involving individual Federal Reserve System employees.

2. Any items carried forward from a previously announced meeting.

FOR FURTHER INFORMATION CONTACT:
Michelle A. Smith, Director, Office of Board Members; 202–452–2955.

SUPPLEMENTARY INFORMATION: You may call 202–452–3206 beginning at approximately 5 p.m. two business days before the meeting for a recorded announcement of bank and bank holding company applications scheduled for the meeting; or you may contact the Board’s Web site at http://www.federalreserve.gov for an electronic announcement that not only lists applications, but also indicates procedural and other information about the meeting.


Robert deV. Frierson, Deputy Secretary of the Board.

[FR Doc. 05–12392 Filed 6–17–05; 4:00 pm]
BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH & HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Incidence, Natural History, and Quality of Life of Diabetes in Youth, Request for Applications (RFA) DP–05–069

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): Incidence, Natural History, and Quality of Life of Diabetes in Youth, Request for Applications (RFA) DP–05–069.

Times and Dates: 7 a.m.–9 p.m., July 21, 2005(Closed); 8:30 a.m.–1:30 p.m., July 22, 2005(Closed).

Place: Double Tree Hotel, Buckhead, 13342 Peachtree Road NE, Atlanta, GA 30326, Telephone Number 404.231.1234.

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 Incidence, Natural History, and Quality of Life of Diabetes in Youth, Request for Applications (RFA) DP–05–069.

Contact Person for More Information: J. Felix Rogers, Ph.D., M.P.H., Scientific Review Administrator, National Immunization Program, CDC, 1600 Clifton Road NE., Mailstop E–05, Atlanta, GA 30333, Telephone 404.639.6101.

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 14, 2005.

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

[FR Doc. 05–12186 Filed 6–20–05; 8:45 am]
BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH & HUMAN SERVICES

Administration for Children and Families

Office of Planning, Research and Evaluation; Secondary Analysis of Data From the National Survey of Child Abuse and Neglect

Funding Opportunity Title: Secondary Analysis of Data from the National Survey of Child Abuse and Neglect (NSCAW).

Announcement Type: Initial.


CFDA Number: 93.647.

Due Date For Letter of Intent or Preapplications: Three weeks prior to due date.

Due Date for Applications: Application is due August 5, 2005.

Executive Summary: Funds are available to support grants for secondary analysis of data available from the National Survey on Child and Adolescent Well-Being. NSCAW provides longitudinal data from multiple informants on the functioning, well-being, and services provided to a national probability sample of children and families who come into contact with the child welfare system through an investigation of child maltreatment. Data are available through licensing agreements from the National Data Archive on Child Abuse and Neglect at Cornell University (http://www.ndacan.cornell.edu). Applicants’ planned analyses should be designed to advance the state of knowledge of child maltreatment, child welfare services, and/or child development for high risk children.

I. Funding Opportunity Description

A. Purpose

The purpose of this priority area is to announce the availability of funds to support grants for secondary analysis of data available from the National Survey on Child and Adolescent Well-Being. The planned analyses should be designed to advance the state of knowledge in child maltreatment, child welfare services, child and family services, and/or child development for high risk children.

B. Background

The National Survey of Child and Adolescent Well-Being, authorized under Section 429A of the Personal Responsibility and Work Opportunities Reconciliation Act, is the first nationally representative study that examines the functioning and well-being of children and families who come to the attention of the child welfare system. Although there has been an increasing emphasis on child well-being as a key outcome of child welfare services, and states are being held accountable for those outcomes, there has been little information, particularly on a national scale, to examine well-being within the context of the family and community environments and the service systems that are likely to affect children’s functioning. NSCAW was designed to begin to address this gap.

Children in the core sample (n=5504) were selected from those investigated by Child Protective Services in 92 primary sampling units (PSUs) during a 15-month sampling period beginning in the fall of 1999. Children are included in the sample and followed up whether or not their investigation resulted in a case opening; thus, NSCAW includes children who remain at home without services; those who remain at home and receive child welfare services; and those who are placed out of home in foster, kinship, or group care. A supplemental sample (n=727) was selected from children who were reaching their first anniversary in foster care during the same sampling period. Extensive information on child and family characteristics, service needs, and service receipt was collected directly from the target children, their caregivers, their caseworkers, and their teachers at baseline, and follow-up data were collected from all respondents at 18 months and 36 months post-baseline. In addition, information about services was collected from caregivers and caseworkers at 12 months post-baseline. Baseline contextual data are available from state administrators and local child welfare administrators in the PSUs.