Event	Number of hours/costs per event and labor category* (per respondent)	Approx. number of respondents**	Approx. annual hours (millions)	Approx. total costs (millions)
Total			1.70	31.75

* Staff calculated labor costs by applying appropriate hourly cost figures to burden hours. The hourly rates used were $50 for managerial/professional time (e.g., compliance evaluation and/or planning), $20 for skilled technical time (e.g., designing and producing notices, reviewing and updating information systems), and $10 for clerical time (e.g., reproduction tasks, filing, and, where applicable to the given event, typing or mailing). Consumers have a continuing right to opt-out, as well as a right to revoke their opt-out at any time. When a respondent changes its information sharing practices, consumers are again given the opportunity to opt-out. Again, staff assumes that the time required of consumer to respond affirmatively to respondent's opt-out program (be it manually or electronically) would be minimal.

** The estimate of respondents is based on the following assumptions: (1) 100,000 respondents, approximately 70% of whom maintain customer relationships exceeding one year (2) no more than 1% (1,000) of whom make additional changes to privacy policies at any time other than the occasion of the annual notice; and (3) such changes will occur no more often than once per year.

As calculated above, the average PRA burden for all affected entities in a given year would be 1,000,000 hours and $19,875,000.

Estimated Capital/Other Non-Labor Costs Burden: Staff estimates that the capital or other non-labor costs associated with the document requests are minimal. Covered entities will already be equipped to provide written notices (e.g., computers with word processing programs, typewriters, copying machines, mailing capabilities.) Most likely, only entities that already have on-line capabilities will offer consumers the choice to receive notices via electronic format. As such, these entities will already be equipped with the computer equipment and software necessary to disseminate the required disclosures via electronic means.

William E. Kovacic,

General Counsel.

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BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality; Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Health Care Policy and Research Special Emphasis Panel (SEP) meeting.

The Health Care Policy and Research Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and quality (AHRQ), and agree to be available, to conduct, or an as needed basis, scientific review or applications for AHRQ support. Individual members of the Panel do not meet regularly and do not serve for fixed terms or long periods of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Grant applications for Cooperative Agreement Awards are to be reviewed and discussed at this meeting. These discussions are likely to include personal information concerning individuals associated with these applications. This information is exempt from mandatory disclosure under the above-cited statutes.

SEP Meeting on: Centers for Education and Research on Therapeutic (Limited Competitive Continuation Projects).

Date: June 10, 2002 (Open on June 10, from 8 a.m. to 8:15 a.m. and closed for remainder of the meeting).

Place: Holiday Inn Bethesda, 8120 Wisconsin Avenue, Georgia Room, 3rd Floor, Bethesda, MD 20814.

Contact Person: Anyone wishing to obtain a roster of members or minutes of this meeting should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Research Review, Education and Policy, AHRQ, 2101 East Jefferson Street, Suite 400, Rockville, Maryland 20852, Telephone (301) 594–1846.

Agenda items for this meeting are subject to change as priorities dictate.

Dated: May 10, 2002.

Carolyn M. Clancy,

Acting Director.

[FR Doc. 02–12310 Filed 5–15–02; 8:45 am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Toxic Substances and Disease Registry

Citizens Advisory Committee on Public Health Service (PHS) Activities and Research at Department of Energy (DOE) Sites: Oak Ridge Reservation Health Effects Subcommittee

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Agency for Toxic Substances and Disease Registry (ATSDR) and the Centers for Disease Control and Prevention (CDC) announce the following meeting.

Name: Citizens Advisory Committee on PHS Activities and Research at DOE Sites: Oak Ridge Reservation Health Effects Subcommittee (ORRHES).

Time and Date: 12 p.m.–8 p.m., June 18, 2002.

Place: YWCA of Oak Ridge, 1660 Oak Ridge Turnpike, Oak Ridge, Tennessee, 37830. Telephone: (865) 482–2008.

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

Background: A Memorandum of Understanding (MOU) signed in October 1990 and renewed in September 2000 between ATSDR and DOE, delineates the responsibilities and procedures for ATSDR's public health activities at DOE sites required under sections 104, 105, 107, and 120 of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA or ''Superfund''). These activities include health consultations and public health assessments at DOE sites listed on, or proposed for, the Superfund National Priorities List and at sites that are the subject of petitions from the public; and other health-related activities such as epidemiologic studies, health surveillance, exposure and disease registries, health education, substance-specific applied research, emergency response, and preparation of toxicological profiles. In addition, under an MOU signed in December 1990 with DOE and replaced by an MOU signed in 2000, the Department of Health and Human Services (HHS) has been given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities, workers at DOE facilities, and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production and use. HHS has delegated program responsibility to CDC.

Purpose: This subcommittee is charged with providing advice and recommendations to the Director, CDC, and the Administrator, ATSDR, pertaining to CDC's and ATSDR's public health activities and research at this DOE site. Activities shall focus on providing the public with a vehicle to express concerns and provide advice and recommendations to CDC and ATSDR. The purpose of this meeting is to receive updates from ATSDR and CDC, and to address other issues and topics, as necessary.