

to the needs of victims of domestic violence.

WomanKind strives to: (1) Increase health care providers' capacity and motivation to identify and refer battered women to WomanKind advocates from several hospital departments, (2) facilitate clients' decisions to alter their circumstances, and (3) work with clients to identify and access existing community services that provide practical support in developing and implementing a plan for change.

The evaluation is being conducted to determine the extent to which the objectives listed above are achieved and to identify the integration and level of contribution made by each specific program element. If proven effective, this program could be used with other domestic violence prevention strategies to reduce the incidence of domestic violence.

Respondents	No. of re-spond-ents	No. of re-sponses/ respond-ent	Avg. bur-den/re-sponse (in hours)
Hospital Staff KABB Survey—Census 1 & 6 month & year .....	950	3	.17
Hospital Staff KABB Survey—Trainees Immediate Post-test .....	250	1	.17
Volunteer Advocate KABB Survey .....	30	4	.17
Womankind Client KABB Survey .....	450	4	.25
Control Client KABB Survey .....	200	4	.25
Hospital Staff Training Evaluation ...	250	1	.08
Volunteer Advocate Training Evaluation .....	30	6	.08
Hospital Staff Trainer Evaluation .....	250	1	.08
Volunteer Trainer Evaluation .....	30	6	.08

The total annual burden is 1262. Send comments to Allison Eydt; Human Resources and Housing Branch, New Executive Office Building, Room 10235; Washington, DC 20503.

Dated: November 22, 1995.

Joseph R. Carter,

*Acting Associate Director for Management and Operations, Centers for Disease Control and Prevention (CDC).*

[FR Doc. 95-29167 Filed 11-28-95; 8:45 am]

BILLING CODE 4163-18-P

**Mine Health Research Advisory Committee; Meeting**

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC), announces the following committee meeting.

*Name:* Mine Health Research Advisory Committee.

*Times and Dates:* 9 a.m.-5 p.m., December 13, 1995. 9 a.m.-12 noon, December 14, 1995.

*Place:* The Washington Court Hotel, Hermitage Room, 525 New Jersey Avenue, NW., Washington, DC 20001.

*Status:* Open to the public, limited only by the space available.

*Purpose:* The Committee is charged with advising the Secretary of Health and Human Services on matters involving or relating to mine health research, including grants and contracts for such research. Additionally, the Committee assesses mine health research needs and advises on the conduct of mine health research.

*Matters to be discussed:* The agenda will include a discussion on current trends and future directions in the mining industries, NIOSH plans for a National Occupational Research Agenda, critical mining research needs for the next decade, NIOSH Recommended Standard for Occupational Exposure to Respirable Coal Mine Dust, subcommittee reports, report from 2nd International Conference on Health Miners and a report from the Director of NIOSH.

Agenda items are subject to change as priorities dictate.

*Contact Person for More Information:* Gregory R. Wagner, M.D., Executive Secretary, Division of Respiratory Disease Studies, NIOSH, CDC, Mailstop 220, 944 Chestnut Ridge Road, Morgantown, West Virginia 26505, telephone 304/291-4474.

Dated: November 21, 1995.

Carolyn J. Russell,

*Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).*

[FR Doc. 95-29168 Filed 11-28-95; 8:45 am]

BILLING CODE 4163-19-M

**Board of Scientific Counselors, National Center for Infectious Diseases; Meeting**

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Centers for Disease Control and Prevention (CDC)

announces the following committee meeting.

*Name:* Board of Scientific Counselors, National Center for Infectious Diseases (NCID).

*Times and Dates:* 8:30 a.m.-5 p.m., December 7, 1995. 8:30 a.m.-2:30 p.m., December 8, 1995.

*Place:* CDC, Auditorium B, 1600 Clifton Road, NE, Atlanta, Georgia 30333.

*Status:* Open to the public, limited only by the space available.

*Purpose:* The Board of Scientific Counselors, NCID, provides advice and guidance to the Director, CDC, and Director, NCID, in the following areas: program goals and objectives; strategies; program organization and resources for infectious disease prevention and control; and program priorities.

*Matters to be Discussed:* The agenda will focus on:

1. Progress and Plans for Implementation of CDC Emerging Infectious Diseases: Prevention Strategy for the U.S.
2. Personnel and Facility Updates.
3. Committee on Science, Engineering, and Technology Working Group Report on Emerging and Re-emerging Infectious Diseases; Recommendations for CDC; Implementation of Interagency Task Force.
4. Update on World Health Organization Emerging Infections Programs.
5. Research on Emerging and Re-emerging Infectious Diseases.
6. Health Communication Strategies.
7. Current Scientific Investigations.

Other agenda items include announcements/introductions; NCID update; late breakers; and follow-up on actions recommended by the Board (May 1995).

Agenda items are subject to change as priorities dictate.

Written comments are welcome and should be received by the contact person listed below prior to the opening of the meeting.

*Contact Person for More Information:* Diane S. Holley, Office of the Director, NCID, CDC, Mailstop C-20, 1600 Clifton Road, NE, Atlanta, Georgia 30333, telephone 404/639-0078.

Dated: November 20, 1995.

Carolyn J. Russell,

*Director, Management Analysis and Services Office, Centers for Disease Control and Prevention (CDC).*

[FR Doc. 95-29191 Filed 11-28-95; 8:45 am]

BILLING CODE 4163-18-M

**Food and Drug Administration**

**Advisory Committee; Notice of Meeting**

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

**ACTION:** Notice.

**SUMMARY:** This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). This notice also summarizes the procedures for the meeting and methods by which

interested persons may participate in open public hearings before FDA's advisory committees.

FDA has established an Advisory Committee Information Hotline (the hotline) using a voice-mail telephone system. The hotline provides the public with access to the most current information on FDA advisory committee meetings. The advisory committee hotline, which will disseminate current information and information updates, can be accessed by dialing 1-800-741-8138 or 301-443-0572. Each advisory committee is assigned a 5-digit number. This 5-digit number will appear in each individual notice of meeting. The hotline will enable the public to obtain information about a particular advisory committee by using the committee's 5-digit number. Information in the hotline is preliminary and may change before a meeting is actually held. The hotline will be updated when such changes are made.

**MEETING:** The following advisory committee meeting is announced:

#### **Oncologic Drugs Advisory Committee**

*Date, time, and place.* December 14, 1995, 8 a.m., Holiday Inn—Bethesda, Versailles Ballrooms I and II, 8120 Wisconsin Ave., Bethesda, MD.

*Type of meeting and contact person.* Open public hearing, 8 a.m. to 9 a.m., unless public participation does not last that long; open committee discussion, 9 a.m. to 5 p.m.; Stephen P. Pollitt or Adele S. Seifried, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-443-4695, or FDA Advisory Committee Information Hotline, 1-800-741-8138 (301-443-0572 in the Washington, DC area), Oncologic Drugs Advisory Committee, code 12542.

*General function of the committee.* The committee reviews and evaluates data on the safety and effectiveness of marketed and investigational human drugs for use in the treatment of cancer.

*Agenda—Open public hearing.* Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Those desiring to make formal presentations should notify the contact person before December 8, 1995, 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 required to make their comments.

*Open committee discussion.* The committee will discuss: (1) FDA

oncology initiatives, (2) new drug application (NDA) 20-587, Sterile Aerosol Talc (Talc of Luzenac PR 784, Bryan Corp.), for treatment of malignant pleural effusion, and (3) product license application (PLA) 94-0308, NR-LU-10 Fab (Karl Thomae GmbH), for primary staging of patients with biopsy confirmed small cell lung cancer.

FDA public advisory committee meetings may have as many as four separable portions: (1) An open public hearing, (2) an open committee discussion, (3) a closed presentation of data, and (4) a closed committee deliberation. Every advisory committee meeting shall have an open public hearing portion. Whether or not it also includes any of the other three portions will depend upon the specific meeting involved. There are no closed portions for the meetings announced in this notice. The dates and times reserved for the open portions of each committee meeting are listed above.

The open public hearing portion of each meeting shall be at least 1 hour long unless public participation does not last that long. It is emphasized, however, that the 1 hour time limit for an open public hearing represents a minimum rather than a maximum time for public participation, and an open public hearing may last for whatever longer period the committee chairperson determines will facilitate the committee's work.

Public hearings are subject to FDA's guideline (subpart C of 21 CFR part 10) concerning the policy and procedures for electronic media coverage of FDA's public administrative proceedings, including hearings before public advisory committees under 21 CFR part 14. Under 21 CFR 10.205, representatives of the electronic media may be permitted, subject to certain limitations, to videotape, film, or otherwise record FDA's public administrative proceedings, including presentations by participants.

Meetings of advisory committees shall be conducted, insofar as is practical, in accordance with the agenda published in this Federal Register notice. Changes in the agenda will be announced at the beginning of the open portion of a meeting.

Any interested person who wishes to be assured of the right to make an oral presentation at the open public hearing portion of a meeting shall inform the contact person listed above, either orally or in writing, prior to the meeting. Any person attending the hearing who does not in advance of the meeting request an opportunity to speak will be allowed to make an oral presentation at the

hearing's conclusion, if time permits, at the chairperson's discretion.

The agenda, the questions to be addressed by the committee, and a current list of committee members will be available at the meeting location on the day of the meeting.

Transcripts of the open portion of the meeting may be requested in writing from the Freedom of Information Office (HFI-35), Food and Drug Administration, rm. 12A-16, 5600 Fishers Lane, Rockville, MD 20857, approximately 15 working days after the meeting, at a cost of 10 cents per page. The transcript may be viewed at the Dockets Management Branch (HFA-305), Food and Drug Administration, rm. 1-23, 12420 Parklawn Dr., Rockville, MD 20857, approximately 15 working days after the meeting, between the hours of 9 a.m. and 4 p.m., Monday through Friday. Summary minutes of the open portion of the meeting may be requested in writing from the Freedom of Information Office (address above) beginning approximately 90 days after the meeting.

This notice is issued under section 10(a)(1) and (2) of the Federal Advisory Committee Act (5 U.S.C. app. 2), and FDA's regulations (21 CFR part 14) on advisory committees.

Dated: November 20, 1995.

David A. Kessler,

*Commissioner of Food and Drugs.*

[FR Doc. 95-29085 Filed 11-28-95; 8:45 am]

BILLING CODE 4160-01-F

#### **Grassroots Regulatory Partnership Meeting; Southwest Region; Human and Veterinary Drug Industry**

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

**ACTION:** Notice of a public meeting.

**SUMMARY:** The Food and Drug Administration (FDA) (Office of External Affairs, Office of Regulatory Affairs, Office of the Southwest Region, Center for Drug Evaluation and Research, Center for Biologics Evaluation and Research, and Center for Veterinary Medicine) is announcing a free public meeting as a followup to a meeting held in April 1995. FDA Office of the Southwest Region will meet with interested persons in the Southwest Region to address specific issues related to the human and veterinary drug industry. The agency is holding this meeting to promote the President's initiative for a partnership approach between front-line regulators and the people affected by the work of this agency.