1	1	51

Send comments to Summer King, SAMHSA Reports Clearance Officer, Room 7–1044, One Choke Cherry Road, Rockville, MD 20857 and e-mail her a copy at summer.king@samhsa.hhs.gov. Written comments should be received within 60 days of this notice.

Dated: July 10, 2009.

Dennis O. Romero,
Deputy Executive Officer and Deputy Director, Office of Program Services.
[FR Doc. E9–16768 Filed 7–14–09; 8:45 am]
BILLING CODE 4162–20–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Recordkeeping/Reporting Requirements Under Emergency Review by the Office of Management and Budget (OMB)

Title: Child Care and Development Fund (CCDF) Financial Report, Form ACF–696 (States and Territories).

OMB No.: 0970–0163.

Description: States and Territories use the Financial Report Form ACF–696 for reporting Child Care and Development Fund (CCDF) expenditures. Authority to collect and report this information is

found in Section 658G of the Child Care and Development Block Grant Act of 1990, as revised. In addition to the Program Reporting Requirements set forth in 45 CFR Part 98, Subpart H, the regulations at 45 CFR 98.65(g) and 98.67(c)(1) authorize the Secretary to require financial reports as necessary.

The form provides specific data regarding claims and provides a mechanism for States to request Child Care grant awards and to certify the availability of State matching funds. Failure to collect this data would seriously compromise ACF’s ability to monitor Child Care and Development Fund expenditures. This information is also used to estimate outlays and may

be used to prepare ACF budget submissions to Congress.

The American Recovery and Reinvestment Act (ARRA) of 2009, (Pub. L. 111–5) provides an additional \$2 billion for the Child Care and Development Fund to help States, Territories, and Tribes provide child care assistance to low income working families. CCDF Program Instruction (CCDF–ACF–PI–2009–03) provided guidance on ARRA spending requirements.

Section 1512 of the ARRA legislation requires recipients to report quarterly spending and performance data on the

public Web site, “Recovery.gov”. Federal agencies are required to collect ARRA expenditure data and performance data and these data must be clearly distinguishable from the regular CCDF (non-ARRA) funds. To ensure transparency and accountability, the ARRA authorizes Federal agencies and grantees to track and report separately on expenditures from funds made available by the stimulus bill. Office of Management and Budget (OMB) guidance implementing the ARRA legislation indicates that agencies requiring additional information for

oversight should rely on existing authorities and reflect these requirements in their award terms and conditions as necessary, following existing procedures. Therefore, to capture ARRA expenditures, the ACF–696 has been modified (by the addition of a column) for reporting ARRA expenditure data. In addition, a new data element will ask States and Territories to estimate the number of child service months funded with ARRA dollars. The collection will not duplicate other information.

Respondents: States and Territories.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
CCDF State and Territory Plan	56	4	5	1,120

Estimated Total Annual Burden Hours: 1,120.

Additional Information:

ACF is requesting that OMB grant a 90 day approval for this information collection under procedures for emergency processing by July 15, 2009. A copy of this information collection, with applicable supporting documentation, may be obtained by calling the Administration for Children and Families, Reports Clearance Officer, Robert Sargis at (202) 690–7275.

Comments and questions about the information collection described above should be directed to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ACF, Office of Management and Budget, Paperwork Reduction Project, 725 17 Street, NW., Washington, DC 20503, FAX (202) 395–6974.

Dated: June 6, 2009.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. E9–16477 Filed 7–14–09; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2009–N–0296]

Agency Information Collection Activities; Proposed Collection; Comment Request; Food Labeling Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions in FDA’s food labeling regulations and on Form FDA 3570, “Model Small Business Nutrition Labeling Exemption Notice,” which small businesses may use to claim the small business exemption from nutrition labeling.

DATES: Submit written or electronic comments on the collection of information by September 14, 2009.

ADDRESSES: Submit electronic comments on the collection of information to <http://www.regulations.gov>. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of Information Management (HFA–710), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–796–3794.

SUPPLEMENTARY INFORMATION: Under the 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. “Collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA’s functions, including whether the information will have practical utility; (2) the accuracy of FDA’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques,