DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panels (SEP): Centers for Excellence To Promote a Healthier Workforce, Request for Application OH–05–006

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): Centers for Excellence to Promote a Healthier Workforce, Request for Application OH–05–006.

Times and Dates: 1 p.m.–5 p.m., November 10, 2005 (Closed).

Place: National Institute for Occupational Safety and Health, CDC, 1600 Clifton Road, NE MS E–74, Atlanta, GA 30333 Telephone Number 404.498.2556.

Status: 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: Centers for Excellence to Promote a Healthier Workforce, Request for Application OH–05–006.

Contact Person For More Information: Pamela J. Wilkerson, MPA, Scientific Review Administrator, National Institute for Occupational Safety and Health, CDC, 1600 Clifton Road, NE., Mailstop E–74, Atlanta, GA 30333, Telephone Number 404.498.2556.

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: October 17, 2005.

Alvin Hall,

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

[FR Doc. 05–21177 Filed 10–21–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Center for Environmental Health/Agency for Toxic Substances and Disease Registry

The Program Peer Review Subcommittee of the Board of Scientific Counselors (BSC), Centers for Disease Control and Prevention (CDC), National Center for Environmental Health (NCEH)/Agency for Toxic Substances and Disease Registry (ATSDR): Teleconference.

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

Name: Program Peer Review Subcommittee (PPRS).

Time and Date: 12:30 p.m.–2 p.m., November 8, 2005.

Place: The teleconference will originate at the National Center for Environmental Health/Agency for Toxic Substances and Disease Registry in Atlanta, Georgia. Please see ''Supplementary Information'' for details on accessing the teleconference.

Status: Open to the public, teleconference access limited only by availability of telephone ports.

Purpose: Under the charge of the Board of Scientific Counselors, NCEH/ATSDR the Program Peer Review Subcommittee will provide the BSC, NCEH/ATSDR with advice and recommendations on NCEH/ATSDR program peer review. They will serve the function of organizing, facilitating, and providing a long-term perspective to the conduct of NCEH/ATSDR program peer review.

Matters To Be Discussed: Update on Air Pollution and Respiratory Health Branch Peer Review; discuss Community Tribal Subcommittee feedback on peer review questionnaire; discuss process for evaluating peer review questionnaires; discuss selection of subcommittee representation on 2006 peer review workgroups, and review of Action Items from this meeting.

Agenda Items are subject to change as priorities dictate.

Supplementary Information: This conference call is scheduled to begin at 12:30 p.m. Eastern Standard Time. To participate in the teleconference, please dial (877) 315–6535 and enter conference code 383520.

Contact Person For More Information: Sandra Malcom, Committee Management Specialist, Office of Science, NCEH/ATSDR, M/S E–28, 1600 Clifton Road, NE., Atlanta, Georgia 30333, telephone 404/498–0003.

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 National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

Alvin Hall,

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

[FR Doc. 05–21175 Filed 10–21–05; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: Online Intergovernmental Referral Guide (IRG).

OMB No. 0970–0209.

Description: The IRG is an essential reference maintained by the Office of Child Support Enforcement (OCSE) that provides states with an effective and efficient way of viewing and updating state profile, address, and FIPS code information by consolidating data available through numerous discrete sources into a single centralized, automated repository.

Respondents: State IV–D Child Support Programs, Other Country Child Support Programs.

ANNUAL BURDEN ESTIMATES

Respondent	Number of respondents	Number of responses per respondent	Average urden hours per response	Total burden hours
States and Territories	54	18	.3	292
Foreign Countries and Canadian provinces	23	2	.1	5

Estimated Total Annual Burden Hours: 297.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. E-mail address: *infocollection@acf.hhs.gov.* All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: October 17, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05–21163 Filed 10–21–05; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Delegation of Authority

Notice is hereby given that, under the authority vested in me by the Secretary of Health and Human Services, I have redelegated to the Commissioner, Administration on Children, Youth and Families, with the authority to further redelegate, the authority to continue the administration of grants and contracts initially awarded in the Fiscal Years 2002, 2003 and 2004 under the Special Projects of Regional and National Significance (SPRANS) Community-based Abstinence Education Program, pursuant to Title V, section 501(a)(2) of the Social Security Act, as amended. The SPRANS Community-based Abstinence Education Program includes Community-based Abstinence Education grants, Abstinence Education Special Congressional Initiative Project grants, and the Abstinence Education Technical Assistance contract with the National Abstinence Clearinghouse. This delegation permits the Commissioner, Administration on Children, Youth and Families, to administer FY 2002, 2003 and FY 2004 SPRANS abstinence education grants under the terms and conditions of the initial awards, thereby allowing the continuation of the existing grants consistent with recent appropriations enactments (Pub. L. 108–477).

This delegation shall be exercised under the Department's policy on regulations and the existing delegation of authority to approve and issue regulations. This delegation excludes the authority to issue reports to Congress, to take final action to withhold funds from States and to act under the nondiscrimination provisions of the Social Security Act.

This delegation also supersedes all prior delegations of authority to the extent that they are inconsistent with the provisions of this delegation.

I hereby ratify any actions taken by the Commission, Administration on Children Youth and Families, or any other Administration on Children, Youth and Families official, which, in effect, involved the exercise of this authority prior to the effective date of this delegation.

This delegation is effective on the date of signature.

Dated: October 6, 2005.

Wade. F. Horn,

Assistant Secretary for Children & Families.

[FR Doc. 05–21162 Filed 10–21–05; 8:45am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N–0395]

Agency Information Collection Activities: Proposed Collection; Comment Request; Guidance for Industry on Formal Meetings With Sponsors and Applicants for Prescription Drug User Fee Act Products

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 to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection contained in the guidance for industry on formal meetings with sponsors and applicants for Prescription Drug User Fee Act (PDUFA) products.

DATES: Submit written or electronic comments on the collection of information by December 23, 2005.

ADDRESSES: Submit electronic comments on the collection of information to *http://www.fda.gov/dockets/ecomments*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Karen Nelson, Office of Management Programs (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–1482.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical