

Shawneetown, Illinois, and thereby indirectly acquire Illinois One Bank, N.A., Shawneetown, Illinois.

Board of Governors of the Federal Reserve System, January 30, 1998.

**Jennifer J. Johnson,**

*Deputy Secretary of the Board.*

[FR Doc. 98-2725 Filed 2-3-98; 8:45 am]

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## FEDERAL RESERVE SYSTEM

### Notice of Proposals to Engage in Permissible Nonbanking Activities or to Acquire Companies that are Engaged in Permissible Nonbanking Activities

The companies listed in this notice have given notice under section 4 of the Bank Holding Company Act (12 U.S.C. 1843) (BHC Act) and Regulation Y, (12 CFR Part 225) to engage *de novo*, or to acquire or control voting securities or assets of a company that engages either directly or through a subsidiary or other company, in a nonbanking activity that is listed in § 225.28 of Regulation Y (12 CFR 225.28) or that the Board has determined by Order to be closely related to banking and permissible for bank holding companies. Unless otherwise noted, these activities will be conducted throughout the United States.

Each notice is available for inspection at the Federal Reserve Bank indicated. The notice also will be available for inspection at the offices of the Board of Governors. Interested persons may express their views in writing on the question whether the proposal complies with the standards of section 4 of the BHC Act.

Unless otherwise noted, comments regarding the applications must be received at the Reserve Bank indicated or the offices of the Board of Governors not later than February 19, 1998.

**A. Federal Reserve Bank of St. Louis** (Randall C. Sumner, Vice President) 411 Locust Street, St. Louis, Missouri 63102-2034:

*1. DeWitt First Bankshares Corporation, DeWitt, Arkansas;* to engage *de novo* in extending credit and servicing loans, pursuant to § 225.28(b)(1) of the Board's Regulation Y.

Board of Governors of the Federal Reserve System, January 30, 1998.

**Jennifer J. Johnson,**

*Deputy Secretary of the Board.*

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

[INFO-98-11]

### Proposed Data Collections Submitted for Public Comment and Recommendations

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call the CDC Reports Clearance Officer on (404) 639-7090.

*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; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques for other forms of information technology. Send comments to Wilma Johnson, CDC Reports Clearance Officer, 1600 Clifton Road, MS-D24, Atlanta, GA 30333. Written comments should be received within 60 days of this notice.

### Proposed Projects

#### *1. A Longitudinal Study of Lead Poisoning from the Maternal Infant Relationship Through Early Childhood—New—*

The Agency for Toxic Substances and Disease Registry (ATSDR) is mandated pursuant to the 1980 Comprehensive Environmental Response Compensation and Liability Act (CERCLA), and its 1986 Amendments, The Superfund Amendments and Reauthorization Act (SARA), to prevent or mitigate adverse human health effects and diminished quality of life resulting from exposure to hazardous substances in the environment. Lead exposure has been associated with negative pregnancy outcomes in humans, including low

birth weight, spontaneous abortion, congenital malformation, and various neurological effects in newborns and young children. The level of lead considered to be toxic has been lowered over the years by major research groups, organizations, and agencies. While lead has been shown to affect all organs, the brain or nervous system seems to be the most sensitive to lead toxicity, especially in young children. Blood lead levels as low as 10 µg/dL have been shown to result in delayed cognitive development, reduced IQ scores, and impaired hearing.

This study, originally approved by OMB in 1995, examines the long-term effects of low and marginal toxic blood lead levels in neonates and preschool African-American children in the Atlanta area. This study is divided into two components, (i) Prevalence of lead exposure in children of preschool age and (ii) longitudinal health effects of low and marginal lead exposure. These studies are conducted concurrently.

The primary focus of the prevalence study is the evaluation of the relationship between socio-economic status, elemental blood lead levels within the home environment, and blood lead levels of preschool aged children. The objective of the longitudinal study is the evaluation of the relationship between lead levels found in maternal and cord blood and adverse health effects in the infant, including deficits in behavioral, cognitive and physical development. To correlate cognitive and behavioral development with varying blood lead levels, each newborn is to undergo a series of psychometric testing at birth, then again at 6 months, 1, and 2 years of age. Evaluations of physician development will be conducted by reviewing the medical records of each newborn within the first year after birth.

This request is for a 3-year extension of the current OMB approval; however we are requesting a new OMB authority (and number) as the old number (0923-0015) will now apply only to the Substance Specific Applied Research Program (AMHPS) [King/Drew Lead Study in-Person Interview, Lead and Hypertension Screening Questionnaire/Risk Factor Questionnaire]. The requests for OMB approval for the two studies has been separated, with the King/Drew investigation retaining the old OMB number (0923-0015).