interpreted by 46 CFR 545.5. Complainant alleges this violation arose from the assessment of detention charges during periods when empty containers could not be returned due to reasons such as dual transaction requirements and appointment unavailability.

An answer to the complaint must be filed with the Commission within 25 days after the date of service.

The full text of the complaint can be found in the Commission's electronic Reading Room at *https://www2.fmc.gov/readingroom/proceeding/24-24/.* This proceeding has been assigned to the Office of Administrative Law Judges. The initial decision of the presiding judge shall be issued by July 21, 2025, and the final decision of the Commission shall be issued by February 4, 2026.

David Eng,

Secretary.

[FR Doc. 2024–16260 Filed 7–23–24; 8:45 am]

BILLING CODE 6730–02–P

FEDERAL RESERVE SYSTEM

Change in Bank Control Notices; Acquisitions of Shares of a Bank or Bank Holding Company

The notificants listed below have applied under the Change in Bank Control Act (Act) (12 U.S.C. 1817(j)) and § 225.41 of the Board's Regulation Y (12 CFR 225.41) to acquire shares of a bank or bank holding company. The factors that are considered in acting on the applications are set forth in paragraph 7 of the Act (12 U.S.C. 1817(j)(7)).

The public portions of the applications listed below, as well as other related filings required by the Board, if any, are available for immediate inspection at the Federal Reserve Bank(s) indicated below and at the offices of the Board of Governors. This information may also be obtained on an expedited basis, upon request, by contacting the appropriate Federal Reserve Bank and from the Board's Freedom of Information Office at *https://www.federalreserve.gov/foia/request.htm.* Interested persons may express their views in writing on the standards enumerated in paragraph 7 of the Act.

Comments received are subject to public disclosure. In general, comments received will be made available without change and will not be modified to remove personal or business information including confidential, contact, or other identifying information. Comments should not include any information such as confidential information that would not be appropriate for public disclosure.

Comments regarding each of these applications must be received at the Reserve Bank indicated or the offices of the Board of Governors, Ann E. Misback, Secretary of the Board, 20th Street and Constitution Avenue NW, Washington, DC 20551–0001, not later than August 8, 2024.

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

1. *John Adam Robertson; Aaron Andrew Robertson; the Edwin G. Robertson Children's Irrevocable Trust, John Adam Robertson, Trustee; and the Craig E. Robertson Children's Irrevocable Trust, Aaron Andrew Robertson, Trustee, all of Speedwell, Tennessee; Emily Alayne King, Powell, Tennessee; Erica Leigh Corum, Harrogate, Tennessee; Matthew Craig Robertson, Tazewell, Tennessee; and Dakota John Robertson Bristol, Tennessee; as a group acting in concert,* to retain voting shares of Robertson Holding Company, L.P., Harrogate, Tennessee. Robertson Holding Company, L.P., controls Commercial Bancgroup, Inc., which controls Commercial Bank, both of Harrogate, Tennessee.

In addition, *Aaron Andrew Robertson; Cynthia Diane Robertson; James Oscar Robertson; John Adam Robertson, all of Speedwell, Tennessee; Sherri Jo Robertson and Noah Bradley Robertson, both of Harrogate, Tennessee; Dakota John Robertson, Bristol, Tennessee; Judith Yvonne Robertson, Cumberland Gap, Tennessee; Matthew Craig Robertson and Matthew Craig Robertson II, both of Tazewell, Tennessee; Olivia Grace Robertson, Hanahan, South Carolina; Emily Alayne King; Halle McLayne King; John McKinley King; and Riley Parker King, all of Powell, Tennessee; as a group acting in concert with Robertson Holding Company, L.P.,* to retain voting shares of Commercial Bancgroup, Inc.

B. Federal Reserve Bank of Richmond (Brent B. Hassell, Assistant Vice President) P.O. Box 27622, Richmond, Virginia 23261. Comments can also be sent electronically to *Comments.applications@rich.frb.org:*

1. *Barry J. Renbaum, Carol E. Renbaum, both individually and of Reisterstown, Maryland, Bryan M. Renbaum, individually, Frederick, Maryland; to form the Renbaum Family Control Group, a group acting in concert,* to acquire voting shares of Farmers and Merchants Bancshares, Inc., Hampstead, Maryland, and thereby indirectly acquire voting shares of Farmers and Merchants Bank, Upperco, Maryland.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2024–16265 Filed 7–23–24; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day–24–24HP; Docket No. CDC–2024–0056]

Proposed Data Collection Submitted for Public Comment and Recommendations

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice with comment period.

SUMMARY: The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden and maximize the utility of government information, invites the general public and other Federal agencies the opportunity to comment on a proposed information collection, as required by the Paperwork Reduction Act of 1995. This notice invites comment on a proposed information collection project titled Compliance Attestation Statement for the Framework for Nucleic Acid Synthesis Screening. The project aims to assist providers and manufacturers of synthetic nucleic acids and benchtop nucleic acid synthesis equipment (providers) in making an attestation that they have instituted a process to screen nucleic acid sequences of concern and verify customer legitimacy, in accordance with the requirements outlaid in the OSTP Framework for Nucleic Acid Synthesis Screening.

DATES: CDC must receive written comments on or before September 23, 2024.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2024–0056 by either of the following methods:

- *Federal eRulemaking Portal: www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600

Clifton Road NE, MS H21–8, Atlanta, Georgia 30329.

Instructions: All submissions received must include the agency name and Docket Number. CDC will post, without change, all relevant comments to *www.regulations.gov.*

Please note: Submit all comments through the Federal eRulemaking portal (*www.regulations.gov*) or by U.S. mail to the address listed above.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the information collection plan and instruments, contact Jeffrey M. Zirger, Information Collection Review Office, Centers for Disease Control and Prevention, 1600 Clifton Road NE, MS H21–8, Atlanta, Georgia 30329; Telephone: 404–639–7570; Email: *omb@cdc.gov.*

SUPPLEMENTARY INFORMATION: Under the Paperwork Reduction Act of 1995 (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. In addition, the PRA also requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each new proposed collection, each proposed extension of existing collection of information, and each reinstatement of previously approved information collection before submitting the collection to the OMB for approval. To comply with this requirement, we are publishing this notice of a proposed data collection as described below.

The OMB is particularly interested in comments that will help:

1. Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;

2. Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

3. Enhance the quality, utility, and clarity of the information to be collected;

4. Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submissions of responses; and

5. Assess information collection costs.

Proposed Project

Compliance Attestation Statement for the Framework for Nucleic Acid Synthesis Screening—New—Office of Science (OS), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

This data collection form was developed pursuant to the Framework for Nucleic Acid Synthesis Screening, which was released by the Office of Science and Technology Policy (OSTP) in April of 2024. This framework was directed by the *Executive Order on the Safe, Secure, and Trustworthy Development of Artificial Intelligence,* and recommends that providers and manufacturers of synthetic nucleic acids screen their sequences and customers before fulfilling orders to prevent potential misuse.

The Attestation Form will collect basic organizational information and an attestation of compliance from providers and manufacturers of synthetic nucleic acids and benchtop nucleic acid synthesis equipment. Data collected includes organization name, location, website, and type of organization. The form also includes primary and secondary contact information such as name, location, phone number and email address to ensure there is a point of contact with the company in case of questions regarding compliance and record keeping. This data is needed to ensure the self-attestation form can be filed and logged correctly, and to ensure the government can reach out to the correct contact if clarification if necessary.

CDC requests OMB approval for an estimated 20 annual burden hours. There is no cost to respondents other than their time to participate.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Providers and manufacturers of synthetic nucleic acids and bench top nucleic acid synthesis equipment.	Annual Provider and Manufacturer Self-Attestation Statement.	60	1	20/60	20
Total					20

Jeffrey M. Zirger,

Lead, Information Collection Review Office, Office of Public Health Ethics and Regulations, Office of Science, Centers for Disease Control and Prevention.

[FR Doc. 2024–16233 Filed 7–23–24; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10434 #66]

Medicaid and Children's Health Insurance Program (CHIP) Generic Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: On May 28, 2010, the Office of Management and Budget (OMB) issued Paperwork Reduction Act (PRA) guidance related to the ''generic'' clearance process. Generally, this is an expedited process by which agencies may obtain OMB's approval of collection of information requests that are ''usually voluntary, low-burden, and uncontroversial collections,'' do not raise any substantive or policy issues, and do not require policy or methodological review. The process requires the submission of an