Location: Center for Drug Evaluation and Research Advisory Committee conference rm. 1066, 5630 Fishers Lane, Rockville, 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, or e-mail: 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 October 21, 2002, the committee will: (1) Receive summary reports and provide direction for the Nonclinical Studies Subcommittee and the Process Analytical Technologies Subcommittee; (2) receive updates on risk-based chemistry manufacturing control review and blend uniformity; and (3) discuss and provide comments on regulatory issues related to crystal habits—polymorphism. On October 22, 2002, the committee will: (1) Discuss and provide direction for future subcommittee—Good Manufacturing Practices/Manufacturing Subcommittee; and (2) discuss manufacturing issues; sterile drug products produced by aseptic processing.

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 October 14, 2002. Oral presentations from the public will be scheduled between approximately 11:45 a.m. and 12:45 p.m. on October 21, 2002, and 1 p.m. and 2 p.m. on October 22, 2002. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before October 14, 2002, 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: September 22, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–24812 Filed 9–30–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Process Analytical Technologies 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: Process Analytical Technologies 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 October 23, 2002, from 8:30 a.m. to 5 p.m.

Location: Ramada Inn, Georgetown and Montrose Conference Rooms, 1775 Rockville Pike, Rockville, 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, or e-mail: 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: The subcommittee will discuss: (1) Computer systems validation—21 CFR part 11 issues pertinent to process analytical technologies (PAT), (2) a PAT case study, and (3) rapid microbiology testing.

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 October 14, 2002. Oral presentations from the public will be scheduled between approximately 11:30 a.m. and 12:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before October 14, 2002, 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.

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

Dated: September 22, 2002.

Linda Arey Skladany,

Senior Associate Commissioner for External Relations.

[FR Doc. 02–24813 Filed 9–30–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

RIN 1018–AI55

Fiscal Year (FY) 2002 Landowner Incentive Program (Non Tribal Portion) for States, Territories and the District of Columbia; Final Policy With Implementation Guidelines, and Request for Proposals

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Final policy with implementation guidelines; notice of request for proposals.

SUMMARY: The Department of the Interior and Related Agencies Appropriations Act 2002 allocated $40 million from the Land and Water Conservation Fund for conservation grants to States, the District of Columbia, Puerto Rico, Guam, the United States Virgin Islands, the Northern Mariana Islands, and American Samoa (hereafter referred to collectively as States), and Tribes under a Landowner Incentive Program (LIP).