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

methodology. Recommendation votes may be scheduled for COVID–19 vaccine injuries and Long-COVID and ACIP recommendation methodology. Agenda items are subject to change as priorities dictate. For more information on the meeting agenda, visit *https://www.cdc.gov/acip/meetings/index.html.*

Meeting Information: The meeting will be webcast live via the World Wide Web. For more information on ACIP, please visit the ACIP website: *https://www.cdc.gov/acip.*

Public Participation

Interested persons or organizations are invited to participate by submitting written views, recommendations, and data. Please note that comments received, including attachments and other supporting materials, are part of the public record and are subject to public disclosure. Comments will be posted on *https://www.regulations.gov.* Therefore, do not include any information in your comment or supporting materials that you consider confidential or inappropriate for public disclosure. If you include your name, contact information, or other information that identifies you in the body of your comments, that information will be on public display. CDC will review all submissions and may choose to redact, or withhold, submissions containing private or proprietary information such as Social Security numbers, medical information, inappropriate language, or duplicate/near-duplicate examples of a mass-mail campaign. CDC will carefully consider all comments submitted into the docket.

Written Public Comment: The docket will be opened to receive written comments March 2–12, 2026. Written comments must be received by March 12, 2026.

Oral Public Comment: This meeting will include time for members of the public to make an oral comment. Oral public comment will occur before any scheduled votes, including all votes relevant to the ACIP's Affordable Care Act and Vaccines for Children Program roles. Priority will be given to individuals who submit a request to make an oral public comment before the meeting according to the procedures below.

Procedure for Oral Public Comment: All persons interested in making an oral public comment at the March 18—19, 2026, ACIP meeting must submit a request at *https://www.cdc.gov/acip/meetings/index.html* between March 2—12, 2026, and no later than 11:59 p.m., EST, March 12, 2026, according to the instructions provided.

If the number of persons requesting to speak is greater than can be reasonably accommodated during the scheduled time, CDC will conduct a random draw to determine the speakers for the scheduled public comment session. CDC staff will notify individuals regarding their request to speak by email by March 16, 2026. To accommodate the significant interest in participation in the oral public comment session of ACIP meetings, each speaker will be limited to three minutes, and each speaker may speak only once per meeting.

The Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Kalwant Smagh,

Director, Office of Strategic Business Initiatives, Office of the Chief Operating Officer, Centers for Disease Control and Prevention.

[FR Doc. 2026–03877 Filed 2–25–26; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10062 and CMS–10691]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare & Medicaid Services, Health and Human 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 (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 a second opportunity for public comment on the notice. Interested persons are invited to send comments regarding the burden estimate 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 on the collection(s) of information must be received by the OMB desk officer by March 30, 2026.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

To obtain copies of a supporting statement and any related forms for the proposed collection(s) summarized in this notice, please access the CMS PRA website by copying and pasting the following web address into your web browser: *https://www.cms.gov/Regulations-and-Guidance/Legislation/PaperworkReductionActof1995/PRA-Listing.*

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

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. 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 (44 U.S.C. 3506(c)(2)(A)) requires federal agencies to publish a 30-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 that summarizes the following proposed collection(s) of information for public comment.

1. *Type of Information Collection Request:* Extension without change of a currently approved collection; *Title of Information Collection:* Collection of Diagnostic Data in the Abbreviated