disease prevention and health promotion goals, and the feasibility of the project in terms of the operational plan.

3. The Qualifications of Program Personnel (20 Points)

a. The extent to which the application provides evidence of the qualifications, experience, and commitment of the principal staff person, and his/her ability to devote adequate time and effort to provide effective leadership.

b. The extent to which the application provides evidence of the competence of associate staff persons, discussion leaders, speakers, and presenters to accomplish conference objectives.

c. The extent to which the application demonstrates the knowledge of nationwide and educational efforts currently underway which may affect, and be affected by, the proposed conference.

4. Evaluation Methods (20 Points)

Evaluation instrument(s) for the conference should adequately assess increased knowledge, attitudes, and behaviors of the target audience.

5. Applicant's Capability (10 Points)

a. The applicant's capability includes the adequacy of the applicant's resources (additional sources of funding, organization's strengths, staff time, proposed physical facilities, *etc.*) available for conducting conference activities.

b. The extent to which the applicant demonstrates a history (at least three years) of managing conferences.

6. Budget Justification and Adequacy of Facilities (Not Scored)

The proposed budget will be evaluated on the basis of its reasonableness, concise and clear justification, and consistent with the intended use of grant funds. The application will also be reviewed as to the adequacy of existing or proposed facilities and resources for conducting conference activities.

J. Other Requirements

Technical Reporting Requirements: Provide the CDC with original plus two copies of:

1. A performance report or, in lieu of a performance report, proceedings of the conference, no later than 90 days after the end of the budget/project period.

2. Financial status report, no later than 90 days after the end of the budget/project period.

Send all reports to the Grants Management Specialist identified in the ''Where to Obtain Additional Information'' section of this announcement.

Additional Requirements: The following additional requirements are applicable to this program. For a complete description of each, see Attachment I of this announcement as posted on the CDC Web site.

AR–7—Executive Order 12372 Review.

AR–9—Paperwork Reduction Act Requirements.

AR–10—Smoke-Free Workplace Requirements.

AR–11—Healthy People 2010.

AR–12—Lobbying Restrictions.

AR–13—Prohibition on Use of CDC Funds for Certain Gun Control Activities.

AR–15—Proof of Non-Profit Status.

AR–20—Conference Support.

K. Where To Obtain Additional Information

This and other CDC announcements, the necessary applications and associated forms can be found on the CDC Web site, Internet address: *http://www.cdc.gov.* Click on ''Funding'' then ''Grants and Cooperative Agreements.''

For general questions about this announcement, contact: Technical Information Management, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341–4146, Telephone: (770) 488–2700.

For business management assistance, contact: Rick Jaeger, Grants Management Specialist, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 2920 Brandywine Road, Atlanta, Georgia 30341–4146, Telephone: (770) 488–2727, e-mail address: *rjaeger@cdc.gov.*

For program technical assistance, contact: Janet Telman, Funding Resource Specialist, Office of the Director Extramural Services Activity, Public Health Practice Program Office (PHPPO), Centers for Disease Control and Prevention (CDC), 4770 Buford Highway, NE, MS K38, Atlanta, Georgia 30341–3714, Telephone: (770) 488–2834, e-mail address: *jtelman@cdc.gov.*

Dated: August 21, 2003.

Sandra R. Manning,

Director, Program and Grants Office, Centers for Disease Control and Prevention.

[FR Doc. 03–22007 Filed 8–27–03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

[ATSDR–195]

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 announces those sites for which ATSDR has completed public health assessments during the period from April 2003 through June 2003. This list includes sites that are on or proposed for inclusion on the National Priorities List (NPL), and includes sites for which assessments were prepared in response to requests from the public.

FOR FURTHER INFORMATION CONTACT: Robert C. Williams, P.E., DEE, Assistant Surgeon General, 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) 498–0007.

SUPPLEMENTARY INFORMATION: The most recent list of completed public health assessments was published in the **Federal Register** on August 8, 2003 [68 FR 47324]. This 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 addenda are available for public inspection at the Division of Health Assessment and Consultation, Agency for Toxic Substances and Disease Registry, Building 1825, Century Blvd., 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) 605–6000. 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, 2003 and June 30, 2003, public health assessments were issued for the sites listed below:

NPL Sites

California

Edwards Air Force Base (PB2003–104567)

Markleeville (PB2003–104568)

Illinois

Circle Smelting Corporation (PB2003–104617)

Hartford Residential Vapor Resources (a/k/a Hartford Residences) (PB2003–105779)

Maine

Callahan Mining Corporation (PB2003–104534)

Massachusetts

General Electric Site—Unkamet Brook (a/k/a GE-Housatonic River) (PB2003–104653)

General Electric Site—Lyman Street (a/k/a GE-Housatonic River) (PB2003–104654)

General Electric Site—Former Oxbows (a/k/a GE-Housatonic River) (PB2003–104655)

Sutton Brook Disposal Area (PB2003–104569)

New York

Cayuga County Groundwater Contamination (PB2003–105778)

Petitioned

Florida

Eglin Air Force Base (a/k/a USAF Eglin Air Force Base Armament Division) (PB2003–104185)

Dated: August 22, 2003.

Georgi Jones,

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

[FR Doc. 03–22002 Filed 8–27–03; 8:45 am]

BILLING CODE 4163–70–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee (CLIAC)

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: Clinical Laboratory Improvement Advisory Committee (CLIAC).

Times and Dates: 8:30 a.m.–5 p.m.; September 17, 2003. 8:30 a.m.–3:30 p.m.; September 18, 2003.

Place: Sheraton Colony Square Hotel, 188 14th Street NE., Atlanta, Georgia 30361.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people.

Purpose: This committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the need for, and the nature of, revisions to the standards under which clinical laboratories are regulated; the impact on medical and laboratory practice of proposed revisions to the standards; and the modification of the standards to accommodate technological advances.

Matters To Be Discussed: The agenda will include updates from CDC, the Centers for Medicare & Medicaid Services, and the Food and Drug Administration; a report on the results of the General Services Administration's Office of Government-wide Policy Federal Advisory Committee Stakeholder Engagement Survey; presentations and discussion on the CLIA waiver criteria and process, previous CLIAC recommendations related to such, and AdvaMed's CLIAC waiver criteria proposal; a report on the Coordinating Council for Clinical Laboratory Workforce's June 2003 meeting; a report on the April 2003 Quality Institute; a summary of the March 2003 CLIAC meeting on direct access testing; a presentation on Lab Tests Online; a report on the first meeting of the Secretary's Advisory Committee on Genetics, Health and Society; and several presentations on CDC's various genetic testing activities.

Agenda items are subject to change as priorities dictate.

Providing Oral or Written Comments: It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments whenever possible. *Oral Comments:* In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes (unless otherwise indicated). Speakers must also submit their comments in writing for inclusion in the meetings Summary Report. *Written Comments:* For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated). However, the comments should be received at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments, one hard copy with original signature, should be provided to the contact person below. Written comments will be included in the meetings Summary Report.

Contact Person for Additional Information: Rhonda Whalen, Chief, Laboratory Practice Standards Branch, Division of Laboratory Systems, Public Health Practice Program Office, CDC, 4770 Buford Highway, NE., Mailstop F–11, Atlanta, Georgia 30341–3717; telephone (770) 488–8042; fax (770) 488–8279; or via e-mail at *RWhalen@cdc.gov.*

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: August 20, 2003.

Alvin Hall,

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

[FR Doc. 03–22008 Filed 8–27–03; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Information Relevant to Toluene Exposure

AGENCY: National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).