

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

Dated: April 7, 2003.

**Linda Arey Skladany,**  
*Associate Commissioner for External Relations.*

[FR Doc. 03-9032 Filed 4-11-03; 8:45 am]

BILLING CODE 4160-01-S

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### Manufacturing Subcommittee of the 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:* Manufacturing Subcommittee of the 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 May 21 and 22, 2003, from 8:30 a.m. to 5 p.m.

*Location:* Marriott Washingtonian Center, Ballrooms A, B, C, and D, 9751 Washingtonian Blvd., Gaithersburg, MD.

*Contact Person:* Kathleen Reedy, Center for Drug Evaluation and Research (HFD-21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301-827-7001, e-mail: [REEDYK@cder.fda.gov](mailto:REEDYK@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 May 21, 2003, the subcommittee will discuss: (1) The mission of the subcommittee; and (2) direction of the Pharmaceutical Current Good Manufacturing Practices (CGMPs) for the 21st Century: A Risk-Based Approach. On May 22, 2003, the subcommittee will discuss: (1) The regulatory approaches regarding aseptic manufacturing; and (2) process analytical technologies and transition from the Advisory Committee for Pharmaceutical Science—Process

Analytical Technologies Subcommittee to Manufacturing Subcommittee.

*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 May 13, 2003. Oral presentations from the public will be scheduled between approximately 11:30 a.m. and 12:30 p.m. on both days. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before May 13, 2003, 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.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact Kathleen Reedy at least 7 days in advance of the meeting.

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

Dated: April 7, 2003.

**Linda Arey Skladany,**  
*Associate Commissioner for External Relations.*

[FR Doc. 03-9029 Filed 4-11-03; 8:45 am]

BILLING CODE 4160-01-S

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 03N-0134]

#### Team Biologics Program Effectiveness; Public Meeting

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

**ACTION:** Notice of public meeting; request for comments.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the following public meeting: Team Biologics Program Effectiveness. The Center for Biologics Evaluation and Research and the Office of Regulatory Affairs, FDA, are sponsoring an open public meeting to solicit views and comments in an effort to measure the effectiveness of the Team Biologics

Program as it relates to the inspections of manufacturers of vaccines, allergenics, fractionated plasma products, licensed in vitro diagnostics, and therapeutic products. The goal of the public meeting is to give stakeholders the opportunity to provide input on how they think the agency should measure the effectiveness of the Team Biologics Program. We will use the information obtained to identify criteria to prospectively evaluate the Team Biologics Program.

**DATES:** The public meeting will be held on Wednesday, May 21, 2003, from 8 a.m. to 12 noon.

Submit requests via fax or e-mail by May 1, 2003, to make an oral presentation. Submit a copy of all presentation materials by May 15, 2003. If you are not making an oral presentation, submit registration information by May 12, 2003.

Submit written or electronic comments by June 10, 2003.

**ADDRESSES:** The public meeting will be held at the Parklawn Bldg., conference room D, 5600 Fishers Lane, Rockville, MD 20857.

Submit requests to make an oral presentation, registration information, and any presentation material to Melanie Whelan (*see FOR FURTHER INFORMATION CONTACT*). The requested registration information is listed in section II of this document.

Submit written comments to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.fda.gov/dockets/ecomments>.

**FOR FURTHER INFORMATION CONTACT:** Melanie N. Whelan, Center for Biologics Evaluation and Research (HFM-43), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448, 301-827-2000, FAX 301-827-3079, or e-mail: [Whelan@cber.fda.gov](mailto:Whelan@cber.fda.gov).

#### SUPPLEMENTARY INFORMATION:

##### I. Scope of Public Meeting

FDA is seeking input on ways to evaluate the Team Biologics Program. The Team Biologics Program, established in 1997, is a partnership between FDA's Center for Biologics Evaluation and Research and the Office of Regulatory Affairs, which uses the diverse skills and knowledge of both organizations to focus resources on inspectional and compliance issues in the biologics area. Comments are sought at this public meeting about specific methods, tools, criteria, and metrics that could be used in this effort. In presentations we ask that you