Authority of New York and New Jersey, NY and NJ, Comment Period Ends: July 21, 2004, Contact: Bernard Cohen (212) 668–1770. The document is available on the Internet at: http://www.panynj.gov/pathrestoration.

EÍS No. 040258, Final EIS, UAF, Air Force Mission at Johnston Atoll Airfield (Installation) Termination, Implementation, Johnston Atoll is an Unincorporated Territory of the United States, Wait Period Ends: July 6, 2004, Contact: Patricia Vokoun (703) 604–5263.

EIS No. 040259, Final EIS, NOA, AK, Programmatic EIS—Alaska Groundfish Fisheries, New Information concerning the Ecosystem and a Preferred Alternative, Fishery Management Plans for the Groundfish Fishery of the Gulf of Alaska and the Groundfish of the Bering Sea and Aleutian Islands Area, North Pacific Fishery Management Council, AK, Wait Period Ends: July 6, 2004, Contact: James W. Balsiger (907) 586–7221. This document is available on the Internet at: http://www.fakr.noaa.gov.

EIS No. 0240260, Final EIS, BLM, WY, Desolation Flats Natural Gas Field Development Project, Drilling Additional Development Wells, Carbon and Sweetwater Counties, WY, Wait Period Ends: July 6, 2004, Contact: David Simons (307) 367—5309. This document is available on the Internet at: http://www.wy.blm.gov/nepadocs.htm.

Amended Notices

EIS No. 040027, Draft EIS, IBR, NE, CO, WY, Programmatic EIS—Platte River Recovery Implementation Program, Assessing Alternatives, Cooperative, Endangered Species Recovery Program, The Four Target Species are Whooping Crane, Interior Least Tern, Piping Plover and Pallid Sturgeon, NE, WY and CO, Comment Period Ends: August 20, 2004, Contact: Curt Brown (303) 445–2096. Revision of FR Notice Published on 1/30/2004: CEQ Comment Period Ending on 06/2/2004 has been Extended to 8/20/2004.

EIS No. 040222, Draft EIS, DOE, CA, Imperial-Mexicali 230-kV Transmission Lines, Construct a Double-Circuit 230-kV Transmission Line, Presidential Permit and Rightof-Way Grants, Imperial Valley Substation to Calexico at the U.S.-Mexico Border, Imperial County, CA and U.S.-Mexico Border, Comment Period Ends: July 30, 2004, Contact: Ellen Russell (202) 586–7624. Revision of FR Notice Published on 5/ 14/2004: CEQ Comment Period Ending 6/30/2004 has been Extended to 7/30/2004.

EIS No. 040233, Revised Final EIS, BLM, NM, Sierra and Otero Counties Resource Management Plan Amendment and Federal Fluid Minerals Leasing and Development, Additional Information to Improve the Public Understanding of the Proposed Plan, Implementation, Sierra and Otero Counties, NM, Wait Period Ends: June 23, 2004, Contact Tom Phillips (505) 525–4377. Revision of FR Notice Published on 5/21/2004: Correction to Contact Person Telephone Number.

Dated: June 1, 2004.

Ken Mittelholtz,

Environmental Protection Specialist, Office of Federal Activities.

[FR Doc. 04–12705 Filed 6–3–04; 8:45 am]

FEDERAL MARITIME COMMISSION

Sunshine Act Meeting

TIME AND DATE: 10 a.m.—June 9, 2004. PLACE: 800 North Capitol Street, NW., First Floor Hearing Room, Washington, DC.

STATUS: Closed.

MATTERS TO BE CONSIDERED: 1.

Licensing, Compliance, Enforcement Process and SHJ International Express, LLC and Gary Yenkok Tan.

CONTACT PERSON FOR MORE INFORMATION: Bryant L. VanBrakle, Secretary, (202) 523–5725.

Bryant L. VanBrakle,

Secretary.

[FR Doc. 04–12817 Filed 6–2–04; 1:56 pm]
BILLING CODE 6730–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel: Occupational Health and Safety Research, Program Announcement Number 04038

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following meeting:

Name: Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Occupational Health and Safety Research, Program Announcement Number 04038. Times and Dates: 2 p.m.-2:15 p.m., June 23, 2004 (Open), 2:15 p.m.-3:30 p.m., June 23, 2004 (Closed).

Place: Teleconference Phone Number 1.888.283.3870 Pass Code 11026.

Status: Portions of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement Number 04038.

For Further Information Contact: Bernadine B. Kuchinski, Ph.D., Occupational Health Consultant, National Institute for Occupational Safety and Health, CDC, 4676 Columbia Parkway MS C7, Cincinnati, OH 45226, Telephone 513.533.8511.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: May 27, 2004.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04–12676 Filed 6–3–04; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect (NTFFASFAE).

Times and Dates: 9 a.m.—4:30 p.m., June 16, 2004. 9 a.m.—12 noon, June 17, 2004.

Place: Doubletree Hotel Atlanta/Buckhead, 3342 Peachtree Road, NE., Atlanta, Georgia 30326, telephone 404/231–1234, fax 404/231–3112.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 65 people.

Purpose: The Secretary is authorized by the Public Health Service Act, Section 399G, (42 U.S.C. Section 280f, as added by Public Law 105–392) to establish a National Task Force on Fetal Alcohol Syndrome and Fetal Alcohol Effect to:

- (1) Foster coordination among all governmental agencies, academic bodies and community groups that conduct or support Fetal Alcohol Syndrome (FAS) and Fetal Alcohol Effect (FAE) research, programs and surveillance; and
- (2) To otherwise meet the general needs of populations actually or potentially impacted by FAS and FAE.

Matters to be Discussed: Agenda items include: a review of the September 2002 Task Force Recommendations and the activities undertaken by Federal and non-governmental agencies and organizations in response to the recommendations; the identification of priority areas for the current Task Force; and the reconvening of the Research working group and the Services and Public Awareness working group. Additional agenda items include: updates from Task Force members on current initiatives; an update on activities from the Interagency Coordinating Committee on Fetal Alcohol Syndrome, the CDC and other Federal agencies; reports from Task Force liaison organizations; future topics, and scheduling the next meeting.

Agenda items are subject to change as priorities dictate.

Due to programmatic issues that had to be resolved, the **Federal Register** notice is being published less than fifteen days before the date of the meeting.

For Further Information Contact: R. Louise Floyd, DSN, RN, Designated Federal Official, National Center on Birth Defects and Developmental Disabilities, CDC, 1600 Clifton Road, NE, (E–86), Atlanta, Georgia 30333, telephone 404/498–3923, fax 404/498–3550.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities for both the CDC and ATSDR.

Dated: May 28, 2004.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04–12675 Filed 6–3–04; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Advisory Committee on Immunization Practices

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following committee meeting.

Name: Advisory Committee on Immunization Practices (ACIP). Times and Dates: 9 am—5 pm, June 23, 2004. 8 am—5 pm, June 24, 2004. Place: Atlanta Marriott Century Center, 2000 Century Boulevard, NE., Atlanta, Georgia 30345–3377.

Status: Open to the public, limited only by the space available.

Purpose: The Committee is charged with advising the Director, CDC, on the appropriate uses of immunizing agents. In addition, under 42 U.S.C. § 1396s, the Committee is mandated to establish and periodically review and, as appropriate, revise the list of vaccines for administration to vaccine-eligible children through the Vaccines for Children (VFC) program, along with schedules regarding the appropriate periodicity, dosage, and contraindications applicable to the vaccines.

Matters to be Discussed: The Agenda will include discussions on influenza; IOM report on autism and vaccines; an update on Hepatitis A vaccine; recommended childhood and adolescent immunization schedules; PCV7 shortage; discussion on meningococcal conjugate vaccine; smallpox pregnancy registry outcomes; pertussis; working group and departmental updates.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Demetria Gardner, Epidemiology and Surveillance Division, National Immunization Program, CDC, 1600 Clifton Road, NE, (E–61), Atlanta, Georgia 30333, telephone 404/639–8096, fax 404/639–8616.

The Director, Management Analysis and Services Office has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both CDC and the Agency for Toxic Substances and Disease Registry.

Dated: May 27, 2004.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04–12677 Filed 6–3–04; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2003N-0360]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Information Program on Clinical Trials for Serious and Life-Threatening Diseases: Maintaining a Databank

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Information Program on Clinical Trials for Serious and Life-Threatening Diseases: Maintaining a Databank" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT:

Jonna Capezzuto, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the Federal Register of February 19, 2004 (69 FR 7753), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0459. The approval expires on May 31, 2007. A copy of the supporting statement for this information collection is available on the Internet at http://www.fda.gov/ ohrms/dockets.

Dated: March 27, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.
[FR Doc. 04–12684 Filed 6–3–04; 8:45 am]
BILLING CODE 4160–01–8

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2004N-0244]

Agency Information Collection Activities; Proposed Collection; Comment Request; Current Good Manufacturing Practice Regulations for Type A Medicated Articles

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and allow 60 days for public comment in response to the notice. This notice solicits comments on the recordkeeping requirements for manufacturers of type A medicated articles.