available to the manufacturer, packer, or distributor of the dietary supplement that bears such a statement on its labeling or in its labeling. This estimate is based on the average number of notification submissions received by the agency in the preceding 18 months.

Dated: October 19, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–26885 Filed 10–24–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Request for Participants at the Process Analytical Technologies Subcommittee of the Advisory Committee for Pharmaceutical Science

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is requesting names of qualified persons to participate on the Process Analytical Technologies Subcommittee (the Subcommittee) of the Advisory Committee for Pharmaceutical Science. The Subcommittee will identify and report to the Advisory Committee for Pharmaceutical Science on scientific issues related to application and validation of online process technologies such as near infrared and Raman spectroscopy and imaging methods for application in the manufacture of drug substances and drug products. The Subcommittee will also report on the potential benefits and risks associated with the application of these new technologies to public health and, as part of this analysis, evaluate the feasibility of the parametric release concept.

FDA has a special interest in ensuring that women, minority groups, and individuals with disabilities are adequately represented and, therefore encourages recommendations of qualified candidates from these groups. Final selections from among qualified candidates will be based on the expertise demonstrated and previous experience with online process technologies.

DATES: All applications should be received by November 30, 2001.

ADDRESSES: Submit applications to David Morley (address below).

FOR FURTHER INFORMATION CONTACT: David Morley, Office of Testing and Research (HFD–900), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–5186, FAX 301–827–3787, e-mail: morleyd@cder.fda.gov.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is seeking qualified persons to participate on the Process Analytical Technologies Subcommittee being formed under the Advisory Committee for Pharmaceutical Science. The Subcommittee will identify and report on the current state of technology, validation procedures, and the mechanistic basis of online process controls in both drug development and scaleup. These participants are not members of the Subcommittee and will not be voting on any issues, but they are encouraged to participate in the discussion of the issues. The Subcommittee will evaluate the potential for enhancing product quality and providing public health benefit.

II. Selection Criteria

Persons from government, industry, academia, and other organizations (such as research institutes) applying to participate on the Subcommittee should have exceptional accomplishments and be leading technical experts in the appropriate fields. In particular, expertise in application of the following scientific disciplines to pharmaceutical development and pharmaceutical manufacturing processes is desired: Process analytical chemistry, pharmaceutics, industrial pharmacy, chemical engineering, pharmaceutical analysis, chemometrics, pattern recognition, computer expert systems, information technology, and statistics.

III. Application Procedures

Any interested person should submit appropriate biographical material and a list of scientific publications relevant to the Subcommittee to the contact person listed above.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to advisory committees.

Dated: October 17, 2001.

Linda A. Suydam,

Senior Associate Commissioner.

[FR Doc. 01–26834 Filed 10–24–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Organ Procurement and Transplantation Network

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice of meeting of the Advisory Committee on Organ Transplantation.

SUMMARY: Pursuant to Public Law 92–463, the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the first meeting of the Advisory Committee on Organ Transplantation (ACOT), Department of Health and Human Services (HHS). The meeting will be held from approximately 8:15 a.m. to 6 p.m. on December 3, 2001, and from 8 a.m. to 5:15 p.m. on December 4, 2001, at the Hyatt Dulles, at Dulles International Airport, 2300 Dulles Corner Boulevard, Herndon, Virginia 20171. The meeting will be open to the public; however, seating is limited and pre-registration is encouraged (see below).

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. section 217a, section 222 of the Public Health Service Act, as amended, and 42 CFR 121.12 (2000), the ACOT was established to assist the Secretary in enhancing organ donation, ensuring that the system of organ transplantation is grounded in the best available medical science, and assuring the public that the system is as effective and equitable as possible, and, thereby, increasing public confidence in the integrity and effectiveness of the transplantation system. The ACOT is composed of 41 members, including the Chair. Members are non-governmental individuals with diverse backgrounds in fields such as organ donation, health care public policy, transplantation medicine and surgery, critical care medicine and other medical specialties involved in the identification and referral of donors, non-physician transplant professions, nursing, epidemiology, immunology, law and bioethics, behavioral sciences, economics and statistics, as well as representatives of transplant candidates, transplant recipients, organ donors, and family members.

The ACOT will consider a number of subjects relating to the means of expanding the donor pool and increasing organ donation; and it will also review the organ allocation policies submitted by the Organ Procurement

and Transplantation Network (OPTN) to HHS for approval. The draft meeting agenda and a registration form are available on the Division of Transplantation's Web site: *http://www.hrsa.gov/osp/dot/whatsnew.htm* or the Department's donation Web site at: *http://www.organdonor.gov/news.htm.* The completed registration form should be submitted by facsimile to McFarland and Associates, Inc., the logistic support contractor for the meeting, at FAX number (301) 589–2567. Individuals without access to the Internet who wish to register may call McFarland and Associates, Inc., at 301–562–5362. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the ACOT Executive Director, Jack Kress, in advance of the meeting. Mr. Kress may be reached by telephone at 301–443–8653, by e-mail at: *jkress2@hrsa.gov,* or in writing at the address of the Division of Transplantation provided below. Management and support services for ACOT functions are provided by the Division of Transplantation, Office of Special Programs, HRSA, Room 7C–22, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857. Telephone 301–443–7577.

There will be a limited period of time for presentation of selected public comments before the Committee considers each allocation policy and donation issue. The public may review the current and proposed modified OPTN policies on the Division of Transplantation's Web site at: *http://www.hrsa.gov/osp/dot/whatsnew.htm* and may also obtain this material, as well as reports of committee discussions of these policies by contacting the Division of Transplantation at 301–443–7577. While public comments are welcome for possible presentation, please note that the Committee will be working with a full agenda and a limited amount of time. Therefore, to facilitate this process, we recommend that individuals interested in providing public comments submit those comments in writing by November 16, 2001, to the Executive Director of the Committee (address above). The Department reserves the right to select comments from among those submitted for oral presentation within the time available, although it will include all comments in the record of the ACOT meeting.

Dated: October 19, 2001.

Elizabeth M. Duke,

Acting Administrator.

[FR Doc. 01–26932 Filed 10–24–01; 8:45 am]

BILLING CODE 4165–15–P

INTER-AMERICAN FOUNDATION

Sunshine Act Meeting

TIME AND DATE: October 29, 2001, 9:00 a.m.–3:30 p.m.

PLACE: Inter-American Foundation, 901 N. Stuart Street, 10th Floor, Arlington, VA 22203.

STATUS: Open session.

MATTERS TO BE CONSIDERED:

- Approval of the Minutes of the April 23, 2001, Meeting of the Board of Directors
- President's Report
- Country Priorities Presentation
- Overview of IAF
- IAF Strategic Plan for Fiscal Years 2002–2007
- IAF 2000 Results Report
- IAF Web Site
- Review of a Sample of Successful Closed-out Grants
- IAF Experience with Corporate Partners
- Improving the Role, Mission, and Operations of the IAF

CONTACT PERSON FOR MORE INFORMATION: Carolyn Karr, General Counsel, (703) 306–4350.

Dated: October 22, 2001.

Carolyn Karr,

General Counsel.

[FR Doc. 01–27061 Filed 10–23–01; 3:02 pm]

BILLING CODE 7025–01–M

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Notice of Receipt of Applications for Permit

Endangered Species

The public is invited to comment on the following application(s) for a permit to conduct certain activities with endangered species. This notice is provided pursuant to section 10(c) of the Endangered Species Act of 1973, *as amended* (16 U.S.C. 1531, *et seq.*). Written data, comments, or requests for copies of these complete applications should be submitted to the Director (address below) and must be received within 30 days of the date of this notice.

Applicant: Karl W. Widl, Los Altos, OH, PRT–048967.

The applicant requests a permit to import the sport-hunted trophy of one male bontebok (*Damaliscus pygargus dorcas*) culled from a captive herd maintained under the management program of the Republic of South Africa, for the purpose of enhancement of the survival of the species.

Applicant: Thomas Heideman, Lockport, NY, PRT–049027.

The applicant requests a permit to import the sport-hunted trophy of one male bontebok (*Damaliscus pygargus dorcas*) culled from a captive herd maintained under the management program of the Republic of South Africa, for the purpose of enhancement of the survival of the species.

Applicant: Scott Starbard, Edmonds, WA, PRT–049031.

The applicant requests a permit to import the sport-hunted trophy of one male bontebok (*Damaliscus pygargus dorcas*) culled from a captive herd maintained under the management program of the Republic of South Africa, for the purpose of enhancement of the survival of the species.

Applicant: Daniel Dienstbier, Omaha, NE, PRT–049032.

The applicant requests a permit to import the sport-hunted trophy of one male bontebok (*Damaliscus pygargus dorcas*) culled from a captive herd maintained under the management program of the Republic of South Africa, for the purpose of enhancement of the survival of the species.

The U.S. Fish and Wildlife Service has information collection approval from OMB through March 31, 2004, OMB Control Number 1018–0093. Federal Agencies may not conduct or sponsor and a person is not required to respond to a collection of information unless it displays a current valid OMB control number.

Documents and other information submitted with these applications are available for review, subject to the requirements of the Privacy Act and Freedom of Information Act, by any party who submits a written request for a copy of such documents within 30 days of the date of publication of this notice to: U.S. Fish and Wildlife Service, Division of Management Authority, 4401 North Fairfax Drive, Room 700, Arlington, Virginia 22203, telephone 703/358–2104 or fax 703/358–2281.

Dated: October 12, 2001.

Anna Barry,

Senior Permit Biologist, Branch of Permits, Division of Management Authority.

[FR Doc. 01–26876 Filed 10–24–01; 8:45 am]

BILLING CODE 4310–55–P