

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. Additional information on all bank holding companies may be obtained from the National Information Center Web site at www.ffiec.gov/nic/.

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 24, 2003.

A. Federal Reserve Bank of Kansas City (Susan Zubradt, Assistant Vice President) 925 Grand Avenue, Kansas City, Missouri 64198-0001:

1. *Sundance State Bank Profit Sharing ESOP and Trust*, Sundance Wyoming; to become a bank holding company by acquiring 28 percent of the voting shares of Sundance Bankshares, Inc., and Sundance State Bank, both of Sundance, Wyoming.

Board of Governors of the Federal Reserve System, January 24, 2003.

Jennifer J. Johnson,
Secretary of the Board.

[FR Doc. 03-2107 Filed 1-29-03; 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 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 (301) 443-7978.

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

Mandatory Guidelines for Federal Workplace Drug Testing Programs (0930-0158, revision)—SAMHSA will request renewal of OMB approval for the Federal Drug Testing Custody and Control Form for Federal agency and federally regulated drug testing programs which must comply with the HHS Mandatory Guidelines for Federal

Workplace Drug Testing Programs (59 FR 29908) dated June 9, 1994, and for the information provided by laboratories for the National Laboratory Certification Program (NLCP).

The Federal Drug Testing Custody and Control Form is used by all Federal agencies and employers regulated by the Department of Transportation to document the collection and chain of custody of urine specimens at the collection site, for laboratories to report results, and for Medical Review Officers to make a determination. The Federal Drug Testing Custody and Control Form approved by OMB three years ago will be submitted for OMB approval without any revision.

Prior to an inspection, a laboratory is required to submit specific information regarding its laboratory procedures. A major change in the submitted information requires a laboratory to provide specific information on its specimen validity testing procedures. Since all certified laboratories are expected to have the capability to conduct specimen validity tests on regulated specimens, collecting this information prior to an inspection allows the inspectors to thoroughly review and understand the laboratory's specimen validity testing procedures before arriving at the laboratory.

The NLCP application form is being revised compared to the previous form. The major change in the NLCP application form includes, where appropriate in each section, a request for specific information on the applicant laboratory's ability to conduct specimen validity testing (*i.e.*, determining if a specimen is adulterated or substituted). Since all certified laboratories are expected to have the capability to conduct specimen validity tests on regulated specimens, it is necessary to ensure that each applicant laboratory has the same capability before being certified.

The annual total burden estimates for the Federal Drug Testing Custody and Control Form, the NLCP application, the NLCP inspection checklist, and NLCP recordkeeping requirements are shown in the following table.

| Form/respondent | Burden/re-sponse (Hrs.) | Number of responses | Total annual burden (Hrs.) |
|---------------------------------------|-------------------------|---------------------|----------------------------|
| Custody and Control Form: | | | |
| Donor | .08 | 7,096,000 | 567,680 |
| Collector | .07 | 7,096,000 | 496,720 |
| Laboratory | .05 | 7,096,000 | 354,800 |
| Medical Review Officer | .05 | 7,096,000 | 354,800 |
| Laboratory Application | 3.00 | 3 | 9 |
| Laboratory Inspection Checklist | 3.00 | 110 | 330 |