FOR FURTHER INFORMATION CONTACT: Titel, Aiden Natural Resources Division (D230–0L), Office of Clean Air Programs, U.S. Environmental Protection Agency, Research Triangle Park, North Carolina, 27711; telephone number: (919) 541–4836; email address: *titel.aiden@epa.gov.*

SUPPLEMENTARY INFORMATION: This is a proposed extension of the ICR, which is currently approved through February 28, 2026. An agency may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number.

Public comments were previously requested via the **Federal Register** on August 6, 2024 during a 60-day comment period (89 FR 63933). This notice allows for an additional 30 days for public comments. Supporting documents, which explain in detail the information that the EPA will be collecting, are available in the public docket for this ICR. The docket can be viewed online at *www.regulations.gov* or in person at the EPA Docket Center, WJC West, Room 3334, 1301 Constitution Ave. NW, Washington, DC. The telephone number for the Docket Center is 202–566–1744. For additional information about EPA's public docket, visit *http://www.epa.gov/dockets.*

Abstract: The New Source Performance Standards (NSPS) for Generic Maximum Achievable Control Technology Standards for Acetal Resin; Acrylic and Modacrylic Fiber; Hydrogen Fluoride and Polycarbonate Production were proposed on October 14, 1998; and promulgated on June 29, 1999; and amended on: November 22, 1999; November 2, 2001; June 7, 2002; July 12, 2002; and October 8, 2014. These regulations apply to new and existing facilities of the following four categories: Polycarbonates (PC) Production, Acrylic and Modacrylic Fibers (AMF) Production, Acetal Resins (AR) Production, and Hydrogen Fluoride (HF) Production. New facilities include those that either commenced construction or reconstruction after the date of proposal. This information is being collected to assure compliance with 40 CFR part 63, subpart YY.

In general, all NESHAP standards require initial notifications, performance tests, and periodic reports by the owners/operators of the affected facilities. They are also required to maintain records of the occurrence and duration of any startup, shutdown, or malfunction in the operation of an affected facility, or any period during which the monitoring system is inoperative. These notifications, reports, and records are essential in determining compliance, and are required of all affected facilities subject to NESHAP.

Form Numbers: None.

Respondents/affected entities: Polycarbonate, acrylic and modacrylic fiber, acetal resin, and hydrogen fluoride production facilities.

Respondent's obligation to respond: Mandatory (40 CFR part 63, subpart YY).

Estimated number of respondents: 7 (total).

Frequency of response: Initially and semiannually.

Total estimated burden: 2,910 hours (per year). Burden is defined at 5 CFR 1320.03(b).

Total estimated cost: $457,000 (per year), includes $59,100 annualized capital or operation & maintenance costs.

Changes in the Estimates: There is no change in burden from the most recently approved ICR as currently identified in the OMB Inventory of Approved Burdens. This is due to two considerations. First, the regulations have not changed over the past three years and are not anticipated to change over the next three years. Second, the growth rate for this industry is very low or non-existent, so there is no significant change in the overall burden. There is an increase in the capital/startup or operation and maintenance (O&M) costs due to an adjustment to increase costs from 2013 to 2024 using the CEPCI CE Index.

Courtney Kerwin,

Deputy Director, Data & Enterprise Programs Division.

[FR Doc. 2026–03904 Filed 2–25–26; 8:45 am]

BILLING CODE 6560–50–P

FEDERAL COMMUNICATIONS COMMISSION

[OMB 3060–1246; FR ID 332620]

Information Collection Being Reviewed by the Federal Communications Commission Under Delegated Authority

AGENCY: Federal Communications Commission.

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork burdens, and as required by the Paperwork Reduction Act (PRA) of 1995, the Federal Communications Commission (FCC or the Commission) invites the general public and other Federal agencies to take this opportunity to comment on the following information collection. Comments are requested concerning: whether the proposed collection of information is necessary for the proper performance of the functions of the Commission, including whether the information shall have practical utility; the accuracy of the Commission's burden estimate; ways to enhance the quality, utility, and clarity of the information collected; ways to minimize the burden of the collection of information on the respondents, including the use of automated collection techniques or other forms of information technology; and ways to further reduce the information collection burden on small business concerns with fewer than 25 employees.

The FCC may not conduct or sponsor a collection of information unless it displays a currently valid control number. No person shall be subject to any penalty for failing to comply with a collection of information subject to the PRA that does not display a valid Office of Management and Budget (OMB) control number.

DATES: Written PRA comments should be submitted on or before April 27, 2026. If you anticipate that you will be submitting comments, but find it difficult to do so within the period of time allowed by this notice, you should advise the contact listed below as soon as possible.

ADDRESSES: Direct all PRA comments to Nicole Ongele, FCC, via email *PRA@fcc.gov* and to *nicole.ongele@fcc.gov.*

FOR FURTHER INFORMATION CONTACT: For additional information about the information collection, contact Nicole Ongele, (202) 418–2991.

SUPPLEMENTARY INFORMATION:

OMB Control Number: 3060–1246.

Title: FCC Reasonable Accommodation forms.

Form Number(s): FCC Form 5626, FCC Form 5627.

Type of Review: Extension of a currently approved collection.

Respondents: Individuals or households; Federal Government.

Number of Respondents and Responses: 164 respondents; 328 responses.

Estimated Time per Response: 0.16 hours–5 hours.

Frequency of Response: One-time reporting requirement.

Obligation to Respond: Voluntary. Statutory authority for these collections is contained in the Rehabilitation Act of 1973, 29 U.S.C. 12101 *et seq.;* see also 29 CFR part 1630; Establishing Procedures to Facilitate the Provision of Reasonable Accommodation; EEOC, Enforcement Guidance on Reasonable

Accommodation and Undue Hardship Under the Americans with Disabilities Act, 29 CFR part 1615.

Total Annual Burden: 850 hours.

Total Annual Cost: $3,400.

Needs and Uses: This information will be used by the FCC to process, track, and maintain the confidentiality of reasonable accommodation requests submitted on FCC Form 5626 and FCC Form 5627.

Federal Communications Commission.

Marlene Dortch,

Secretary, Office of the Secretary.

[FR Doc. 2026–03808 Filed 2–25–26; 8:45 am]

BILLING CODE 6712–01–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, Benjamin W. McDonough, Deputy Secretary of the Board, 20th Street and Constitution Avenue NW, Washington DC 20551–0001, not later than March 13, 2026.

A. Federal Reserve Bank of Minneapolis (Mark Nagle, Assistant Vice President) 90 Hennepin Avenue, Minneapolis, Minnesota 55480–0291. Comments can also be sent electronically to *MA@mpls.frb.org:*

1. *The Leon E. Langemeier GST Exempt Bridger Company Stock Family Trust for the benefit of Brian L. Langemeier, Bart Langemeier, as trustee, and the Leon E. Langemeier GST Exempt Bridger Company Stock Family Trust for the benefit of Brenda Langemeier, Bart Langemeier, as trustee, all of Red Lodge, Montana;* to join the Langemeier Family Control Group, a group acting in concert, to acquire voting shares of The Bridger Company, and thereby indirectly acquire voting shares of Bank of Bridger, National Association, both of Bridger, Montana. Bart Langemeier is a member of the Langemeier Family Control Group and was previously permitted by the Federal Reserve System to acquire voting shares of The Bridger Company in his individual capacity.

2. *Brian Schoenborn, Saint Joseph, Minnesota;* to acquire voting shares of Kensington Bancorp., Inc., and thereby indirectly acquire voting shares of Kensington Bank, both of Kensington, Minnesota.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Associate Secretary of the Board.

[FR Doc. 2026–03884 Filed 2–25–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2026–0199]

Meeting of the Advisory Committee on Immunization Practices

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

ACTION: Notice of meeting and request for comment.

SUMMARY: In accordance with the Federal Advisory Committee Act, the Centers for Disease Control and Prevention (CDC) announces the following meeting of the Advisory Committee on Immunization Practices (ACIP). Time will be available for public comment.

DATES: The meeting will be held on March 18, 2026, from 8 a.m. to 5 p.m., EST, and March 19, 2025, from 8 a.m. to 5 p.m., EST, (times subject to change; see the ACIP website for updates: *https://www.cdc.gov/vaccines/acip/index.html*). The meeting is expected to be held at the Centers for Disease Control and Prevention, with virtual option. Written comments must be received between March 2—12, 2026.

ADDRESSES: You may submit comments, identified by Docket No. CDC–2026–0199, by either of the methods listed below. CDC does not accept comments by email.

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments.
- *Mail:* ACIP Secretariat, ACIP Meeting, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H21–12, Atlanta, Georgia 30329–4027. Attn: Docket No. CDC–2026–0199.

Instructions: All submissions received must include the Agency name and docket number. For access to the docket to read background documents or comments received, go to *https://www.regulations.gov.*

The meeting will be webcast live via the World Wide Web. The webcast link can be found on the ACIP website at *https://www.cdc.gov/acip.*

FOR FURTHER INFORMATION CONTACT: ACIP Secretariat, Advisory Committee on Immunization Practices, Centers for Disease Control and Prevention, 1600 Clifton Road NE, Mailstop H21–12, Atlanta, Georgia 30329–4027. Email: *ACIP@cdc.gov.*

SUPPLEMENTARY INFORMATION: *Purpose:* The Advisory Committee on Immunization Practices is charged with advising the Director, Centers for Disease Control and Prevention (CDC), on the use of immunizing agents. In addition, under 42 U.S.C. 1396s, the Committee is mandated to establish and periodically review and, as appropriate, revise the list of vaccines for administration to vaccine-eligible children through the Vaccines for Children program, along with schedules regarding dosing interval, dosage, and contraindications to administration of vaccines. Further, under applicable provisions of the Affordable Care Act and section 2713 of the Public Health Service Act, immunization recommendations of ACIP that have been adopted by the Director, CDC, and appear on CDC immunization schedules generally must be covered by applicable health plans.

Matters to be Considered: The agenda will include updates on ACIP Workgroups and discussions on COVID–19 vaccine injuries and Long-COVID and ACIP recommendation