POLICIES AND PRACTICES FOR RETRIEVAL OF RECORDS:

Records are retrieved by the name or other programmatic identifier, including the date of the training or FMCS service.

POLICIES AND PRACTICES FOR RETENTION AND DISPOSAL OF RECORDS:

Temporarily stored data or records received by the project manager is deleted by the end of the fiscal year unless there is a specific need to retain it longer.

Records are retained and disposed of in accordance with General Records Schedule 4.2, issued by the National Archives and Records Administration.

ADMINSTRATIVE, TECHNICAL, AND PHYSICAL SAFEGUARDS:

FMCS maintains the FacilitatePro data and user profiles on its own servers and have an electronic backup system in place in the event of a system failure, as well as an alternative system consistent with requirements of Continuing of Operations Plan. The system requires a username and password which can only be created by FMCS. FMCS employee access to these systems is on a limited license basis and requires use of internal agency network and drives. Access is restricted, and accessible to limited FMCS Personnel such as the Project Manager, System Administrator, IT, and/or individuals in a need-to-know capacity. The other platforms mentioned above are web-based programs and require either FMCS Office 365 credentials, usernames and passwords, or both, in order to be used by an employee of FMCS.

RECORD ACCESS PROCEDURES:

FMCS employees, both current and former, may request access to their own records used as the basis for their performance evaluations through the Office of Human Resources. For external users, Privacy Act requests may be completed pursuant to 29 CFR 1410.3, Individual access requests. Individuals must provide the following information for their records to be located and identified: (1) Full name, (2) Address, and (3) A specific description of the record content requested. Also, see *https://www.fmcs.gov/privacy-policy/*.

CONTESTING RECORDS PROCEDURES:

See 29 CFR 1410.6, Requests for correction or amendment of records, on how to contest the content of any records. Privacy Act requests to amend or correct records may be submitted to the Privacy Office at *privacy@fmcs.gov* or via mail at Federal Mediation and Conciliation Service, 250 E Street SW, Washington, DC 20427. Also, see *https://www.fmcs.gov/privacy-policy/*.

NOTIFICATION PROCEDURES:

See 29 CFR 1410.3(a), Individual access requests.

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

None.

[FR Doc. 2022–11617 Filed 5–27–22; 8:45 am]

BILLING CODE 6732–01–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 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 June 13, 2022.

A. Federal Reserve Bank of Cleveland (Bryan S. Huddleston, Vice President) 1455 East Sixth Street, Cleveland, Ohio 44101–2566. Comments can also be sent electronically to *Comments.applications@clev.frb.org:*

1. *Jodi Hillyer and Kim Hillyer, both of Dennison, Ohio; Kurt Shelley, New Philadelphia, Ohio; Kim Shelley, Belmont, Maine; Tina Floyd, North Canton, Ohio; Todd Scott, Strasburg, Ohio; the Connolly, Hillyer and Ong Law Firm, Uhrichsville, Ohio*; to join the Hillyer Family Control Group, a group acting in concert, to retain voting shares of FNB, Inc., Dennison, Ohio.

Board of Governors of the Federal Reserve System.

Michele Taylor Fennell,

Deputy Associate Secretary of the Board.

[FR Doc. 2022–11545 Filed 5–27–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0905]

Vaccines and Related Biological Products Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, Health and Human Services (HHS).

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA) announces a forthcoming public advisory committee meeting of the Vaccines and Related Biological Products Advisory Committee (VRBPAC). The general function of the committee is to provide advice and recommendations to FDA on regulatory issues. The committee will meet in open session to discuss whether and how the SARS–CoV–2 strain composition of COVID–19 vaccines should be modified. The meeting will be open to the public. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held virtually on June 28, 2022, from 8:30 a.m. to 5 p.m. Eastern Time.

ADDRESSES: Please note that due to the impact of this COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. The online web conference meeting will be available at *https://youtu.be/BFdzNUus_CE* on the day of the meeting. Answers to commonly asked questions may be accessed at: *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm408555.htm.*

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2022–N–0905. The docket will close on June 27, 2022. Submit either electronic or written comments on this public meeting by June 27, 2022. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before June 27, 2022. The *https://www.regulations.gov* electronic filing system will accept

comments until 11:59 p.m. Eastern Time at the end of June 27, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Comments received on or before June 22, 2022, will be provided to the committee. Comments received after June 22, 2022, and by June 27, 2022, will be taken into consideration by FDA. In the event that the meeting is cancelled, FDA will continue to evaluate any relevant applications or information, and consider any comments submitted to the docket, as appropriate.

You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov*.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2022–N–0905 for ''Vaccines and Related Biological Products Advisory Committee (VRBPAC); Notice of Meeting; Establishment of a Public Docket; Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m. Eastern Time, Monday through Friday, 240–402–7500.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' FDA will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify the information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

FOR FURTHER INFORMATION CONTACT: Prabhakara Atreya or Sussan Paydar, Center for Biologics Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 71, Rm. 1226, Silver Spring, MD 20993–0002, 240–506–4946, *CBERVRBPAC@fda.hhs.gov;* or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area). A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice. Therefore, you should always check the FDA's website at *https://www.fda.gov/AdvisoryCommittees/default.htm* and scroll down to the appropriate advisory committee meeting link, or call the advisory committee information line to learn about possible modifications before coming to the meeting.

SUPPLEMENTARY INFORMATION:

Agenda: The meeting presentations will be heard, viewed, captioned, and recorded through an online teleconferencing platform. On June 28, 2022, the committee will meet in open session to discuss whether and how the SARS–CoV–2 strain composition of COVID–19 vaccines should be modified.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its website prior to the meeting, the background material will be made publicly available at the time of the advisory committee meeting, and the background material will be posted on FDA's website after the meeting. Background material is available at *https://www.fda.gov/AdvisoryCommittees/Calendar/default.htm.* Scroll down to the appropriate advisory committee meeting link. The meeting will include slide presentations with audio components to allow the presentation of materials in a manner that most closely resembles an in-person advisory committee meeting.

Procedure: On June 28, 2022, from 8:30 a.m. to 5 p.m. Eastern Time, the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. All electronic and written submissions submitted to the Docket (see **ADDRESSES**) on or before June 22, 2022, will be provided to the committee. Comments received after June 22, 2022, and by June 27, 2022, will be taken into consideration by FDA. Oral presentations from the public will be scheduled between approximately 1:30 p.m. and 2:30 p.m. Eastern Time. Those individuals interested in making formal oral presentations should notify the contact person and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and email addresses

of proposed participants, and an indication of the approximate time requested to make their presentation on or before 6 p.m. Eastern Time on June 21, 2022. Time allotted for each presentation may be limited. If the number of registrants requesting to speak is greater than can be reasonably accommodated during the scheduled open public hearing session, FDA may conduct a lottery to determine the speakers for the scheduled open public hearing session. The contact person will notify interested persons regarding their request to speak by June 23, 2022.

For press inquiries, please contact the Office of Media Affairs at *fdaoma@fda.hhs.gov* or 301–796–4540.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with disabilities. If you require accommodations due to a disability, please contact Prabhakara Atreya or Sussan Paydar (see **FOR FURTHER INFORMATION CONTACT**) at least 7 days in advance of the meeting.

FDA is committed to the orderly conduct of its advisory committee meetings. Please visit our website at *https://www.fda.gov/AdvisoryCommittees/AboutAdvisoryCommittees/ucm111462.htm* for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: May 25, 2022.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2022–11670 Filed 5–26–22; 11:15 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0904]

Vaccines and Related Biological Products Advisory Committee; Notice of Meeting; Establishment of a Public Docket; Request for Comments

AGENCY: Food and Drug Administration, Health and Human Services (HHS).

ACTION: Notice; establishment of a public docket; request for comments.

SUMMARY: The Food and Drug Administration (FDA or Agency) announces a forthcoming public advisory committee meeting of the Vaccines and Related Biological Products Advisory Committee. The general function of the committee is to provide advice and recommendations to the Agency on FDA's regulatory issues. Members will participate via teleconference. This 2-day virtual meeting will be held to discuss recent requests to amend the Emergency Use Authorization (EUA) of the Moderna COVID–19 mRNA vaccine to include the administration of a primary series to infants, children, and adolescents 6 months through 17 years of age and to amend the EUA of the Pfizer-BioNTech COVID–19 mRNA vaccine to include the administration of a primary series to infants and children 6 months through 4 years of age. The meeting will be open to the public on both days. FDA is establishing a docket for public comment on this document.

DATES: The meeting will be held June 14 and June 15, 2022, from 8:30 a.m. to 5 p.m. Eastern Time (ET). Comments received on or before June 7, 2022, will be provided to the committee. Comments received after June 7, 2022, and by June 13, 2022, be taken into consideration by FDA.

ADDRESSES: Please note that due to the impact of this COVID–19 pandemic, all meeting participants will be joining this advisory committee meeting via an online teleconferencing platform. The online web conference meeting will be available at the following separate links on the respective days of the meeting:

Day 1: *https://youtu.be/GbNpaZeDPiA*
Day 2: *https://youtu.be/Ixm4UmldTGQ*

FDA is establishing a docket for public comment on this meeting. The docket number is FDA–2022–N–0904. The docket will close on June 13, 2022. Submit either electronic or written comments on this public meeting by June 13, 2022. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before June 13, 2022. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. ET at the end of June 13, 2022. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are received on or before that date.

Comments received on or before June 7, 2022, will be provided to the committee. Comments received after June 7, 2022, and by June 13, 2022, will be taken into consideration by FDA. If the meeting is canceled, FDA will continue to evaluate any relevant applications, submissions, or information and consider any comments submitted to the docket, as appropriate.

You may submit comments as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov*. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov*.
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2022–N–0904 for ''Vaccines and Related Biological Products; Notice of Meeting; Establishment of a Public Docket; Request for Comments.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- *Confidential Submissions*—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two