Grants Office (PGO) staff and for responsiveness by ATSDR. Incomplete applications and applications that are non-responsive to the eligibility criteria will not advance through the review process. Applicants will be notified that their application did not meet submission requirements.

An objective review panel will evaluate your application according to the criteria listed section ''V.1. Criteria'' section above.

In addition, the following factor may affect the funding decision: Ability to provide site-specific educational consultation on environmental medicine and pediatric health concerns in locations such as superfund sites where ATSDR or the EPA is assisting communities to cope with hazardous contamination.

*V.3. Anticipated Announcement Award Date*

August 1, 2004

**VI. Award Administration Information**

*VI.1. Award Notices*

Successful applicants will receive a Notice of Grant Award (NGA) from the CDC PGO. The NGA shall be the only binding, authorizing document between the recipient and CDC. The NGA will be signed by an authorized Grants Management Officer, and mailed to the recipient fiscal officer identified in the application.

Unsuccessful applicants will receive notification of the results of the application review by mail.

*VI.2. Administrative and National Policy Requirements*

45 CFR Parts 74 and Part 92.

For more information on the Code of Federal Regulations, *see* the National Archives and Records Administration at the following Internet address: *http://www.access.gpo.gov/nara/cfr/cfr-table-search.html.*

The following additional requirements apply to this project:

- AR–1 Human Subjects Requirements
- AR–2 Requirements for Inclusion of Women and Racial and Ethnic Minorities in Research
- AR–8 Public Health System Reporting Requirements
- AR–9 Paperwork Reduction Act Requirements
- AR–10 Smoke-Free Workplace Requirements
- AR–11 Healthy People 2010
- AR–12 Lobbying Restrictions
- AR–14 Accounting System Requirements
- AR–18 Cost Recovery-ATSDR
- AR–19 Third Party Agreements-ATSDR

Additional information on these requirements can be found on the CDC Web site at the following Internet address: *http://www.cdc.gov/od/pgo/funding/ARs.htm.*

*VI.3. Reporting Requirements*

You must provide CDC with an original, plus two copies of the following reports:

1. Interim progress report, no less than 90 days before the end of the budget period. The progress report will serve as your non-competing continuation application, and must contain the following elements:

a. Current Budget Period Activities Objectives.

b. Current Budget Period Financial Progress.

c. New Budget Period Program Proposed Activity Objectives.

d. Detailed Line-Item Budget and Justification.

e. Additional Requested Information.

f. Measures of Effectiveness.

2. Financial status report and annual progress report, no more than 90 days after the end of the budget period.

3. Final financial and performance reports, no more than 90 days after the end of the project period.

These reports must be sent to the Grants Management Specialist listed in the ''Agency Contacts'' section of this announcement.

**VII. Agency Contacts**

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

For program technical assistance, contact:

Robert H. Johnson, MD, Medical Officer, Division of Health Education and Promotion, 1600 Clifton Road, N.E., Mailstop E–33, Atlanta, GA 30333, Telephone: (404) 498–0498, e-mail: *rdj2@cdc.gov.*

For budget assistance, contact: Edna Green, Grants Management Specialist, CDC Procurement and Grants Office, 2920 Brandywine Road, Atlanta, GA 30341, Telephone: (770) 488–2743, e-mail: *ecg4@cdc.gov.*

Dated: June 4, 2004.

**William P. Nichols,**

*Acting Director, Procurement and Grants Office, Centers for Disease Control and Prevention.*

[FR Doc. 04–13140 Filed 6–9–04; 8:45 am]

**BILLING CODE 4163–70–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### Breast and Cervical Cancer Early Detection and Control Advisory Committee

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:* Breast and Cervical Cancer Early Detection and Control Advisory Committee (BCCEDCAC).

*Times and Dates:* 8 a.m.–5 p.m., June 23, 2004. 8 a.m.–3:30 p.m., June 24, 2004.

*Place:* Hyatt Regency New Orleans, Poydras Plaza at Loyola Ave., New Orleans, Louisiana, 70113–1805. Phone: 1–504–561–1234.

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

*Purpose:* The committee is charged with advising the Secretary, Department of Health and Human Services, and the Director, CDC, regarding the early detection and control of breast and cervical cancer. The committee makes recommendations regarding national program goals and objectives; implementation strategies; and program priorities including surveillance, epidemiologic investigations, education and training, information dissemination, professional interactions and collaborations, and policy.

*Matters To Be Discussed:* The agenda will include discussion and review of National Breast and Cervical Early Detection Program (NBCCEDP) Programmatic issues related to the NBCCEDP Manual review/update, IMS (Information Management Services) update, Cervical cancer policy and new technologies, recruitment issues, Models of cancer registry, New mammography and CAD, Breast and Cervical issues, and Clinical Breast Exams issues; Comprehensive and Integrated Approaches Cancer Control; Health disparities within NBCCEDP; Building Better Partnerships; and discussion with NBCCEDP Program Directors related to implementation of the National Breast and Cervical Program.

Agenda items are subject to change as priorities dictate.

*Contact Person for More Information:* Debra Younginer, Executive Secretary, BCCEDCAC, Division of Cancer Prevention and Control, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway, Mailstop K–57, Chamblee, Georgia 30316, telephone: 770–488–1074.

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: June 4, 2004.

**Alvin Hall,**

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

[FR Doc. 04–13133 Filed 6–9–04; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Disease Control and Prevention

### National Institute for Occupational Safety and Health; Public Meeting

The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) announces the following public meeting and request for information:

*Name:* Public Meeting to Seek Input on Gaps in Chronic Lymphocytic Leukemia Radiogenicity Research.

*Time and Date:* 9 a.m.–12 noon, July 21, 2004.

*Place:* Best Western Skyline Inn, 10 I Street, SW., Washington, DC 20024.

*Status:* Forum will include scientists and representatives from various government agencies, industry, labor, and other stakeholders, and is open to the public, limited only by the space available. The meeting room accommodates up to 100 people. Due to limited space, notification of intent to attend the meeting should be made with Patty Gudlewski, no later than Friday, July 16, 2004. Ms. Gudlewski can be reached by telephone at 513–841–4419, or by e-mail at *pkg1@cdc.gov*. Access to the meeting will be accommodated on a first-come basis.

*Purpose:* To discuss possible scientific research strategies to evaluate any relationship between exposure to ionizing radiation and chronic lymphocytic leukemia (CLL). Current scientific opinion, based largely on epidemiological data, holds that the incidence of CLL is not related to exposure to ionizing radiation. The U.S. Congress directed NIOSH to conduct epidemiological research and other activities to establish the scientific link between radiation exposure and the occurrence of CLL.

The public is invited to attend and will have an opportunity to provide limited comments. Written comments may be submitted to the address listed below by August 16, 2004, so that they may be considered by NIOSH in planning its research priorities.

*Summary:* CLL is the most common adult leukemia in the Western world, but its etiology is largely unknown. Exposures to some herbicides have been implicated in epidemiologic studies. Yet other studies to date largely have shown no evidence of an association between external ionizing radiation and CLL; however, a number of uncertainties remain and additional studies may be informative. Recent laboratory studies have identified sub-types of CLL and at least one familial form of B-cell CLL has been identified. In addition, new technologies including interphase fluorescence *in situ* hybridization, expression microarrays and flow cytometric analysis provide diagnostic and prognostic indicators of disease. This meeting will assist in identifying gaps in existing research needed to address the radiogenicity of CLL.

*Addresses:* Comments should be submitted to David F. Utterback, 4676 Columbia Parkway, M/S R–44, Cincinnati, Ohio 45226, or by e-mail to *dutterback@cdc.gov*. Any attachments should be formatted in Microsoft Word.

All information received in response to this notice will be available for public examination and copying.

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: June 4, 2004.

**Alvin Hall,**

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

[FR Doc. 04–13134 Filed 6–9–04; 8:45 am]

**BILLING CODE 4163–19–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Centers for Medicare & Medicaid Services

### Privacy Act of 1974; Report of New System

**AGENCY:** Department of Health and Human Services (HHS), Centers for Medicare & Medicaid Services (CMS).

**ACTION:** Notice of new system of records (SOR).

**SUMMARY:** In accordance with the requirements of the Privacy Act of 1974, we are proposing to establish a new SOR, titled ''MMA Section 641 Prescription Drug Benefit Demonstration'' (MMA641) System NO. 09–70–0545, HHS/CMS/ORDI. The primary purposes of the system of records are to maintain information on individual Medicare beneficiaries who voluntarily enroll in a demonstration project for coverage of certain prescription drugs and biologicals. This demonstration project is mandated in the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 under section 641. The system of records will enable CMS to: Enroll and communicate with eligible Medicare beneficiaries who volunteer to participate in the demonstration project, communicate with clinicians and other providers and suppliers who submit claims payable under the demonstration project, review submitted claims and pay those conforming to applicable payment criteria and federal law, and develop, maintain, and analyze research information showing the potential impact of providing certain prescription drugs and biologicals.

Information retrieved from this system of records will also be disclosed to support regulatory, reimbursement, and policy functions performed within the agency or by a contractor or consultant; support constituent requests made to a Congressional representative; support litigation involving the agency; support activities reasonably necessary to fulfill the provisions of the demonstration project and ensure appropriate use of Medicare trust fund and program funds; and third parties where the contact is expected to have information relating to the individual's capacity to manage his or her own affairs.

We have provided background information about the proposed system in the ''Supplementary Information'' section, below. CMS invites comments on all portions of this notice. See ''Effective Dates'' section for comment period.

**EFFECTIVE DATES:** CMS filed a new system report with the Chair of the House Committee on Government Reform and Oversight, the Chair of the Senate Committee on Governmental Affairs, and the Administrator, Office of Information and Regulatory Affairs, Office of Management and Budget (OMB) on June 4, 2004. In any event, we will not disclose any information under a routine use until forty (40) calendar days after publication. We may defer implementation of this system of records or one or more of the routine use statements listed below if we receive comments that persuade us to defer implementation.

**ADDRESSES:** The public should address comments to: Director, Division of Privacy Compliance Data Development (DPCDD), CMS, Room N2–04–27, 7500 Security Boulevard, Baltimore, Maryland 21244–1850. Comments received will be available for review at this location, by appointment, during regular business hours, Monday through Friday from 9 a.m.–3 p.m., eastern time zone.

**FOR FURTHER INFORMATION CONTACT:** James Coan, Division of Health Promotion and Disease Prevention Demonstrations (DHPDPD), Office of Research, Development, and Information, CMS, MS–S3–02–01, 7500 Security Boulevard, Baltimore,