

CERCLA costs for a cleanup action performed by the EPA at the Site.

**DATES:** The Agency will consider public comments on the settlement until July 24, 2017. The Agency will consider all comments received and may modify or withdraw its consent to the proposed settlement if comments received disclose facts or considerations which indicate that the proposed settlement is inappropriate, improper, or inadequate.

**ADDRESSES:** Copies of the settlement are available from the Agency by contacting Ms. Paula V. Painter, Program Analyst, using the contact information provided in this document. Comments may also be submitted by referencing the site's name through one of the following methods:

*Internet:* <https://www.epa.gov/aboutepa/about-epa-region-4-southeast#r4-public-notices>.

• *U.S. Mail:* U.S. Environmental Protection Agency, Superfund Division, Attn: Paula V. Painter, 61 Forsyth Street SW., Atlanta, Georgia 30303.

• *Email:* [painter.paula@epa.gov](mailto:painter.paula@epa.gov).

**FOR FURTHER INFORMATION CONTACT:** Paula V. Painter at (404) 562-8887.

Dated: February 8, 2017.

Anita L. Davis,

Chief, Enforcement and Community Engagement Branch, Superfund Division.

[FR Doc. 2017-13063 Filed 6-21-17; 8:45 am]

**BILLING CODE 6560-50-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 July 18, 2017.

*A. Federal Reserve Bank of Philadelphia* (William Spaniel, Senior Vice President) 100 North 6th Street, Philadelphia, Pennsylvania 19105-1521. Comments can also be sent electronically to

[Comments.applications@phil.frb.org](mailto:Comments.applications@phil.frb.org):

1. *Seneca Financial MHC*, Baldwinsville, New York; to become a federal mutual holding company, and Seneca Financial Corp., Baldwinsville, New York, to become a savings and loan holding company, by acquiring 100 percent of Seneca Savings Bank, Baldwinsville, New York, following the conversion of Seneca Federal Savings and Loan Association, Baldwinsville, New York, from a federal mutual savings association to a federal stock savings association to be called Seneca Savings Bank, Baldwinsville, New York.

Board of Governors of the Federal Reserve System, June 19, 2017.

Ann E. Misback,

Secretary of the Board.

[FR Doc. 2017-13046 Filed 6-21-17; 8:45 am]

**BILLING CODE 6210-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

[Document Identifier: CMS-116]

### Agency Information Collection Activities: Proposed Collection; Comment Request

**AGENCY:** Centers for Medicare & Medicaid Services, HHS.

**ACTION:** Notice.

**SUMMARY:** The Centers for Medicare & Medicaid Services (CMS) is announcing an opportunity for the public to comment on CMS' intention to collect information from the public. Under the Paperwork Reduction Act of 1995 (the PRA), federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information (including each proposed

extension or reinstatement of an existing collection of information) and to allow 60 days for public comment on the proposed action. Interested persons are invited to send comments regarding our burden estimates or any other aspect of this collection of information, including the necessity and utility of the proposed information collection for the proper performance of the agency's functions, the accuracy of the estimated burden, ways to enhance the quality, utility, and clarity of the information to be collected, and the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

**DATES:** Comments must be received by August 21, 2017.

**ADDRESSES:** When commenting, please reference the document identifier or OMB control number. To be assured consideration, comments and recommendations must be submitted in any one of the following ways:

1. *Electronically.* You may send your comments electronically to <http://www.regulations.gov>. Follow the instructions for "Comment or Submission" or "More Search Options" to find the information collection document(s) that are accepting comments.

2. *By regular mail.* You may mail written comments to the following address: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Document Identifier/OMB Control Number \_\_\_\_, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, you may make your request using one of following:

1. Access CMS' Web site address at <https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.html>.

2. Email your request, including your address, phone number, OMB number, and CMS document identifier, to [Paperwork@cms.hhs.gov](mailto:Paperwork@cms.hhs.gov).

3. Call the Reports Clearance Office at (410) 786-1326.

**FOR FURTHER INFORMATION CONTACT:** William Parham at (410) 786-4669.

### SUPPLEMENTARY INFORMATION:

#### Contents

This notice sets out a summary of the use and burden associated with the following information collections. More detailed information can be found in each collection's supporting statement

and associated materials (see **ADDRESSES**).

**CMS-116 Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations**

Under the PRA (44 U.S.C. 3501–3520), federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. The term “collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA requires federal agencies to publish a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension or reinstatement of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, CMS is publishing this notice.

**Information Collection**

1. *Type of Information Collection Request*: Revision of a currently approved collection; *Title of Information Collection*: Clinical Laboratory Improvement Amendments (CLIA) Application Form and Supporting Regulations; *Use*: The application must be completed by entities performing laboratory’s testing specimens for diagnostic or treatment purposes. This information is vital to the certification process. In this revision, the majority of changes were minor changes to the form and

accompanying instructions to facilitate the completion and data entry of the form. However, we added the collection of identifying the non-waived testing to be performed to section VIII of the form. We anticipate that the change to section VIII will take an average of 15 additional minutes to complete. *Form Number*: CMS-116 (OMB Control Number: 0938–0581); *Frequency*: Biennially and Occasionally; *Affected Public*: Private Sector—Business or other for-profits and Not-for-profit institutions; *Number of Respondents*: 42,000; *Total Annual Responses*: 51,000; *Total Annual Hours*: 51,000. (For policy questions regarding this collection contact Kathleen Todd at 410–786–3385.)

Dated: June 19, 2017.

**William N. Parham, III,**

*Director, Paperwork Reduction Staff, Office of Strategic Operations and Regulatory Affairs.*

[FR Doc. 2017–13070 Filed 6–21–17; 8:45 am]

**BILLING CODE 4120–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Children and Families**

[OMB No.: New]

**Proposed Information Collection Activity; Comment Request; Medical Complaint Form, Contact Investigation Form: Non-TB Illness, and Contact Investigation Form: Active/Suspect TB**

*Description*: The Administration for Children and Families’ Office of Refugee Resettlement (ORR) places unaccompanied minors in their custody

in licensed care provider facilities until reunification with a qualified sponsor. Care provider facilities are required to provide children with services such as classroom education, mental health services, and health care. Pursuant to Exhibit 1, part A.2 of the Flores Settlement Agreement (Jenny Lisette Flores, et al., v. Janet Reno, Attorney General of the United States, et al., Case No. CV 85–4544–RJK (C.D. Cal. 1996), care provider facilities, on behalf of ORR, shall arrange for appropriate routine medical and dental care, family planning services, and emergency health care services, including a complete medical examination (including screening for infectious disease) within 48 hours of admission, excluding weekends and holidays, unless the minor was recently examined at another facility; appropriate immunizations in accordance with the U.S. Public Health Service (PHS), Center for Disease Control; administration of prescribed medication and special diets; appropriate mental health interventions when necessary for each minor in their care.

The forms are to be used as worksheets for healthcare providers and health departments to compile information that would otherwise have been collected during a medical evaluation. Once completed, the forms will be given to care provider program staff for data entry into ORR’s electronic data repository known as ‘The UAC Portal’. Data will be used to record UC health conditions/illnesses and for case management of any identified illnesses/conditions.

*Respondents*: Office of Refugee Resettlement Grantee staff.

**ANNUAL BURDEN**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Medical Complaint Form .....	120	2,507	.13	39,109
Contact Investigation Form: Non-TB Illness .....	120	4	.08	38
Contact Investigation Form: Suspect or Active TB .....	120	2	.08	19

*Estimated Total Annual Burden Hours*: 39,166.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap 35), 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, 330 C Street SW., Washington DC 20201. Attn: ACF Reports Clearance Officer. Email address: [infocollection@acf.hhs.gov](mailto: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.