DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Toxicology Program

Notice of Establishment

Pursuant to the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), the Director, National Toxicology Program (NTP), announces the establishment of the Advisory Committee on Alternative Toxicological Methods by the Secretary, DHHS.

The Committee will advise the NIEHS Interagency Center for the Evaluation of Alternative Toxicological Methods and the Director of the Environmental Toxicology Program on the activities and directives both present and future, as they relate to the Center, including advice on fostering interactions with all stakehoders.

Duration of this Committee is continuing unless formally determined by the Secretary, DHHS, that termination would be in the best public interest.

Dated: September 4, 1997.

Kenneth Olden,

Director, National Toxicology Program.

[FR Doc. 97–24273 Filed 9–12–97; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[ATSDR–122]

Quarterly Public Health Assessments Completed

AGENCY: Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: This notice is a quarterly announcement of sites for which ATSDR has completed public health assessments during the period April–June 1997. This list includes sites that are on or proposed for inclusion on the National Priorities List (NPL), and a site for which an assessment was prepared in response to a request from the public.

FOR FURTHER INFORMATION CONTACT: Robert C. Williams, P.E., DEE, Director, Division of Health Assessment and Consultation, Agency for Toxic Substances and Disease Registry, 1600 Clifton Road, NE., Mailstop E–32, Atlanta, Georgia 30333, telephone (404) 639–0610.

SUPPLEMENTARY INFORMATION: The most recent list of completed public health assessments was published in the **Federal Register** on June 10, l997, [62 FR 31603]. The quarterly announcement is the responsibility of ATSDR under the regulation, Public Health Assessments and Health Effects Studies of Hazardous Substances Releases and Facilities [42 CFR Part 90]. This rule sets forth ATSDR's procedures for the conduct of public health assessments under section 104(i) of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA), as amended by the Superfund Amendments and Reauthorization Act (SARA) [42 U.S.C. 9604(i)].

Availability—The completed public health assessments and addendum are available for public inspection at the Division of Health Assessment and Consultation, Agency for Toxic Substances and Disease Registry, Building 33, Executive Park Drive, Atlanta, Georgia (not a mailing address), between 8 a.m. and 4:30 p.m., Monday through Friday except legal holidays. The completed public health assessments are also available by mail through the U.S. Department of Commerce, National Technical Information Service (NTIS), 5285 Port Royal Road, Springfield, Virginia 22161, or by telephone at (703) 487–4650. NTIS charges for copies of public health assessments and addenda. The NTIS order numbers are listed in parentheses following the site names.

Public Health Assessments Completed or Issued

Between April 1, 1997 and June 1, 1997, public health assessments were issued for the sites listed below:

NPL Sites

Idaho
- Triumph Mine Tailings Piles—Hailey—(PB97–171722)

Illinois
- Dupage County Landfill (Blackwell Forest Preserve)—Warrenville—(PB97–162598)
- Woodstock Municipal Landfill—Woodstock—(PB97–161459)

Missouri
- Weldon Spring Quarry/Plant/Pits (USDOE)—St. Charles—(PB97–180509)

New York
- Pollution Abatement Services—Oswego—(PB97–171748)

Ohio
- Miami County Incinerator—Troy—(PB97–157770)

Oklahoma
- Kerr-McGee Corporate Cushing Refinery—Cushing—(PB97–155204)

Pennsylvania
- Breslube-Penn Incorporated—Coraopolis—(PB97–164891)
- USA Tobyhanna Army Depot—Coolbaugh Township—(PB97–161491)

Tennessee
- ICG Iselin Railroad Yard—Jackson—(PB97–159511)

Non NPL Petition Site

Connecticut
- Connecticut Correctional Institution (a/k/a Somers Correctional Facility)—Somers—(PB97–155212)

Dated: September 9, 1997.

Georgi Jones,

Director, Office of Policy and External Affairs Agency for Toxic Substances and Disease Registry.

[FR Doc. 97–24336 Filed 9–12–97; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Cooperative Agreements to Conduct Research on the Diagnosis and Pathogenesis of Lyme Disease in the United States, Program Announcement 800: 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: Disease, Disability, and Injury Prevention and Control SEP: Cooperative Agreements to Conduct Research on the Diagnosis and Pathogenesis of Lyme Disease in the United States, Program Announcement 800.

Time and Dates: 8:30 a.m.–4:30 p.m., October 6–7, 1997.

Place: Holiday Inn Conference Center, 130 Clairemont Avenue, Decatur, Georgia 30030.

Status: Open: 8:30 a.m.–9:15 a.m., October 6, 1997. Closed: 9:15 a.m. October 6, 1997, through 4:30 p.m. October 7, 1997.

Matters to be Discussed: The meeting will include the review, discussion, and evaluation of applications received in response to Program Announcement 800.

Portions of this 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 Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463.

Contact Person For More Information: Edwarda Lee, M.P.A., Division of Vector-Borne Infectious Diseases, National Center

for Infectious Diseases, CDC, M/S P02/MLR, P.O. Box 2087 (Foothills Campus), Fort Collins, Colorado 80522, telephone 970/221–6415.

Dated: September 8, 1997.

Carolyn J. Russell,

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

[FR Doc. 97–24335 Filed 9–12–97; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: Child Care Quarterly Case Record Report; Republication, In **Federal Register** Document 97–23469 (Volume 62, Number 171), Page 46743 the word ''Disaggregate'' replaces the word ''Desegregate'' and ''Respondents: States and Territories'' replaces ''Respondents: State, Local or Tribal Govt.'' For the convenience of the reader, the document is being republished in its entirety.

OMB No.: New Request.

Description: This legislatively-mandated report collects program and participants data on children receiving direct CCDF funds. Disaggregate data will be collected and will be used to determine the participants and program characteristics, as well as cost and level of child care services. The data will be used to provide a report to Congress.

Respondents: States and Territories.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF–801	56	4	20	4,360

Estimated Total Annual Burden Hours: 4,360.

Additional Information: Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register.** Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Laura Oliven.

Dated: September 9, 1997.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 97–24280 Filed 9–12–97; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N–0022]

Agency Information Collection Activities Submission for OMB Review; Comment Request

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (the PRA).

DATES: Submit written comments on the collection of information by October 15, 1997.

ADDRESSES: Submit written comments on the collection of information to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW., rm. 10235, Washington, DC 20503, Attention: Desk Officer for FDA.

FOR FURTHER INFORMATION CONTACT: Judith V. Bigelow, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1479.

SUPPLEMENTARY INFORMATION: In compliance with section 3507 of the PRA (44 U.S.C. 3507), FDA has submitted the following proposed collection of information to OMB for review and clearance.

Hearing Aid Devices: Professional and Patient Package Labeling and Conditions for Sale—21 CFR 801.420 and 801.421 (OMB Control No. 0910–0171—Reinstatement)

Under section 520(e) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360j(e)), the Secretary of the Department of Health and Human Services (the Secretary) may, under certain conditions, require by regulation that a device be restricted to sale, distribution, or use only upon authorization of a licensed practitioner or upon other prescribed conditions. Sections 801.420 and 801.421 (21 CFR 801.420 and 801.421) implement this authority for hearing aids, which are restricted devices. The regulations require that the manufacturer or distributor provide to the user data useful in selecting, fitting, and checking the performance of a hearing aid through distribution of a User Instructional Brochure. The User Instructional Brochure must also contain technical data about the device, instructions for its use, maintenance and care, a warning statement, a notice about the medical evaluation requirement, and a statement if the aid is rebuilt or used.

Hearing aid dispensers are required to provide the prospective user, before the sale of a hearing aid, with a copy of the User Instructional Brochure for the hearing aid model that has been, or may be, selected for the prospective user and to review the contents of the brochure with the buyer. In addition, upon