

Governors not later than September 6, 2016.

A. Federal Reserve Bank of Atlanta (Chapelle Davis, Assistant Vice President) 1000 Peachtree Street NE., Atlanta, Georgia 30309. Comments can also be sent electronically to Applications.Comments@atl.frb.org:

1. *Pinnacle Financial Corporation, Elberton, Georgia*, to acquire 100 percent of the outstanding voting stock of Independence Bank of Georgia, Braselton, Georgia.

Board of Governors of the Federal Reserve System, August 5, 2016.

Margaret McCloskey Shanks,
Deputy Secretary of the Board.

[FR Doc. 2016–19059 Filed 8–10–16; 8:45 am]

BILLING CODE 6210–01–P

FEDERAL RESERVE SYSTEM

Formations of, Acquisitions by, and Mergers of Savings and Loan Holding Companies

The companies listed in this notice have applied to the Board for approval, pursuant to the Home Owners' Loan Act (12 U.S.C. 1461 *et seq.*) (HOLA), Regulation LL (12 CFR part 238), and Regulation MM (12 CFR part 239), and all other applicable statutes and regulations to become a savings and loan holding company and/or to acquire the assets or the ownership of, control of, or the power to vote shares of a savings association and nonbanking companies owned by the savings and loan 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 HOLA (12 U.S.C. 1467a(e)). 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 10(c)(4)(B) of the HOLA (12 U.S.C. 1467a(c)(4)(B)). 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 September 6, 2016.

A. Federal Reserve Bank of Cleveland (Nadine Wallman, Vice President) 1455

East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to

Comments.applications@clev.frb.org:

1. *Tinka K. Powell 2016 Family Trust; Tinka K. Powell 2016 Family Trust fbo John W. Powell; Tinka K. Powell 2016 Family Trust fbo Mark W. Powell; Tinka K. Powell 2016 Family Trust fbo Ryan J. Powell; and James R. Powell 2016 Family Trust*, all of Dayton, Ohio all to become savings and loan holding companies by acquiring of more than 25 percent of the total equity of Liberty Capital, Inc., Wilmington, Ohio, and thereby acquire control of Liberty Savings Bank, FSB, Wilmington, Ohio.

Board of Governors of the Federal Reserve System, August 5, 2016.

Margaret McCloskey Shanks,
Deputy Secretary of the Board.

[FR Doc. 2016–19060 Filed 8–10–16; 8:45 am]

BILLING CODE 6210–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket Number CDC–2016–0074; NIOSH 156–B]

National Institute for Occupational Safety and Health Draft Immediately Dangerous to Life or Health (IDLH) Value Profile for Peracetic Acid (CAS #79–21–0)

AGENCY: National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of draft document for public comment.

SUMMARY: On May 1, 2015, the Director of the National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC), published a notice in the **Federal Register** [80 FR 24930] announcing the availability of and a request for comments for the draft immediately dangerous to life or health (IDLH) values and support technical documents, entitled IDLH Values Profiles, for 14 chemicals. Written comments were to be received before the end of the comment period on June 30, 2015. Due to subsequent requests from the public, this Notice announces that NIOSH is seeking further comments on the draft IDLH Value Profile for peracetic acid (CAS #79–21–0) <http://www.cdc.gov/>

[niosh/docket/review/docket156a/pdfs/g1-013-peracetic-acid-cas-79-21-0.pdf](http://www.cdc.gov/niosh/docket/review/docket156a/pdfs/g1-013-peracetic-acid-cas-79-21-0.pdf) for an additional 60 days.

DATES: Electronic or written comments on the draft IDLH Value Profile for peracetic acid must be received by October 11, 2016.

ADDRESSES: You may submit comments, identified by CDC–2016–0074 and docket number NIOSH 156–B, by either of the two following methods:

- **Federal eRulemaking Portal:** www.regulations.gov. Follow the instructions for submitting comments.
- **Mail:** National Institute for Occupational Safety and Health, NIOSH Docket Office, 1090 Tusculum Avenue, MS C–34, Cincinnati, OH 45226.

Instructions: All information received in response to this notice must include the agency name and docket number [CDC–2016–0074; NIOSH 156–B]. All relevant comments received will be posted without change to www.regulations.gov, including any personal information provided. For access to the docket to read background documents or comments received, go to www.regulations.gov.

FOR FURTHER INFORMATION CONTACT: G. Scott Dotson, NIOSH, Education and Information Division, Robert A. Taft Laboratories, 1090 Tusculum Avenue, MS C–32, Cincinnati, Ohio 45226, telephone (513) 533–8540 (not a toll free number).

SUPPLEMENTARY INFORMATION: The proposed IDLH value and draft IDLH Value Profile for peracetic acid is based on the process outlined in the NIOSH Current Intelligence Bulletin 66—Derivation of Immediately Dangerous to Life or Health (IDLH) Values <http://www.cdc.gov/niosh/docs/2014-100/pdfs/2014-100.pdf>. The draft IDLH Value Profile was developed to provide the scientific rationale behind the derivation of the proposed IDLH value for peracetic acid. This includes a detailed summary of the health hazards of acute exposure to high airborne concentrations of peracetic acid and the rationale for the proposed IDLH value for peracetic acid.

To facilitate the review of this draft document, NIOSH requests that the following questions be taken into consideration:

1. Does this document clearly outline the health hazards associated with acute (or short-term) exposures to peracetic acid? If not, what specific information is missing from the document?
2. Are the rationale and logic behind the derivation of an IDLH value for peracetic acid clearly explained? If not, what specific information is needed to clarify the basis of the IDLH value?

3. Are the conclusions supported by the data?

4. Are the tables clear and appropriate?

5. Is the document organized appropriately? If not, what improvements are needed?

6. Are you aware of any scientific data reported in governmental publications, databases, peer-reviewed journals, or other sources that should be included within this document?

John Howard,

Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention.

[FR Doc. 2016-19051 Filed 8-10-16; 8:45 am]

BILLING CODE 4163-19-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2013-N-0115]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Manufactured Food Regulatory Program Standards

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 September 12, 2016.

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 0910-0601. 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, Three White Flint North, 10A63, 11601 Landsdown St., North Bethesda, MD 20852, 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.

Manufactured Food Regulatory Program Standards—OMB Control Number 0910-0601—Extension

In the **Federal Register** of July 20, 2006 (71 FR 41221), FDA announced the availability of a draft document entitled “Manufactured Food Regulatory Program Standards (MFRPS).” These program standards have since been finalized and updated multiple times. The current standards are the framework that States should use to design and manage their manufactured food programs. The current version expires on September 30, 2016, and FDA is proposing to update and submit for issuance with a new expiration date. The current and proposed versions of the standards are available at the docket number identified in brackets at the heading of this document. Persons with

access to the Internet may submit email requests for a single copy of the draft manufactured food standards to OP-ORA@fda.hhs.gov. There are 42 State programs enrolled, in which each State may receive up to \$300,000 each year for a period of 5 years provided there is significant conformance with the 10 standards.

In the first year of implementing the program standards, the State program conducts a baseline self-assessment to determine if it meets the elements of each standard. The State program should use the worksheets and forms contained in the draft program standards; however, it can use alternate forms that are equivalent. The State program maintains the documents and verifies records required for each standard. The information contained in the documents must be current and fit-for-use. If the State program fails to meet all program elements and documentation requirements of a standard, it develops a strategic improvement plan that includes the following: (1) The individual program element or documentation requirement of the standard that was not met, (2) improvements needed to meet the program element or documentation requirement of the standard, and (3) projected completion dates for each task.

In the **Federal Register** of February 12, 2016 (81 FR 7544), FDA published a 60-day notice requesting public comment on the proposed collection of information. We received two comments. However, these comments did not address the information collection.

FDA estimates the burden of this collection of information as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN ¹

Respondent	Number of respondents	Number of responses per respondent	Total annual responses	Average burden per response	Total hours
State Departments of Agriculture or Health	42	1	42	376	15,792

¹ There are no capital costs or operating and maintenance costs associated with this collection of information.