

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 fifth meeting of the Advisory Committee on Organ Transplantation (ACOT), Department of Health and Human Services (HHS). The meeting will be held from approximately 9 a.m. to 5:30 p.m. on November 6, 2003, and from 9 a.m. to 5 p.m. on November 7, 2003, at the Renaissance Washington, DC Hotel, 999 Ninth Street, NW, Washington, DC. 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. 217a, Section 222 of the Public Health Service Act, as amended, and 42 CFR 121.12 (2000), 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. ACOT is composed of up to 41 members, including the Chair. Members are serving as Special Government Employees and have 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.

ACOT will hear and discuss reports from the following ACOT subcommittees: Organ Supply Concerns, Recipient Concerns, Public Concerns, and Allocation Concerns.

The draft meeting agenda will be available on October 15 on the

Department's donation Web site at <http://www.organdonor.gov/acot.html>.

A registration form is available on the Department's donation Web site at <http://www.organdonor.gov/acot.html>. The completed registration form should be submitted by facsimile to Professional and Scientific Associates (PSA), the logistical support contractor for the meeting, at fax number (703) 234-1701. Individuals without access to the Internet who wish to register may call Lora Robinson with PSA at (703) 234-1753. 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: 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, Health Resources and Services Administration, 5600 Fishers Lane, Parklawn Building, Room 16C-17, Rockville, Maryland 20857; telephone number 301-443-7577.

After the presentation of the subcommittee reports, members of the public will have an opportunity to provide comments on the subcommittee reports. Because of the Committee's full agenda and the time frame in which to cover the agenda topics, public comment will be limited. All public comments will be included in the record of the ACOT meeting.

Dated: September 23, 2003.

Jane M. Harrison,

Director, Division of Policy Review and Coordination.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

The National Institutes of Health

Submission for OMB Review; Comment Request; Customer/Partner Satisfaction Surveys The Effectiveness of the National Institute on Drug Abuse's Publications Project

Summary: Under the provisions of Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institute on Drug Abuse (NIDA), the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information

collection listed below. This proposed information collection was previously published in the **Federal Register** on June 3, 2003, page 33168 and allowed 60-days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institutes of Health may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a current valid OMB control number.

Proposed Collection: Title: The Effectiveness of NIDA's Publications Project. **Type of Information Collection Request:** New. **Need and Use of Information Collection:** This is a request for a three-year generic clearance to study the level of customer satisfaction in relation to public health information publications produced by the Institute. This effort is made according to Executive Order 12862, which directs Federal agencies that provide significant services directly to the public to survey customers to determine the kind and quality of services they want and their level of satisfaction with existing services. The primary purpose of the Project is to assess NIDA's effectiveness in developing and disseminating selected public health information publications designed to promote the use of science-based evidence to improve drug abuse and addiction prevention, treatment, and policy. A multi-method approach (survey, in-person interviews, focus groups) will be used to determine the use and usefulness of selected NIDA public health information publications for several of NIDA's key audiences. Measures will include outcomes associated with the following variables: Knowledge/awareness of the publications, receipt of the publications, reading of the publications, use of the publications, perceived utility of the publications, and the impact of the publications on the use of science-based evidence to improve drug abuse and addiction prevention, treatment, and policy. **Frequency of Response:** Respondents will not be asked to respond more frequently than annually or biennially. **Affected Public:** Individuals or households; state or local governments; organizations; businesses or educational institutions. **Type of Respondents:** Community coalition leaders, drug abuse treatment and prevention service providers, drug abuse researchers, Native Americans, middle school science and health educators, public health policy makers