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[FR Doc. 2018–16090 Filed 7–26–18; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

#### Advisory Committee to the Director (ACD), Centers for Disease Control and Prevention (CDC)—Health Disparities Subcommittee (HDS)

**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 Advisory Committee to the Director, Centers for Disease Control and Prevention—Health Disparities Subcommittee (ACD, CDC–HDS). This meeting is open to the public, limited only by the 50 audio phone lines. The public is also welcome to listen to the meeting by teleconference. Please dial (866) 918–8397 and enter code 9346283. There are 50 lines available. The public comment period is from 3:15 p.m.–3:20 p.m.

**DATES:** The meeting will be held on October 9, 2018, 1:30 p.m. to 3:30 p.m., EDT.

**ADDRESSES:** Teleconference phone (866) 918–8397 and enter code 9346283.

**FOR FURTHER INFORMATION CONTACT:** Leandris Liburd, Ph.D., M.P.H., M.A., Designated Federal Officer, Health Disparities Subcommittee, Advisory Committee to the Director, CDC, 1600 Clifton Road NE, M/S K–77, Atlanta, Georgia 30329. Telephone (404) 498–6482; Email: [ACDDirector@cdc.gov](mailto:ACDDirector@cdc.gov).

#### **SUPPLEMENTARY INFORMATION:**

*Purpose:* The Subcommittee will provide counsel to the CDC Director through the ACD on strategic and other health disparities and health equity issues and provide guidance on opportunities for CDC.

*Matters to be Considered:* The agenda will include discussions on new member orientation. This meeting will provide information to new members regarding their role & duties on this subcommittee. 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.

**Dia Taylor,**

*Acting Chief Operating Officer, Centers for Disease Control and Prevention.*

[FR Doc. 2018–16103 Filed 7–26–18; 8:45 am]

**BILLING CODE 4163–19–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. FDA–2018–P–1283]

#### Determination That Metaxalone Tablets, 640 Milligrams, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA or Agency) has determined that metaxalone tablets, 640 milligrams (mg), were not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDAs) for metaxalone tablets, 640 mg, if all other legal and regulatory requirements are met.

**FOR FURTHER INFORMATION CONTACT:** Glen Cheng, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave. Bldg. 51, Rm. 6217, Silver Spring, MD 20993–0002, 301–796–1494.

**SUPPLEMENTARY INFORMATION:** In 1984, Congress enacted the Drug Price Competition and Patent Term Restoration Act of 1984 (Pub. L. 98–417) (the 1984 amendments), which authorized the approval of duplicate versions of drug products under an ANDA procedure. ANDA applicants must, with certain exceptions, show that the drug for which they are seeking approval contains the same active ingredient in the same strength and dosage form as the “listed drug,” which is a version of the drug that was previously approved. ANDA applicants do not have to repeat the extensive clinical testing otherwise necessary to gain approval of a new drug application (NDA).

The 1984 amendments include what is now section 505(j)(7) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C.

355(j)(7)), which requires FDA to publish a list of all approved drugs. FDA publishes this list as part of the “Approved Drug Products With Therapeutic Equivalence Evaluations,” which is known generally as the “Orange Book.” Under FDA regulations, drugs are removed from the list if the Agency withdraws or suspends approval of the drug’s NDA or ANDA for reasons of safety or effectiveness or if FDA determines that the listed drug was withdrawn from sale for reasons of safety or effectiveness (21 CFR 314.162).

A person may petition the Agency to determine, or the Agency may determine on its own initiative, whether a listed drug was withdrawn from sale for reasons of safety or effectiveness. This determination may be made at any time after the drug has been withdrawn from sale, but must be made prior to approving an ANDA that refers to the listed drug (§ 314.161 (21 CFR 314.161)). FDA may not approve an ANDA that does not refer to a listed drug.

Metaxalone tablets, 640 mg, are the subject of NDA 22–503, held by Primus Pharmaceuticals, Inc., and initially approved on June 1, 2015. Metaxalone tablets, 640 mg, are indicated as an adjunct to rest, physical therapy, and other measures for the relief of discomfort associated with acute, painful musculoskeletal conditions.

In a letter dated September 30, 2015, the previous NDA holder CorePharma, LLC notified FDA that metaxalone tablets, 640 mg, were discontinued, and FDA moved the drug product to the “Discontinued Drug Product List” section of the Orange Book.

Sovereign Pharmaceuticals, LLC submitted a citizen petition dated March 26, 2018 (Docket No. FDA–2018–P–1283), under 21 CFR 10.30, requesting that the Agency determine whether metaxalone tablets, 640 mg, were withdrawn from sale for reasons of safety or effectiveness.

After considering the citizen petition and reviewing Agency records and based on the information we have at this time, FDA has determined under § 314.161 that metaxalone tablets, 640 mg, were not withdrawn for reasons of safety or effectiveness. The petitioner has identified no data or other information suggesting that metaxalone tablets, 640 mg, were withdrawn for reasons of safety or effectiveness. We have carefully reviewed our files for records concerning the withdrawal of metaxalone tablets, 640 mg, from sale. We have also independently evaluated relevant literature and data for possible postmarketing adverse events. We have reviewed the available evidence and determined that this drug product was