DATES: Authority for the Arthritis Advisory Committee will expire on April 5, 2026, unless the Commissioner formally determines that renewal is in the public interest.

FOR FURTHER INFORMATION CONTACT: Jessica Seo, Center for Drug Evaluation Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–9001, *AAC@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Pursuant to 41 CFR 102–3.65 and approval by the Department of Health and Human Services and by the General Services Administration, FDA is announcing the renewal of the Arthritis Advisory Committee (the Committee). The Committee is a discretionary Federal advisory committee established to provide advice to the Commissioner. The Committee advises the Commissioner or designee in discharging responsibilities as they relate to helping to ensure safe and effective drugs for human use and, as required, any other product for which FDA has regulatory responsibility.

The Committee reviews and evaluates data concerning the safety and effectiveness of marketed and investigational human drug products for use in the treatment of arthritis, rheumatism, and related diseases, and makes appropriate recommendations to the Commissioner.

Pursuant to its Charter, the Committee shall consist of a core of 11 voting members including the Chair. Members and the Chair are selected by the Commissioner or designee from among authorities knowledgeable in the fields of arthritis, rheumatology, orthopedics, epidemiology or statistics, analgesics, and related specialties. Members will be invited to serve for overlapping terms of up to 4 years. Non-Federal members of this committee will serve as Special Government Employees, representatives, or Ex-Officio members. Federal members will serve as Regular Government Employees or Ex-Officios. The core of voting members may include one technically qualified member, selected by the Commissioner or designee, who is identified with consumer interests and is recommended by either a consortium of consumer-oriented organizations or other interested persons. In addition to the voting members, the Committee may include one non-voting representative member who is identified with industry interests. There may also be an alternate industry representative.

The Commissioner or designee shall have the authority to select members of other scientific and technical FDA advisory committees (normally not to exceed 10 members) to serve temporarily as voting members and to designate consultants to serve temporarily as voting members when: (1) expertise is required that is not available among current voting standing members of the Committee (when additional voting members are added to the Committee to provide needed expertise, a quorum will be based on the combined total of regular and added members) or (2) to comprise a quorum when, because of unforeseen circumstances, a quorum is or will be lacking. Because of the size of the Committee and the variety in the types of issues that it will consider, FDA may, in connection with a particular committee meeting, specify a quorum that is less than a majority of the current voting members. The Agency's regulations (21 CFR 14.22(d)) authorize a committee charter to specify quorum requirements.

If functioning as a medical device panel, an additional non-voting representative member of consumer interests and an additional non-voting representative member of industry interests will be included in addition to the voting members.

Further information regarding the most recent charter and other information can be found at *https://www.fda.gov/advisory-committees/human-drug-advisory-committees/arthritis-advisory-committee* or by contacting the Designated Federal Officer (see **FOR FURTHER INFORMATION CONTACT**). In light of the fact that no change has been made to the committee name or description of duties, no amendment will be made to 21 CFR 14.100.

This notice is issued under the Federal Advisory Committee Act as amended (5 U.S.C. 1001 et seq). For general information related to FDA advisory committees, please visit us at *http://www.fda.gov/AdvisoryCommittees/default.htm.*

Dated: June 4, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–12525 Filed 6–6–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2022–N–0335]

Advisory Committee; Obstetrics, Reproductive and Urologic Drugs Advisory; Renewal

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; renewal of Federal advisory committee.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is announcing the renewal of the Obstetrics, Reproductive and Urologic Drugs Advisory Committee by the Commissioner of Food and Drugs (the Commissioner). The Commissioner has determined that it is in the public interest to renew the Obstetrics, Reproductive and Urologic Drugs Advisory Committee for an additional 2 years beyond the charter expiration date. The new charter will be in effect until the March 23, 2026, expiration date.

DATES: Authority for the Obstetrics, Reproductive and Urologic Drugs Advisory Committee will expire on March 23, 2026, unless the Commissioner formally determines that renewal is in the public interest.

FOR FURTHER INFORMATION CONTACT: Joyce Frimpong, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301–796–7973, email: *ORUDAC@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Pursuant to 41 CFR 102–3.65 and approval by the Department of Health and Human Services and by the General Services Administration, FDA is announcing the renewal of the Obstetrics, Reproductive and Urologic Drugs Advisory Committee (the Committee). The Committee is a discretionary Federal advisory committee established to provide advice to the Commissioner. The Committee advises the Commissioner or designee in discharging responsibilities as they relate to helping to ensure safe and effective drugs for human use and, as required, any other product for which FDA has regulatory responsibility.

The Committee reviews and evaluates data on the safety and effectiveness of marketed and investigational human drug products for use in the practice of obstetrics, gynecology, urology, and related specialties, and makes

appropriate recommendations to the Commissioner.

Pursuant to its Charter, the Committee shall consist of a core of 11 voting members including the Chair. Members and the Chair are selected by the Commissioner or designee from among authorities knowledgeable in the fields of obstetrics, gynecology, urology, pediatrics, epidemiology, or statistics and related specialties. Members will be invited to serve for overlapping terms of up to 4 years. Non-Federal members of this committee will serve as Special Government Employees, representatives, or Ex-Officio members. Federal members will serve as Regular Government Employees or Ex-Officios. The core of voting members may include one technically qualified member, selected by the Commissioner or designee, who is identified with consumer interests and is recommended by either a consortium of consumer-oriented organizations or other interested persons. In addition to the voting members, the Committee may include one non-voting representative member who is identified with industry interests. There may also be an alternate industry representative.

The Commissioner or designee shall have the authority to select members of other scientific and technical FDA advisory committees (normally not to exceed 10 members) to serve temporarily as voting members and to designate consultants to serve temporarily as voting members when: (1) expertise is required that is not available among current voting standing members of the Committee (when additional voting members are added to the Committee to provide needed expertise, a quorum will be based on the combined total of regular and added members), or (2) to comprise a quorum when, because of unforeseen circumstances, a quorum is or will be lacking. Because of the size of the Committee and the variety in the types of issues that it will consider, FDA may, in connection with a particular committee meeting, specify a quorum that is less than a majority of the current voting members. The Agency's regulations (21 CFR 14.22(d)) authorize a committee charter to specify quorum requirements.

If functioning as a medical device panel, an additional non-voting representative member of consumer interests and an additional non-voting representative member of industry interests will be included in addition to the voting members.

Further information regarding the most recent charter and other information can be found at *https://www.fda.gov/advisory-committees/human-drug-advisory-committees/obstetrics-reproductive-and-urologic-drugs-advisory-committee-formerly-bone-reproductive-and* or by contacting the Designated Federal Officer (see **FOR FURTHER INFORMATION CONTACT**). In light of the fact that no change has been made to the committee name or description of duties, no amendment will be made to 21 CFR 14.100.

This notice is issued under the Federal Advisory Committee Act as amended (5 U.S.C. 1001 *et seq.*). For general information related to FDA advisory committees, please visit us at *https://www.fda.gov/Advisory Committees/default.htm.*

Dated: June 4, 2024.

Lauren K. Roth,

Associate Commissioner for Policy.

[FR Doc. 2024–12528 Filed 6–6–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Privacy Act of 1974; System of Records

AGENCY: National Institutes of Health (NIH), Department of Health and Human Services (HHS).

ACTION: Notice of a new system of records.

SUMMARY: In accordance with the requirements of the Privacy Act of 1974, as amended (Privacy Act, or Act), the Department of Health and Human Services (HHS) is establishing a new System of Records (SOR), 09–25–0224, ''NIH Police Records,'' to be maintained by the National Institutes of Health (NIH). The new system of records will contain records about individuals who are the subject of investigations of crime, civil disturbances, and traffic accidents occurring on or otherwise affecting the protection of life and property on NIH property. Because the records will constitute law enforcement investigatory material, elsewhere in the **Federal Register** the agency has published a notice of proposed rulemaking (NPRM) to exempt this system of records from certain requirements of the Privacy Act based on subsections (j)(2) and (k)(2) of the Act. The system of records is more fully described in the system of records notice (SORN) published in this notice.

DATES: The comment period for this SORN is co-extensive with the 60-day comment period provided in the NPRM; *i.e.,* written comments on the SORN should be submitted by August 6, 2024. The new system of records, including the routine uses and the exemptions, will become effective when NIH publishes a Final Rule, which will not occur until the 60-day comment period provided in the NPRM has expired and any comments received on the NPRM (or on this SORN) have been addressed.

ADDRESSES: The public should address written comments, identified by the Privacy Act System of Records (PA SOR) Number 09–25–0224, by any of the following methods:

- *Federal eRulemaking Portal: https://regulations.gov.* Follow the instructions for submitting comments.
- *Email: privacy@mail.nih.gov* and include PA SOR number 09–25–0224 in the subject line of the message.
- *Phone:* (301) 402–6469 (not a toll-free number).
- *Fax:* (301) 402–0169.
- *Mail:* NIH Privacy Act Officer, Office of Management Assessment, National Institutes of Health, 6705 Rockledge Drive (RK1) 601, Rockville, MD 20892–7901.
- *Hand Delivery/Courier:* 6705 Rockledge Drive (RK1) 601, Rockville, MD 20892–7901.

Comments received will be available for inspection and copying at this same address from 9:00 a.m. to 3:00 p.m., Monday through Friday, Federal holidays excepted.

FOR FURTHER INFORMATION CONTACT: General questions about the system of records may be submitted to Dustin Close, NIH Privacy Act Officer, by email at *privacy@mail.nih.gov* or mail at the Office of Management Assessment (OMA), Office of the Director (OD), National Institutes of Health (NIH), 6705 Rockledge Drive (RK1) 601, Rockville, MD 20892–7901. Telephone: 301–402–6469.

SUPPLEMENTARY INFORMATION: The Privacy Act (5 U.S.C. 552a) governs the means by which the United States Government collects, maintains, and uses records in a system of records. A ''system of records'' is a group of any records under the control of a federal agency from which information about individuals is retrieved by name or other personal identifier. The Privacy Act requires each agency to publish in the **Federal Register** a SORN identifying and describing each system of records the agency maintains, including the purposes for which the agency uses records in the system of records, the routine uses for which the agency discloses, or may disclose, such information outside the agency without the subject individual's prior written consent, and procedures explaining how subject individuals can exercise their rights under the Privacy Act (*e.g.,* to