TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
200	30 (maximum)	6,000	0.5	3,000

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA projects 30 emergency risk related surveys per year with a sample of between 50 and 200 respondents per survey. FDA also projects a response time of 0.5 hours per response. These estimates are based on the maximum sample size per questionnaire that FDA can analyze in a timely manner. The annual frequency of response was determined by the maximum number of questionnaires that will be sent to any individual respondent. Some respondents may be contacted only one time per year, while other respondents may be contacted several times annually, depending on the human drug, biologic, or medical device under evaluation. It is estimated that, given the expected type of issues that will be addressed by the surveys, it will take 0.5 hours for a respondent to gather the requested information and fill in the answers.

Dated: October 17, 2005.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 05–21240 Filed 10–24–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Clinical Pharmacology 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: Clinical Pharmacology 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 November 14, 2005, from 8:30 a.m. to 5:30 p.m. and on November 15, 2005, from 8:30 a.m. to 1:30 p.m.

Location: Food and Drug Administration, Center for Drug Evaluation and Research Advisory Committee Conference Room, rm. 1066, 5630 Fishers Lane, Rockville, MD.

Contact Person: Mimi Phan, 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, FAX: 301–827–6776, e-mail: *phanm@cder.fda.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512539. Please call the Information Line for up-to-date information on this meeting.

Agenda: On November 14, 2005, the subcommittee will: (1) Receive an update on previous Clinical Pharmacology Subcommittee meeting recommendations and an introduction to the topics of this meeting, (2) discuss and provide comments on the evidence and process for translation of pharmacogenetic information (e.g., Cytochrome P 2C9 polymorphisms) into label updates for approved products, (3) discuss current evidence related to the pharmacogenetics of warfarin as a potential basis for label updates, and (4) discuss and provide comments on the critical path pilot project, the End-of-Phase 2A meetings which will include a case study. On November 15, 2005, the subcommittee will discuss and and provide comments on: (1) An update on the critical path biomarker-surrogate endpoint project, (2) the use of biomarker information in labels to facilitate individualizing pharmacotherapy, and (3) the analytical and clinical validation criteria for approving a clinical assay (''diagnostic test''). The background material will become available no later than the day before the meeting and will be posted on FDA's Web site at *http://www.fda.gov/ohrms/dockets/ac/acmenu.htm* under the heading ''Advisory Committee for Pharmaceutical Science.'' (Click on the year 2005 and scroll down to the Advisory Committee for Pharmaceutical Science meetings.)

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 November 4, 2005. Oral presentations from the public will be scheduled between approximately 11:15 a.m. and 11:30 a.m. and 3:35 p.m. and 3:50 p.m. on November 14, 2005, and between approximately 11:20 a.m. and 11:50 a.m. on November 15, 2005. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 4, 2005, 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 Mimi Phan 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: October 18, 2005.

Jason Brodsky,

Acting Associate Commissioner for External Relations.

[FR Doc. 05–21241 Filed 10–24–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Infant Mortality; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: Advisory Committee on Infant Mortality (ACIM).

Dates and Times: November 29, 2005, 9 a.m.–5 p.m., November 30, 2005, 8:30 a.m.–3 p.m.

Place: Washington, DC area hotel yet to be determined.

Status: The meeting is open to the public with attendance limited to space availability.

Purpose: The Committee provides advice and recommendations to the Secretary of Health and Human Services on the following: Department programs that are directed at reducing infant mortality and improving the health status of pregnant women and infants; factors affecting the continuum of care with respect to maternal and child health care, including outcomes following childbirth; strategies to coordinate the variety of Federal, State, local and private programs and efforts that are designed to deal with the health and social problems impacting on infant mortality; and the implementation of the Healthy Start program and *Healthy People 2010* infant mortality objectives.

Agenda: Topics that will be discussed include the following: Health Disparities in the Maternal and Child Health (MCH) Population and A Life-course Perspective for Perinatal Health and Preconception Care. Substantial time will be spent in Subcommittee and full Committee discussions aimed at formulating the ACIM issues agenda. Proposed agenda items are subject to change as priorities indicate.

Time will be provided for public comments limited to five minutes each; comments are to be submitted no later than November 14, 2005.

For Further Information Contact: Anyone requiring information regarding the Committee should contact Peter C. van Dyck, M.D., M.P.H., Executive Secretary, ACIM, Health Resources and Services Administration (HRSA), Room 18–05, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857, Telephone: (301) 443–2170.

Individuals who are submitting public comments or who have questions regarding the meeting and location should contact Ann M. Koontz, C.N.M., Dr.P.H., HRSA, Maternal and Child Health Bureau, telephone: (301) 443–6327, e-mail: *ann.koontz@hrsa.hhs.gov.*

Dated: October 18, 2005.

Tina M. Cheatham,

Director, Division of Policy Review and Coordination.

[FR Doc. 05–21245 Filed 10–24–05; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Advisory Council on Nurse Education and Practice; Notice of Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), notice is hereby given of the following meeting:

Name: National Advisory Council on Nurse Education and Practice (NACNEP).

Dates and Times: November 2, 2005, 8:30 a.m.–5 p.m., November 3, 2005, 8:30 a.m.–5 p.m., November 4, 2005, 8:30 a.m.–3 p.m.

Place: DoubleTree Hotel and Executive Center, 1750 Rockville Pike, Rockville, Maryland 20852.

Status: The meeting will be open to the public.

Agenda: Agency and Bureau administrative updates will be provided. The purpose of the meeting will be to address issues related to the status of the nursing workforce. Two leading educators/researchers will present on their findings related to nursing workforce data, trends and issues. In addition, representatives from four nursing organizations representing nursing education, practice and licensing will present their data, trends, and workforce issues. During this meeting and the subsequent April 2006 meeting, Council workgroups will deliberate on content presented and formulate recommendations to the Secretary of Health and Human Services and the Congress on nursing workforce issues based on the latest data and trends. This meeting will form the basis for NACNEP's mandated Sixth Annual Report.

For Further Information Contact: Anyone interested in obtaining a roster of members, minutes of the meeting, or other relevant information should write or contact Ms. Donna English, M.P.H., R.N., Executive Secretary, National Advisory Council on Nurse Education and Practice, Parklawn Building, Room 9–35, 5600 Fishers Lane, Rockville, Maryland 20857, telephone (301) 443–5688.

Dated: October 19, 2005.

Tina M. Cheatham,

Director, Division of Policy Review and Coordination.

[FR Doc. 05–21243 Filed 10–24–05; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Advisory Committee on Organ Transplantation; Notice of Meeting

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice; amendment.

SUMMARY: The Health Resources and Services Administration published a notice in the **Federal Register** of September 29, 2005 (70 FR 56926–56927) announcing an Advisory Committee on Organ Transplantation meeting on November 3–4 2005. The notice is being amended to change type of meeting, place, dates, and time.

FOR FURTHER INFORMATION CONTACT: Ms. Sherry Whipple at 301–443–2764 or e-mail: *swhipple@hrsa.gov.*

Correction

In the **Federal Register** of September 29, 2005, in FR Doc. 05–19431, on page 56927, 1ST column, under the heading Summary, 8th line, change to read:

The meeting will be an Audio Conference Call on November 1, 2005, from 12 noon to 4 p.m. EST. To access the conference call, call the USA Toll Free Number 888–791–2132 and enter the Passcode ''ACOT.'' The conference call leader is Dr. James Burdick.

Dated: October 18, 2005.

Tina M. Cheatham,

Director, Division of Policy Review and Coordination.

[FR Doc. 05–21244 Filed 10–24–05; 8:45 am]

BILLING CODE 4165–15–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

[USCG–2005–21722]

Collection of Information Under Review by Office of Management and Budget (OMB): 1625–0089

AGENCY: Coast Guard, DHS.

ACTION: Request for comments.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, this request for comments announces that the Coast Guard has forwarded one Information Collection Request (ICR)—(1) 1625–0089, The National Recreational Boating Survey—abstracted below, to the Office of Information and Regulatory Affairs (OIRA) of the Office of Management and Budget (OMB) for review and comment. Our ICR describes the information we seek to collect from the public. Review and comment by OIRA ensures that we impose only paperwork burdens commensurate with our performance of duties.

DATES: Please submit comments on or before November 25, 2005.

ADDRESSES: To make sure that your comments and related material do not reach the docket [USCG–2005–21722] or OIRA more than once, please submit them by only one of the following means:

(1)(a) By mail to the Docket Management Facility, U.S. Department of Transportation (DOT), room PL–401, 400 Seventh Street, SW., Washington, DC 20590–0001. (b) By mail to OIRA, 725 17th St. NW., Washington, DC 20503, to the attention of the Desk Officer for the Coast Guard.

(2)(a) By delivery to room PL–401 at the address given in paragraph (1)(a)