state health departments in developing customized interventions tailored to the local context. Culturally appropriate interventions will increase tuberculin skin testing and patient adherence to treatment for active TB disease and latent TB infection. In addition, the results can be used to develop targeted outreach, as well as customized communication protocols, patient education materials, incentives, and enablers. Finally, the study will produce a valid interview instrument that TB clinics can adopt for their own assessments of TB beliefs and attitudes among the local communities they serve. There are no cost to respondents.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs.)	Total burden (in hrs.)
Foreign Born Persons (interviewed)	100	1	1	100
Total				100

Dated: October 11, 2001.

John Moore,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention.

[FR Doc. 01–26322 Filed 10–18–01; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Pediatric Oncology Subcommittee of the Oncologic Drugs 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: Pediatric Oncology Subcommittee of the Oncologic Drugs Advisory Committee.

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

Date and Time: The meeting will be held on November 28, 2001, from 8 a.m. to 5 p.m.

Location: Holiday Inn, Ballrooms A and B, Two Montgomery Village Ave., Gaithersburg, MD.

Contact: Kimberly Littleton Topper or Karen M. Templeton-Somers, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7001, or by e-mail at TopperK@cder.fda.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12542. Please call the Information Line for up-to-date information on this meeting.

Agenda: The subcommittee will discuss the implementation of the pediatric rule with regard to study designs, ethical and developmental considerations, and extrapolation of findings from adult to pediatric cancer patients.

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the subcommittee. Written submissions may be made to the contact person by November 16, 2001. Oral presentations from the public will be scheduled between approximately 8:15 a.m. and 8:45 a.m., and 1 p.m. and 1:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 16, 2001, 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.

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

Dated: October 12, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–26314 Filed 10–18–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Advisory Committee for Pharmaceutical Science; 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: Advisory Committee for Pharmaceutical Science.

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

Date and Time: The meeting will be held on November 28 and 29, 2001, from 8:30 a.m. to 5:30 p.m.

Location: Center for Drug Evaluation and Research Advisory Committee conference room 1066, 5630 Fishers Lane, Rockville, MD.

Contact: Nancy Chamberlin, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1076), Rockville, MD 20857, 301–827–7001, e-mail: CHAMBERLINN@cder.fda.gov, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12539. Please call the Information Line for up-to-date information on this meeting.

Agenda: On November 28, 2001, the committee will: (1) Discuss the current status of, and future plans for, the FDA draft guidance entitled ''ANDAs: Blend Uniformity Analysis;'' (2) discuss and provide direction for the Process Analytical Technology Subcommittee; (3) discuss and provide comments on stability testing and shelf life; and (4) receive updates from subcommittees and on other Center for Drug Evaluation and Research guidance documents. On November 29, 2001, the committee will: (1) Receive updates on FDA research in dermatopharmacokinetics, and (2) discuss and provide comments on bioequivalence issues.

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 November 15, 2001. Oral presentations from the public will be scheduled between approximately 1:30 p.m. and 2:30 p.m. on November 28, 2001, and between approximately 11 a.m. and 12 noon on November 29,