present to the committee commercial information regarding various medical devices used in obstetrics and gynecology that are currently being evaluated by FDA.

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

Dated: June 12, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–16125 Filed 6–18–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Nonprescription Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on July 14, 1997, 8:30 a.m. to 5 p.m.

Location: Holiday Inn, Versailles Ballrooms I and II, 8120 Wisconsin Ave., Bethesda, MD.

Contact Person: Andrea G. Neal or Angie Whitacre, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–5455, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12541. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will hear presentations and discuss the proposed labeling requirements for over-the-counter (OTC) drug products that will enable consumers to better read and understand OTC drug product labeling and to apply this information to the safe and effective use of OTC drug products. Elsewhere in this issue of the **Federal Register**, FDA is also extending the comment period on a proposed rule regarding labeling requirements for OTC drug products that appeared in the **Federal Register** of February 27, 1997 (62 FR 9024).

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by July 3, 1997. Oral presentations from the public will be scheduled between approximately 1:15 p.m. to 2:15 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before July 3, 1997, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.I11Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: June 12, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–16067 Filed 6–18–97; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Public Law 104–13), the Health Resources and Services Administration (HRSA) will publish periodic summaries of proposed projects being developed for submission to OMB under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans, call the HRSA Reports Clearance Officer on (301) 443–1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology.

Proposed Project: Grantee Reporting Requirements for the Rural Telemedicine Grant Program

New—The Rural Telemedicine Grant Program is authorized by Section 330A of the Public Health Service Act as amended by the Health Centers Consolidation Act of 1996 (Public Law 104–229). The goal of the program is to improve access to quality health services for rural residents and reduce the isolation of rural practitioners through the use of telemedicine technologies. The two objectives of the Rural Telemedicine Grant Program are: 1) to demonstrate how telemedicine can be used as a tool in developing integrated systems of health care, which would improve access to health services for rural individuals across the lifespan and reduce the isolation of rural health care practitioners; and 2) to evaluate the feasibility, costs, appropriateness and acceptability of rural telemedicine services and technologies. Such evaluation is needed to determine how best to organize and provide telemedicine services in a sustainable manner.

Grantees will be responsible for submitting the data collection instruments listed in the burden table below. Grantees will gather information from sources involved with their telemedicine program, including patients, providers, health administrators and site coordinators. Information gathered on the data collection instruments will be entered into a database which will communicate with a central server storing all of the data from the grantee sites. Standardized data collection across all grantee sites is essential to drawing meaningful conclusions about the progress and direction of telemedicine.

The estimated burden is as follows: