information: 1 (866) 659–0537, Participant Pass Code 9933701.

Background: The Advisory Board was established under the Energy Employees Occupational Illness Compensation Program Act of 2000 to advise the President on a variety of policy and technical functions required to implement and effectively manage the new compensation program. Key functions of the Advisory Board include providing advice on the development of probability of causation guidelines that have been promulgated by the Department of Health and Human Services (HHS) as a final rule; advice on methods of dose reconstruction, which have also been promulgated by HHS as a final rule; advice on the scientific validity and quality of dose estimation and reconstruction efforts being performed for purposes of the compensation program; and advice on petitions to add classes of workers to the Special Exposure Cohort.

In December 2000, the President delegated responsibility for funding, staffing, and operating the Advisory Board to HHS, which subsequently delegated this authority to CDC. NIOSH implements this responsibility for CDC. The charter was issued on August 3, 2001, renewed at appropriate intervals, and will expire on August 3, 2013.

Purpose: The Advisory Board is charged with (a) Providing advice to the Secretary, HHS, on the development of guidelines under Executive Order 13179; (b) providing advice to the Secretary, HHS, on the scientific validity and quality of dose reconstruction efforts performed for this program; and (c) upon request by the Secretary, HHS, advise the Secretary on whether there is a class of employees at any Department of Energy facility who were exposed to radiation but for whom it is not feasible to estimate their radiation dose, and on whether there is reasonable likelihood that such radiation doses may have endangered the health of members of this class. The Subcommittee for Dose Reconstruction Reviews was established to aid the Advisory Board in carrying out its duty to advise the Secretary, HHS, on dose reconstruction.

Matters to be Discussed: The agenda for the Subcommittee meeting includes: discussion of dose reconstruction cases under review (sets 7–10); DCAS dose reconstruction quality management and assurance activities; and dose reconstruction issues from NIOSH 10-year review.

The agenda is subject to change as priorities dictate.

In the event an individual cannot attend, written comments may be submitted. Any written comments received will be provided at the meeting and should be submitted to the contact person below well in advance of the meeting.

Contact Person for More Information: Theodore Katz, Executive Secretary, NIOSH, CDC, 1600 Clifton Road, Mailstop E–20, Atlanta, Georgia 30333, *Telephone:* (513) 533–6800, *Toll Free:* 1 (800) CDC–INFO, *Email:* ocas@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 the Centers for Disease Control and Prevention, and the Agency for Toxic Substances and Disease Registry.

Dated: November 17, 2011.

Elaine L. Baker,

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

[FR Doc. 2011–30233 Filed 11–22–11; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

National Health and Nutrition Examination Survey (NHANES) DNA Samples

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).

ACTION: Notice.

SUMMARY: The National Health and Nutrition Examination Survey (NHANES) will not be receiving DNA proposals in 2012. NHANES is changing its plan for making DNA available for genetic research and its proposal guidelines. NHANES anticipates that the DNA Bank will be open for proposals approximately January 2013.

DATES: Effective date is date of publication in the **Federal Register.**

ADDRESSES: Geraldine McQuillan, PhD, Division of Health and Nutrition Examination Surveys, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 4204, Hyattsville, MD 20782, *Phone:* (301) 458–4371, *Fax:* (301) 458–4028, *E–Mail: NHANESgenetics@cdc.gov.*

FOR FURTHER INFORMATION CONTACT: Dr. Geraldine McQuillan, Division of Health and Nutrition Examination Surveys, National Center for Health Statistics, Centers for Disease Control and Prevention, 3311 Toledo Road, Room 4204, Hyattsville, MD 20782,

Phone: (301) 458–4371,

Fax: (301) 458–4028,

E-Mail: NHANESgenetics@cdc.gov.

Juliana K. Cyril,

Deputy Director, Office of Science Quality, Office of the Associate Director for Science, Centers for Disease Control and Prevention.

[FR Doc. 2011–30204 Filed 11–22–11; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[Docket No. CDC–2011–0011]

Public Health Service Guideline for Reducing Transmission of Human Immunodeficiency Virus (HIV), Hepatitis B Virus (HBV), and Hepatitis C Virus (HCV) Through Solid Organ Transplantation

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (DHHS).

ACTION: Extension of the public comment period.

SUMMARY: On September 21, 2011, the Centers for Disease Control and Prevention (CDC), located within the Department of Health and Human Services (HHS), published a notice in the **Federal Register** requesting public comment on the draft ''Public Health Service (PHS) Guideline for Reducing Transmission of Human Immunodeficiency Virus (HIV), Hepatitis B Virus (HBV), and Hepatitis C Virus (HCV) through Solid Organ Transplantation'' (76 FR 58517). Written and electronic comments were to be received on or before November 21, 2011. However, HHS/CDC has received requests for a 30 day extension of the comment period. In consideration of those requests, HHS/CDC is extending the comment period by 30 days to December 23, 2011.

CDC also published a supporting document for reference, the *Evidence Report.* The *Evidence Report* includes primary evidence, studies, and data tables that were used by the *Guideline* authors in developing the recommendations in the *Guideline.*

The draft *Guideline* is for use by organ procurement organizations (OPOs); transplant centers, including physicians, nurses, administrators, and clinical coordinators; laboratory personnel responsible for testing and storing donor and recipient specimens; and persons responsible for developing, implementing, and evaluating infection prevention and control programs for OPOs and transplant centers. This *Guideline* provides evidence-based recommendations for reducing unexpected transmission of HIV, HBV and HCV from deceased and living organ donors.

DATES: Written comments must be received on or before December 23, 2011.

ADDRESSES: Written comments may be submitted electronically or by mail. To