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 website 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 October 13, 2006.

A. Federal Reserve Bank of Atlanta (Andre Anderson, Vice President) 1000 Peachtree Street, N.E., Atlanta, Georgia 30309:

1. CP Capital Asset Acquisition, Inc., Miami, Florida; to become a bank holding company by acquiring 100 percent of the voting shares of Security Bank, National Association, North Lauderdale, Florida.

B. Federal Reserve Bank of Chicago (Patrick M. Wilder, Assistant Vice President) 230 South LaSalle Street, Chicago, Illinois 60690-1414:

1. Citizens Banking Corporation, Flint, Michigan; to acquire 100 percent of the voting shares of Republic Bancorp, Inc., Owosso, Michigan, and thereby indirectly acquire voting shares of Republic Bank, Lansing, Michigan.

Board of Governors of the Federal Reserve System, September 13, 2006.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. 06–7979 Filed 9–19–06; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Board of Scientific Counselors, National Institute for Occupational Safety and Health: Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting:

Name: Board of Scientific Counselors (BSC), National Institute for Occupational Safety and Health (NIOSH).

Time and Date: 9 a.m.–2:45 p.m., October 18, 2006.

Place: The Watergate Hotel, 2650 Virginia Avenue, NW., Washington, DC 20037.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: The Board shall provide advice to the Director, NIOSH on research and prevention programs. Specifically, the Board shall provide guidance on the Institute's research activities related to developing and evaluating hypotheses, systematically documenting findings and disseminating results. The Board shall evaluate the degree to which the activities of NIOSH conform to appropriate scientific standards; address current and relevant needs; and produce intended results.

Matters To Be Discussed: Agenda items include a health hazard evaluation program review, an update on the economics of occupational safety and health, and an update on flavorings-related lung disease prevention efforts.

Agenda items are subject to change as priorities dictate.

Contact Person For More Information: Roger Rosa, Executive Secretary, BSC, NIOSH, 200 Independence Avenue, SW., Room 715H, Washington, DC 20201, telephone (202) 205–7856, fax (202) 260–4464.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: September 13, 2006.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 06–7984 Filed 9–19–06; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects

Title: DHHS/ACF/ASPE/DOL Enhanced Services for the Hard-to-Employ Demonstration and Evaluation: Rhode Island 15-Month Survey Amendment.

OMB No.: 0970–0276.

Description: The Enhanced Services for the Hard-to-Employ Demonstration and Evaluation Project (HtE) seeks to learn what works in this area to date and is explicitly designed to build on past research by rigorously testing a wide variety of approaches to promote employment and improve family functioning and child well-being. The HtE project is designed to help Temporary Assistance for Needy Families (TANF) recipients, former TANF recipients, or low-income parents who are hard-to-employ. The project is sponsored by the Office of Planning, Research and Evaluation (OPRE) of the Administration for Children and Families (ACF), the Office of the Assistant Secretary for Planning and Evaluation (ASPE) in the U.S. Department of Health and Human Services (HHS), and the U.S. Department of Labor (DOL).

The evaluation involves an experimental, random assignment design in four sites, testing a diverse set of strategies to promote employment for low-income parents who face serious obstacles to employment. The four include: (1) Intensive care management to facilitate the use of evidence-based treatment for major depression among parents receiving Medicaid in Rhode Island; (2) job readiness training, worksite placements, job coaching, job development and other training opportunities for recent parolees in New York City; (3) pre-employment services and transitional employment for long-term TANF participants in Philadelphia; and (4) home- and center-based care, enhanced with self-sufficiency services, for low-income families who have young children or are expecting in Kansas and Missouri.

Materials for follow-up surveys for each of these sites were previously submitted to OMB and were approved. The purpose of this submission is to add physiological measures to the follow-up effort to the Rhode Island study.

Respondents: The respondents to this component of the Rhode Island follow-

up survey will be low-income parents and their children from the Rhode Island site currently participating in the HtE Project. As described in the prior OMB submission, these parents are Medicaid recipients between the ages of 18 and 45 receiving Medicaid through the managed care provider United Behavioral Health (UBH) in Rhode Island who meet study criteria with regard to their risk for depression. Children are the biological, adopted, and stepchildren of these parents, between 1 and 18 years of age.

The annual burden estimates are detailed below, and the substantive content of each component will be detailed in the supporting statement attached to the forthcoming 30-day notice.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
RI 15-month, parent physiological component	400	8	5 minutes or .08 hrs	266.66
RI 15-month, young child physiological component	160	8	5 minutes or .08 hrs	106.66
RI 15-month, youth physiological component	242	8	5 minutes or .08 hrs	161.33

Estimated Total Annual Burden Hours: 534.65.

In compliance with the requirements of Section 3506(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 information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: September 13, 2006.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 06–7763 Filed 9–19–06; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Office of Family Assistance; Single-Source Program Expansion Supplement

AGENCY: Office of Family Assistance, Administration for Children and Families, HHS.

CFDA#: 93.575.

Legislative Authority: Child Care and Development Block Grant Act of 1990, as amended.

Amount of Award: $101,774.00 for one year.

Project Period: 09/30/2006–09/29/2007.

Justification for the Exception to Competition: Oregon State University (the grantee) is currently conducting data analyses with funding from a research grant awarded in FY 2004 to validate methodologies used to conduct State market rate surveys on the price for child care and early education programs at the State and local levels. The supplemental funds will allow the grantee to include additional datasets in the ongoing analyses representing sampling methodologies that include a more diverse care provider sample, a broader geographical coverage, and several additional data collection methods, and will in turn make the findings from the project more generalizable to States, Tribes and Territories implementing the Child Care and Development Fund program.

CONTACT FOR FURTHER INFORMATION: Ivelisse Martinez-Beck, Research Coordinator, Child Care Bureau, Portals Building, Suite 800, 1250 Maryland Avenue, SW., Washington, DC 20024.

Telephone: 202–690–7885.

Dated: September 1, 2006.

Sidonie Squier,

Director, Office of Family Assistance.

[FR Doc. E6–15559 Filed 9–19–06; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004E–0040]

Determination of Regulatory Review Period for Purposes of Patent Extension; CYDECTIN

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) has determined the regulatory review period for CYDECTIN and is publishing this notice of that determination as required by law. FDA has made the determination because of the submission of an application to the Director of Patents and Trademarks, Department of Commerce, for the extension of a patent which claims that animal drug product.

ADDRESSES: Submit written comments and petitions to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments.*

FOR FURTHER INFORMATION CONTACT: Beverly Friedman, Office of Regulatory Policy (HFD–7), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane,Rockville, MD 20857, 301–594–2041.

SUPPLEMENTARY INFORMATION: The Drug Price Competition and Patent Term Restoration Act of 1984 (Public Law 98–417) and the Generic Animal Drug and