#### Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2018–08383 Filed 4–20–18; 8:45 am] BILLING CODE 4163–18–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ ATSDR)

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

**ACTION:** Notice of meeting.

SUMMARY: In accordance with the Federal Advisory Committee Act, the CDC announces the following meeting for the Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ ATSDR). This meeting is open to the public, limited only by available seating. The meeting room accommodates approximately 60 people. The public is also welcome to listen to the meeting by calling 888-989-4501, passcode 9885805, limited by 100 lines. The deadline for notification of attendance is May 14, 2018. The public comment period is scheduled on June 5, 2018 from 2:30 p.m. until 2:45 p.m., EDT and June 6, 2018 from 10:10 a.m. until 10:25 a.m., EDT. Individuals wishing to make a comment during Public Comment period, please email your name, organization, and phone number by May 7, 2018 to Amanda Malasky at amalasky@cdc.gov.

**DATES:** The meeting will be held on June 5, 2018, 8:30 a.m. to 4:00 p.m., EDT and June 6, 2018, 8:30 a.m. to 11:30 a.m., EDT.

ADDRESSES: CDC, 4770 Buford Highway, CDC Building 106, Room 1B, Atlanta, GA 30341.

### FOR FURTHER INFORMATION CONTACT:

Shirley Little, Program Analyst, NCEH/ATSDR, CDC, 4770 Buford Hwy., Mail Stop F–45, Atlanta, GA 30341, telephone (770) 488–0577; email snl7@cdc.gov.

#### SUPPLEMENTARY INFORMATION:

Purpose: The Secretary, Department of Health and Human Services (HHS) and by delegation, the Director, CDC

and Administrator, NCEH/ATSDR, are authorized under Section 301 (42 U.S.C. 241) and Section 311 (42 U.S.C. 243) of the Public Health Service Act, as amended, to: (1) Conduct, encourage, cooperate with, and assist other appropriate public authorities, scientific institutions, and scientists in the conduct of research, investigations, experiments, demonstrations, and studies relating to the causes, diagnosis, treatment, control, and prevention of physical and mental diseases and other impairments; (2) assist states and their political subdivisions in the prevention of infectious diseases and other preventable conditions and in the promotion of health and wellbeing; and (3) train state and local personnel in health work. The BSC, NCEH/ATSDR provides advice and guidance to the Secretary, HHS; the Director, CDC and Administrator, ATSDR; and the Director, NCEH/ATSDR, regarding program goals, objectives, strategies, and priorities in fulfillment of the agency's mission to protect and promote people's health. The board provides advice and guidance that will assist NCEH/ATSDR in ensuring scientific quality, timeliness, utility, and dissemination of results. The board also provides guidance to help NCEH/ATSDR work more efficiently and effectively with its various constituents and to fulfill its mission in protecting America's health.

Matters to be Considered: The agenda will include discussions on Recovery Efforts to Address Environmental Health Impacts after the 2017 Hurricanes, NCEH/ATSDR Program Responses to BSC Guidance and Action Items, PFAS multi-site study, PEASE, Biomonitoring for PFAS in children, Use of Citizen Science for Assessment of Health Risks, Statistical Inferences in Environmental Epidemiology, and NCEH/ASTDR work with tribes/Tribal Programs. Agenda items are subject to change as priorities dictate.

The Director, Management Analysis and Services Office, 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.

#### Elizabeth Millington,

Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

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### DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration [Docket No. FDA-2018-N-1242]

# Advisory Committee; Arthritis Advisory Committee, Renewal

**AGENCY:** Food and Drug Administration, HHS

**ACTION:** Notice; renewal of advisory committee.

SUMMARY: The Food and Drug Administration (FDA) is announcing the renewal of the Arthritis 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 Arthritis Advisory Committee for an additional 2 years beyond the charter expiration date. The new charter will be in effect until April 5, 2020.

**DATES:** Authority for the Arthritis Advisory Committee will expire on April 5, 2020, unless the Commissioner formally determines that renewal is in the public interest.

#### FOR FURTHER INFORMATION CONTACT:

Yinghua Wang, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993–0002, 301– 796–9001, email: AAC@fda.hhs.gov.

**SUPPLEMENTARY INFORMATION:** Pursuant to 41 CFR 102–3.65 and approval by the Department of Health and Human Services pursuant to 45 CFR part 11 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.

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