FEDERAL DEPOSIT INSURANCE CORPORATION

Notice to All Interested Parties of the Termination of the Receivership of 10304, The First National Bank of Barnesville, Barnesville, GA

Notice is hereby given that the Federal Deposit Insurance Corporation (''FDIC'') as Receiver for The First National Bank of Barnesville, Barnesville, GA (''the Receiver'') intends to terminate its receivership for said institution. The FDIC was appointed receiver of The First National Bank of Barnesville on October 22, 2010. The liquidation of the receivership assets has been completed. To the extent permitted by available funds and in accordance with law, the Receiver will be making a final dividend payment to proven creditors.

Based upon the foregoing, the Receiver has determined that the continued existence of the receivership will serve no useful purpose. Consequently, notice is given that the receivership shall be terminated, to be effective no sooner than thirty days after the date of this Notice. If any person wishes to comment concerning the termination of the receivership, such comment must be made in writing and sent within thirty days of the date of this Notice to: Federal Deposit Insurance Corporation, Division of Resolutions and Receiverships, Attention: Receivership Oversight Department 32.1, 1601 Bryan Street, Dallas, TX 75201.

No comments concerning the termination of this receivership will be considered which are not sent within this time frame.

Dated: June 26, 2015.

Federal Deposit Insurance Corporation.

Robert E. Feldman,

Executive Secretary.

[FR Doc. 2015–16161 Filed 6–30–15; 8:45 am]

BILLING CODE 6714–01–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 July 27, 2015.

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

1. *Atlantic Capital Bancshares, Inc.,* Atlanta, Georgia; to merge with First Security Group, Inc., and thereby acquire FSGBank, NA, both in Chattanooga, Tennessee.

In connection with this applicantion, Atlantic Capital Bancshares' parent companies BankCap Equity Fund, LLC; BankCap Partners GP L.P.; BankCap Partners Fund I, L.P.; and BCP Fund I Southeast Holdings, LLC, all in Dallas, Texas, will indirectly acquire First Security Group, Inc., and FSGBank, NA, both in Chattanooga, Tennessee.

Board of Governors of the Federal Reserve System, June 26, 2015.

Margaret McCloskey Shanks,

Deputy Secretary of the Board.

[FR Doc. 2015–16157 Filed 6–30–15; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: Goal-Oriented Adult Learning in Self-Sufficiency Study.

OMB No.: New Collection.

Description: The Administration for Children and Families (ACF) is proposing a data collection activity as part of the Goal-Oriented Adult Learning in Self-Sufficiency (GOALS) study. The purpose of the GOALS project is to address the nexus between the growing knowledge base in the psychological sciences and long-standing approaches to self-sufficiency programs targeted to adults and young adults. The project will explore the programmatic implications of existing research on psychological processes associated with goal-directed behaviors, including socio-emotional regulation and cognitive skills, executive functioning, and related areas. The project will synthesize current research on these topics; address how insights gained from research can be used to promote economic advancement among low-income populations, identify promising strategies, or strengthen underlying skills in these areas; and inform measurement of changes and developments in skill acquisition.

The proposed information collection activity consists of exploratory calls with program directors and administrators, semi-structured interviews with key program staff and community partner organization staff, and focus group discussions with program participants. ACF seeks to gain an in-depth, systematic understanding of program administration and implementation, service delivery and operation, outputs and outcomes, and identify promising practices and other areas for further study.

Respondents: Key program directors and administrators, program staff and community partner organization staff, and program participants at selected program sites.

ANNUAL BURDEN ESTIMATES

Instrument	Total Number of Respondents	Annual Number of Respondents	Number of Responses per Respondent	Average Burden Hours per Response	Annual Burden Hours
Exploratory telephone call semi-structured interview—program directors and administrators	24	12	1	1	12

ANNUAL BURDEN ESTIMATES—Continued

Instrument	Total Number of Respondents	Annual Number of Respondents	Number of Responses per Respondent	Average Burden Hours per Response	Annual Burden Hours
Site visit semi-structured interview—program staff and community partner organization staff	180	90	1	1.25	113
Site visit group discussion—program participants	84	42	1	1.25	53
Estimated Total Annual Burden Hours					178

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 Planning, Research and Evaluation, 370 L'Enfant Promenade SW., Washington, DC 20447, Attn: OPRE Reports Clearance Officer. Email address: *OPREinfocollection@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.

Karl Koerper,

Reports Clearance Officer.

[FR Doc. 2015–16073 Filed 6–30–15; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0742]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by July 31, 2015.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0045. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Registration of Producers of Drugs and Listing of Drugs in Commercial Distribution—21 CFR Part 207

OMB Control Number 0910–0045—Extension

Requirements for drug establishment registration and drug listing are set forth in section 510 of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 360), section 351 of the Public Health Service Act (42 U.S.C. 262), and part 207 (21 CFR part 207). Fundamental to FDA's mission to protect the public health is the collection of this information, which is used for important activities such as postmarket surveillance for serious adverse drug reactions, inspection of drug manufacturing and processing facilities, and monitoring of drug products imported into the United States. Comprehensive, accurate, and up to date information is critical to conducting these activities with efficiency and effectiveness.

Under section 510 of the FD&C Act, FDA is authorized to establish a system for registration of producers of drugs and for listing of drugs in commercial distribution. To implement section 510 of the FD&C Act, FDA issued part 207. Under current § 207.20, manufacturers, repackers, and relabelers that engage in the manufacture, preparation, propagation, compounding, or processing of human or veterinary drugs and biological products, including bulk drug substances and bulk drug substances for prescription compounding, and drug premixes as well as finished dosage forms, whether prescription or over-the-counter, are required to register their establishment. In addition, manufacturers, repackers, and relabelers are required to submit a listing of every drug or biological product in commercial distribution. Owners or operators of establishments that distribute under their own label or trade name a drug product manufactured by a registered establishment are not required either to register or list. However, distributors may elect to submit drug listing information in lieu of the registered