

*Dates of Meeting:* September 29–30, 1997.  
*Time of Meeting:* 9:00 a.m.–recess,  
 September 29. 8:30 a.m.–adjournment,  
 September 30.

*Place of Meeting:* II Rockledge Center,  
 Room 9104, 6701 Rockledge Drive, Bethesda,  
 Maryland 20892.

*Agenda:* Review and prioritize new  
 research concepts and proposed initiatives  
 developed by the Division of Blood Disease  
 Scientific Research Groups.

*Contact Person:* Carol H. Letendre, Ph.D.,  
 NHLBI/DBDR, II Rockledge Center, 6701  
 Rockledge Drive, Rm. 10162, MSC 7950,  
 Bethesda, Maryland 20892, (301) 435-0080.  
 (Catalog of Federal Domestic Assistance  
 Programs Nos. 93.837, Heart and Vascular  
 Diseases Research; 93.838, Lung Diseases  
 Research; and 93.839, Blood Diseases and  
 Resources Research, National Institutes of  
 Health)

Dated: August 28, 1997.

**LaVeen M. Ponds,**

*Policy Analyst, NIH CMO.*

[FR Doc. 97-23481 Filed 9-3-97; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute of Child Health and Human Development; Notice of Closed Meetings

Pursuant to Section 10(d) of the  
 Federal Advisory Committee Act, as  
 amended (5 U.S.C. Appendix 2), notice  
 is hereby given of the following  
 National Institute of Child Health and  
 Human Development Special Emphasis  
 Panel (SEP) meeting:

*Name of SEP:* Neurodevelopmental Battery  
 for the Maternal Lifestyle Study  
 (Teleconference).

*Date:* September 3, 1997.

*Time:* 1:00 p.m. (ET)–adjournment.

*Place:* 61E Building, Room 5E03, 6100  
 Executive Boulevard, Rockville, MD 20852.

*Contact Person:* Hameed A. Khan, Ph.D.,  
 61E Building, Rm. 5E03, Rockville, Maryland  
 20852, Telephone: 301-496-1485.

*Purpose/Agenda:* To evaluate and review a  
 contract proposal.

This meeting will be closed in accordance  
 with the provisions set forth in sections  
 552b(c)(4) and 552b(c)(6), Title 5 U.S.C. The  
 discussions of this proposal could reveal  
 confidential trade secrets or commercial  
 property such as patentable material and  
 personal information concerning individuals  
 associated with the proposal, the disclosure  
 of which would constitute a clearly  
 unwarranted invasion of personal privacy.

This notice is being published less than 15  
 days prior to the meeting due to the urgent  
 need to meet timing limitations imposed by  
 the review and funding cycle.

(Catalog of Federal Domestic Assistance  
 Program Nos. [93.864, Population Research  
 and No. 93.865, Research for Mothers and  
 Children], National Institutes of Health)

Dated: August 27, 1997.

**LaVeen M. Ponds,**

*Policy Analyst, NIH CMO.*

[FR Doc. 97-23476 Filed 9-3-97; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute of Diabetes and Digestive and Kidney Diseases; Notice of Closed Meeting

Pursuant to Section 10(d) of the  
 Federal Advisory Committee Act, as  
 amended (5 U.S.C. Appendix 2), notice  
 is hereby given of the following  
 National Institute of Diabetes and  
 Digestive and Kidney Diseases Special  
 Emphasis Panel meeting:

*Name of SEP:* ZDK1-GRB-6-J2.

*Date:* November 19, 1997.

*Time:* 7:30 pm.

*Place:* Indiana University, School of  
 Medicine, 635 Barnhill Drive, Indianapolis,  
 IN 46202-5120.

*Contact Person:* Sharee Pepper, Ph.D.,  
 Scientific Review Administrator, Review  
 Branch, NIDDK, Natcher Building, Room  
 6as-25E, National Institutes of Health,  
 Bethesda, Maryland 20892-6600, Phone:  
 (301) 594-7798.

*Purpose/Agenda:* To review and evaluate  
 grant applications.

This meeting will be closed in accordance  
 with the provisions set forth in sections  
 552b(c)(4) and 552b(c)(6), Title 5 U.S.C.  
 Applications and/or proposals and the  
 discussions could reveal confidential trade  
 secrets or commercial property such as  
 patentable material and personal information  
 concerning individuals associated with the  
 applications and/or proposals, the disclosure  
 of which would constitute a clearly  
 unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance  
 Program No. 93.847-849, Diabetes, Endocrine  
 and Metabolic Diseases; Digestive Diseases  
 and Nutrition; and Kidney Diseases, Urology  
 and Hematology Research, National Institutes  
 of Health)

Dated: August 28, 1997.

**LaVeen M. Ponds,**

*Policy Analyst, NIH CMO.*

[FR Doc. 97-23480 Filed 9-3-97; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute on Drug Abuse; Notice of Closed Meeting

Pursuant to Section 10(d) of the  
 Federal Advisory Committee Act, as  
 amended (5 U.S.C. Appendix 2), notice

is hereby given of the following  
 National Institute on Drug Abuse  
 (NIDA) Special Emphasis Panel  
 meeting:

*Purpose/Agenda:* To evaluate and review  
 contract proposals.

*Name of Committee:* NIDA Special  
 Emphasis Panel (Identification and Analysis  
 of THC, Cocaine, and Other Drugs of Abuse  
 and Their Metabolites in Meconium).

*Date:* September 8, 1997.

*Time:* 10:00 a.m.

*Place:* Contracts Management Branch,  
 National Institute on Drug Abuse, NIH, 5600  
 Fishers Lane, Room 10-49, Rockville, MD  
 20857, (Telephone Conference).

*Contact Person:* Mr. Eric Zatman, Contract  
 Review Specialist, Office of Extramural  
 Program Review, National Institute on Drug  
 Abuse, 5600 Fishers Lane, Room 10-42,  
 Rockville, MD 20857, Telephone (301) 443-  
 1644.

This notice is being published less than 15  
 days prior to the meeting due to the urgent  
 need to meet timing limitations imposed by  
 the review and funding cycle.

The meeting will be closed in accordance  
 with provisions set forth in sections  
 552b(c)(4) and 552b(c)(6), Title 5, U.S.C. The  
 applications and/or proposals and the  
 discussions could reveal confidential trade  
 secrets or commercial property such as  
 patentable material and personal information  
 concerning individuals associated with the  
 applications and/or proposals, the disclosure  
 of which would constitute a clearly  
 unwarranted invasion of personal privacy.

(Catalog of Federal Domestic Assistance  
 Program Numbers: 93.277, Drug Abuse  
 Scientist Development, Research Scientist  
 Development, and Research Scientist  
 Awards; 93.278, Drug Abuse National  
 Research Service Awards for Research  
 Training; 93.279, Drug Abuse Research  
 Programs, National Institutes of Health)

Dated: August 28, 1997.

**LaVeen M. Ponds,**

*Policy Analyst, NIH CMO.*

[FR Doc. 97-23482 Filed 9-3-97; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Workshop on Thalidomide: Potential Benefits and Risks

Notice is hereby given of the NIH  
 workshop on "Thalidomide: Potential  
 Benefits and Risks," which will be held  
 September 9–10, 1997, in the Natcher  
 Conference Center of the National  
 Institutes of Health, 9000 Rockville  
 Pike, Bethesda, Maryland 20892. The  
 conference begins at 8:30 a.m. on  
 September 9 and at 8 a.m. on September  
 10.

Thalidomide was associated in 1961  
 with serious human teratogenicity. Its  
 use in the treatment of morning sickness

and insomnia was abandoned, and it became infamous as an example of a drug with major toxic effects. Thalidomide is now being studied as a treatment for many serious diseases, including erythema nodosum leprosum, chronic graft-versus-host disease, and aphthous ulcers in patients with and without HIV infection.

The purpose of the workshop is to provide a public forum to assess the emerging research opportunities, potential clinical applications, and accompanying risks associated with the use of thalidomide. The meeting is open to researchers, academic and community-based physicians, nurses, pharmacists, other health care professionals, industry personnel, patients, and other interested individuals.

The workshop is sponsored by the Office of Rare Diseases, the Office of Research on Women's Health, the Office of Medical Applications of Research, the National Institute of Allergy and Infectious Diseases, the National Institute of Arthritis and Musculoskeletal and Skin Diseases, the National Institute of Dental Research, and the National Institute of Child Health and Human Development of the National Institutes of Health; by the Center for Drug Evaluation and Research and the Office of Special Health Issues of the Food and Drug Administration; and by the Centers for Disease Control and Prevention.

Advance information on the conference program and conference registration materials may be obtained from Prospect Associates, 1801 Rockville Pike, Suite 500, Rockville, Maryland 20852, (301) 468-MEET; by e-mail to [raredisease@ProspectAssoc.com](mailto:raredisease@ProspectAssoc.com); or at <http://rarediseases.info.nih.gov/ord> on the World Wide Web.

Dated August 12, 1997.

**Ruth L. Kirschstein,**  
Deputy Director, National Institutes of Health.  
[FR Doc. 97-23383 Filed 9-3-97; 8:45 am]  
BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Public Health Service

#### National Toxicology Program; Availability of Technical Report on Toxicology and Carcinogenesis Studies of 1-Trans-Delta<sup>9</sup>- Tetrahydrocannabinol

The HHS' National Toxicology Program announces the availability of the NTP Technical Report on the

toxicology and carcinogenesis studies of 1-trans-delta<sup>9</sup>-tetrahydrocannabinol which is a major psychoactive component of marijuana and a widely used Schedule I substance.

Toxicology and carcinogenicity studies were conducted by administering 1-trans-delta<sup>9</sup>-tetrahydrocannabinol (THC) in corn oil to groups of 62 vehicle control male rats, 60 low-dose male rats, 70 mid- and high-dose male rats, and 60 female rats at doses of 0, 12.5, 25, or 50 mg THC/kg body weight by gavage for 104 to 105 weeks. Groups of 62 vehicle control male mice, 60 low-dose male mice, 61 mid-dose male mice, and 60 high-dose male mice and 60 female mice were administered 0, 125, 250, or 500 mg THC/kg body weight in corn oil by gavage for 104 to 105 weeks (males) or 105 to 106 weeks (females).

Under the conditions of these 2-year gavage studies, there was no evidence of carcinogenic activity<sup>1</sup> of 1-trans-delta<sup>9</sup>-tetrahydrocannabinol in male or female F344/N rats administered 12.5, 25, or 50 mg/kg. There was equivocal evidence of carcinogenic activity of THC in male and female B6C3F1 mice based on the increased incidences of thyroid gland follicular cell adenomas in 125 mg/kg groups.

Increased incidences of thyroid gland follicular cell hyperplasia occurred in male and female mice, and increased incidences of hyperplasia and ulcers of the forestomach were observed in male mice.

The incidences of mammary gland fibroadenomas and uterine stromal polyps were decreased in dosed groups of female rats, as were the incidences of pancreatic adenomas, pituitary gland adenomas, and interstitial cell adenomas of the testis in dosed male rats and liver neoplasms in dosed mice. These decreases were likely related to lower body weights in dosed animals.

Questions or comments about the Technical Report should be directed to Central Data Management at PO Box 12233, Research Triangle Park, NC 27709 or telephone (919) 541-3419.

Copies of Toxicology and Carcinogenesis Studies of 1-Trans-Delta<sup>9</sup>-Tetrahydrocannabinol (CAS No. 1972-08-3) (TR-446) are available from Central Data Management, NIEHS, MD E1-02, PO Box 12233, Research Triangle

Park, NC 27709; telephone (919) 541-3419.

Dated: August 14, 1997.

**Samuel H. Wilson,**

NIEHS Deputy Director.

[FR Doc. 97-23381 Filed 9-3-97; 8:45 am]

BILLING CODE 4140-01-M

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Public Health Service

#### Centers for Disease Control and Prevention; Statement of Organization, Functions, and Delegations of Authority

Part C (Centers for Disease Control and Prevention) of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (45 FR 67772-76, dated October 14, 1980, and corrected at 45 FR 69296, October 20, 1980, as amended most recently at 54 FR 13389-90, dated March 30, 1997) is amended to reflect the transfer of international emergency and refugee functions from the International Health Program Office (IHPO) to the National Center for Environmental Health and the transfer of child survival functions associated with Integrated Case Management activities in IHPO to the National Center for Infectious Diseases.

*Section C-B, Organization and Functions*, is hereby amended as follows:

Revise the mission and function statement for the *International Health Program Office (CG)* by deleting item (6) and renumbering the remaining items accordingly.

Revise the functional statement for the *Office of the Director (CG1)*, by deleting item (7) and renumbering the remaining items accordingly.

Revise the functional statement for the *Division of Field Services (CG6)* by deleting item (5) and renumbering the remaining items accordingly.

Delete in their entirety the title and functional statement for *Child Survival Activity (CG62)*.

Revise the functional statement for the *Division of Technical Support (CG7)* by deleting item (6).

Revise the functional statement for the *Epidemiologic Support Branch (CG72)* by deleting items (2) and (8) and renumbering the remaining items accordingly.

<sup>1</sup> The NTP uses five categories of evidence of carcinogenic activity observed in each animal study: two categories for positive results ("clear evidence" and "some evidence"), one category for uncertain findings ("equivocal evidence"), one category for no observable effect ("no evidence"), and one category for studies that cannot be evaluated because of major flaws ("inadequate study").