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

1. Community Bank Investors of America, LP, and FA Capital, LLC, both of Midlothian, Virginia; to acquire 22.95 percent of the outstanding voting shares of Silvergate Capital Corporation, and thereby indirectly acquire Silvergate Bank, both of La Jolla, California.

2. Silvergate Capital Corporation; to become a bank holding company by acquiring 100 percent of the voting shares of Silvergate Bank, both of La Jolla, California.

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

1. First Fontanelle Employee Stock Ownership Plan and Trust, Fontanelle, Iowa; to become a bank holding company by acquiring 30.44 percent of First Fontanelle Bancorporation, Fontanelle, Iowa, and thereby indirectly acquire First National Bank, Fontanelle, Iowa.

Board of Governors of the Federal Reserve System, February 13, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E8–2998 Filed 2–15–08; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Head Start Program Information Report.

OMB No.: 0980–0017.

Description: The Office of Head Start within the Administration for Children and Families, United States Department of Health and Human Services, is proposing to renew authority to collect information using the Head Start Program Information Report (PIR). The PIR provides information about Head Start and Early Head Start services received by the children and families enrolled in Head Start programs. The information collected in the PIR is used to inform the public about these programs and to make periodic reports to Congress about the status of children in Head Start programs as required by the Head Start Act.

Respondents: Head Start and Early Head Start program grant recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Head Start Program Information Report	2,690	1	4	10,760

Estimated Total Annual Burden Hours: 10,760

Additional Information

Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment

OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: February 11, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. 08–696 Filed 2–15–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB review; comment request

Title: Required Data Elements for Voluntary Establishment of Paternity Affidavits.

OMB No.: 0970–0171.

Description: Section 466(a)(5)(C)(iv) of the Social Security Act (the Act) requires States to develop and use an affidavit for the voluntary acknowledgement of paternity. The affidavit for the voluntary acknowledgement of paternity must include the minimum requirements specified by the Secretary under section 452(a)(7) of the Act. The affidavits will be used by hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program.

Respondents: State and Tribal IV–D agencies, hospitals, birth record agencies and other entities participating in the voluntary paternity establishment program.

Annual Burden Estimates

Estimated Total Annual Burden Hours:

Additional Information

Copies of the proposed collection may be obtained 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. All requests should be identified by the title of the information collection. E-mail address: *infocollection@acf.hhs.gov.*

OMB Comment

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Management and Budget, Paperwork Reduction Project, Fax: 202–395–6974, Attn: Desk Officer for the Administration for Children and Families.

Dated: February 11, 2008.

Janean Chambers,

Reports Clearance Officer.

[FR Doc. 08–697 Filed 2–15–08; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[FDA No. 225–08–8002]

Memorandum of Understanding Between the Division of Select Agents and Toxins Center for Disease Control and Prevention and the Food and Drug Administration

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is providing notice of a memorandum of understanding (MOU) between FDA and the Division of Select Agents and Toxins (DSAT) of the Centers for Disease Control and Prevention (CDC). The purpose of this MOU is to establish a procedure to allow CDC/DSAT to confirm that FDA has accepted or approved, under the authority of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 301 et seq), an Investigational New Drug application (IND), a request to establish an Investigational New Animal Drug file (INAD), or an Investigational Device Exemption application (IDE) for a clinical trial involving the use of an investigational product that is, bears, or contains a select agent or toxin.

DATES: The agreement became effective January 25, 2008.

FOR FURTHER INFORMATION CONTACT: Jarilyn Dupont, Director of Regulatory Policy, Office of Policy and Planning (HF–11), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–5906.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 20.108(c), which states that all written agreements and MOUs between FDA and others shall be published in the **Federal Register**, the agency is publishing notice of this MOU.

Dated: February 8, 2008.

Jeffrey Shuren,

Assistant Commissioner for Policy.

BILLING CODE 4160–01–S