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SUPPLEMENTARY INFORMATION:

I. General Information

A. Does this action apply to me?

This action is directed to the public in general. As such, the Agency has not attempted to describe the specific entities that this action may apply to. Although others may be affected, this action applies directly to the submitter of the request for waiver. If you have any questions regarding the applicability of this action to a particular entity, consult the technical person listed under **FOR FURTHER INFORMATION CONTACT**.

B. What should I consider as I prepare my comments for EPA?

1. *Submitting CBI.* Do not submit this information to EPA through regulations.gov or e-mail. Clearly mark the part or all of the information that you claim to be CBI. For CBI information in a disk or CD–ROM that you mail to EPA, mark the outside of the disk or CD–ROM as CBI and then identify electronically within the disk or CD–ROM the specific information that is claimed as CBI. In addition to one complete version of the comment that includes information claimed as CBI, a copy of the comment that does not contain the information claimed as CBI must be submitted for inclusion in the public docket. Information so marked will not be disclosed except in accordance with procedures set forth in 40 CFR part 2.

2. *Tips for preparing your comments.* When submitting comments, remember to:

- i. Identify the document by docket ID number and other identifying information (subject heading, **Federal Register** date and page number).
- ii. Follow directions. The Agency may ask you to respond to specific questions or organize comments by referencing a Code of Federal Regulations (CFR) part or section number.

iii. Explain why you agree or disagree; suggest alternatives and substitute language for your requested changes.

iv. Describe any assumptions and provide any technical information and/or data that you used.

v. If you estimate potential costs or burdens, explain how you arrived at your estimate in sufficient detail to allow for it to be reproduced.

vi. Provide specific examples to illustrate your concerns and suggest alternatives.

vii. Explain your views as clearly as possible, avoiding the use of profanity or personal threats.

viii. Make sure to submit your comments by the comment period deadline identified.

II. Background

A. What action is the agency taking?

EPA is announcing receipt of a request for waiver from testing from 3M. EPA will accept comments on this request and will publish another **Federal Register** notice on or before November 14, 2011, announcing its decisions on this request. See 40 CFR 766.32(c).

B. What is the Agency's authority for taking this action?

Under 40 CFR part 766, EPA requires testing of certain chemical substances to determine whether they may be contaminated with HDDs and HDFs. Under 40 CFR 766.32(a)(2)(i), a waiver may be granted if a responsible company official certifies that the chemical substance is produced only in quantities of 100 kilograms (kg) or less per year, and only for research and development purposes.

Under 40 CFR 766.32(b), a request for a waiver must be made 60 days before resumption of manufacture or importation of a chemical substance not being manufactured, imported, or processed as of June 5, 1987.

On September 14, 2011, EPA received a waiver request from 3M under 40 CFR 766.32(a)(2)(i). The request indicates that 3M intends to import tetrabromobisphenol A (CASRN 79–94–7), a chemical substance subject to testing under 40 CFR part 766, as part of an experimental formulation for research and development purposes. 3M will import less than 100 kg of tetrabromobisphenol A.

List of Subjects

Environmental protection, Dibenzofurans, Dioxins, Hazardous substances, Tetrabromobisphenol A.

Dated: September 22, 2011.

Wendy C. Hamnett,

Director, Office of Pollution Prevention and Toxics.

[FR Doc. 2011–24870 Filed 9–23–11; 11:15 am]

BILLING CODE 6560–50–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 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.

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 21, 2011.

A. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) P.O. Box 442, St. Louis, Missouri 63166–2034:

1. *First Clover Leaf Financial Corp.*, Edwardsville, Illinois; to become a bank holding company through the conversion of First Clover Leaf Bank, Edwardsville, Illinois, from a federally chartered savings bank to a state chartered commercial bank.

Board of Governors of the Federal Reserve System.

Dated: September 22, 2011.

Jennifer J. Johnson,

Secretary of the Board.

[FR Doc. 2011-24753 Filed 9-26-11; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS-0990-New; 30-day notice]

Agency Information Collection Request. 30 Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of

automated collection techniques or other forms of information technology to minimize the information collection burden.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to Sherette.funncoleman@hhs.gov, or call the Reports Clearance Office on (202) 690-5683. Send written comments and recommendations for the proposed information collections within 30 days of this notice directly to the OS OMB Desk Officer; faxed to OMB at 202-395-5806.

Proposed Project: The Children's Health Insurance Program Reauthorization Act (CHIPRA) 10-State Evaluation (New)—OMB No. 0990-NEW—Assistant Secretary Planning and Evaluation (ASPE).

Abstract: The Office of the Assistant Secretary for Planning and Evaluation (ASPE) is requesting the Office of Management and Budget (OMB) approval on a new collection to provide the federal government with new and detailed insights into how the Children's Health Insurance Program (CHIP) has evolved since its early years, what impacts on children's coverage and access to care have occurred, and

what new issues have arisen as a result of policy changes related to CHIPRA and the Patient Protection and Affordable Care Act (The Affordable Care Act) of 2010 (Pub. L. 111-148). The evaluation will address numerous key questions regarding the structure and impact of CHIP and Medicaid programs for children. To answer these questions, ASPE will draw on three new primary data collection efforts, including a survey of selected CHIP enrollees and disenrollees in 10 states (and Medicaid enrollees and disenrollees in 3 of these states), qualitative case studies in the 10 states, and a survey of State Program Administrators in all 50 States and the District of Columbia. This current request seeks clearance for the first two information collections; ASPE will seek clearance for the third information collection at a later date. All data collection will take place one time only over a three year period. The survey component includes a sample of children in 10 selected states, recently enrolled or disenrolled in CHIP or Medicaid. Survey data will be collected using computer-assisted telephone interviewing with an in-person follow-up. The qualitative case studies will include site visit interviews with CHIP and Medicaid administrators and public and child health stakeholders, plus focus groups with parents or family members of CHIP enrollees.

ESTIMATED ANNUALIZED BURDEN TABLE

Forms	Type of respondent	Number of respondents	Number of responses per respondent	Average burden (in hours) per response	Total burden hours
Survey of CHIP Enrollees and Disenrollees.	CHIP enrollees and disenrollees	15,000	1	30/60	7,500
Survey of Medicaid Enrollees and Disenrollees.	Medicaid enrollees and disenrollees	4,500	1	30/60	2,250
Site Visits	CHIP and Medicaid personnel—1 ...	300	1	1	300
Focus Groups	Parents and other family members of children—2	80	1	2	160
Total Burden					10,210

Keith Tucker,

*Paperwork Reduction Act Clearance Officer,
Office of the Secretary.*

[FR Doc. 2011-24721 Filed 9-26-11; 8:45 am]

BILLING CODE 4150-05-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Final Effect of Designation of a Class of Employees for Addition to the Special Exposure Cohort

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: HHS gives notice concerning the final effect of the HHS decision to designate a class of employees from the Sandia National Laboratories, Albuquerque, New Mexico, as an addition to the Special Exposure Cohort (SEC) under the Energy Employees Occupational Illness Compensation Program Act of 2000. On July 29, 2011, as provided for under 42 U.S.C. 7384q(b), the Secretary of HHS designated the following class of employees as an addition to the SEC: