maintain the assets that are being divested (as well as the ''crown jewel'' assets) in their current condition and provide gas gathering services on the same terms and conditions available to customers on March 1, 2000, until the assets are sold.

The purpose of this analysis is to invite public comment concerning the consent order. This analysis is not intended to constitute an official interpretation of the agreement and order or to modify their terms in any way.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 00–8771 Filed 4–7–00; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Special Emphasis Panel meeting.

A Special Emphasis panel (SEP) is a committee of a few experts selected to conduct scientific reviews of applications related to their areas of expertise. The committee members are drawn from a list of experts and designated to serve for particular individual meetings rather than for extended fixed terms of services.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C., 552b (c)(6). Grant applications are to be reviewed and discussed at this meeting. These discussions are likely to include personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure under the above-cited statutes.

Name of SEP: Understanding the Eliminating Minority Health Disparities.

Date: May 1–2, 2000 (Open from 8 a.m. to 8:15 a.m. and closed for remainder of the meeting).

Place: Doubletree Hotel, 1750 Rockville Pike, Conference TBD, Rockville, Maryland 20852.

Contact Person: Anyone wishing to obtain a roster of members or minutes of the meeting should contact Ms. Jenny Griffith, Committee management Officer, Office of Research Review, Education and Policy, AHRQ, 2101 East Jefferson Street, Suite 400, Rockville, Maryland 20852, Telephone (301) 594–1847.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: March 29, 2000.

John M. Eisenberg,

Director.

[FR Doc. 00–8842 Filed 4–7–00; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99P–4848]

Determination That Carbinoxamine Maleate 4 Milligrams per 5 Cubic Centimeters Elixir Was 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) has determined that carbinoxamine maleate (Clistin) 4 milligrams (mg) per 5 cubic centimeters (cc) elixir was not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDA's) for carbinoxamine maleate 4 mg per 5 cc elixir.

FOR FURTHER INFORMATION CONTACT: Mary E. Catchings, Center for Drug Evaluation and Research (HFD–7), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20855, 301–594–2041.

SUPPLEMENTARY INFORMATION: In 1984, Congress enacted the Drug Price Competition and Patent Term Restoration Act of 1984 (Public Law 98–417) (the 1984 amendments), which authorized the approval of duplicate versions of drug products approved under an ANDA procedure. ANDA sponsors 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. Sponsors of ANDA's do not have to repeat the extensive clinical testing otherwise necessary to gain approval of a new drug application (NDA). The only clinical data required in an ANDA are data to show that the drug that is the subject of the ANDA is bioequivalent to the listed drug.

The 1984 amendments included what is now section 505(j)(6) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 355(j)(6)), 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,'' generally known as the ''Orange Book.'' Under FDA regulations, drugs are withdrawn 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). Under § 314.161(a)(1) (21 CFR 314.161(a)(1)) the agency must make a determination as to whether a listed drug was withdrawn from sale for reasons of safety or effectiveness before an ANDA that refers to that listed drug may be approved. FDA may not approve an ANDA that does not refer to a listed drug.

In a citizen petition dated October 8, 1999 (Docket No. 99P–4848/CP1), submitted under 21 CFR 314.122, Mikart, Inc., requested that the agency determine whether carbinoxamine maleate (Clistin) 4 mg per 5 cc elixir was withdrawn from sale for reasons of safety or effectiveness. Carbinoxamine maleate (Clistin) 4 mg per 5 cc elixir was the subject of approved NDA 8–955. In the **Federal Register** of April 5, 1985 (50 FR 13661), FDA withdrew approval of NDA 8–955 for Clistin Elixir after McNeil Pharmaceutical notified the agency that Clistin Elixir was no longer being marketed under NDA 8–955 and requested the withdrawal of that application.

FDA has reviewed its records and, under § 314.161, has determined that carbinoxamine maleate 4 mg per 5 cc elixir was not withdrawn from sale for reasons of safety or effectiveness. Accordingly, the agency will list carbinoxamine maleate 4 mg per 5 cc elixir in the ''Discontinued Drug Product List'' section of the Orange Book. The ''Discontinued Drug Product List'' identifies, among other items, drug products that have been discontinued from marketing for reasons other than safety or effectiveness. ANDA's that refer to carbinoxamine maleate 4 mg per 5 cc elixir as the listed drug may be approved by the agency.

Dated: April 3, 2000.

Margaret M. Dotzel,

Acting Associate Commissioner for Policy.

[FR Doc. 00–8715 Filed 4–7–00; 8:45 am]

BILLING CODE 4160–01–F