country areas in the State of South Dakota.

# C. Requesting a Hearing and Submitting Written Comments

Any request for a public hearing shall include the following: (1) The name, address, and telephone number of the individual, organization, or other entity requesting a hearing; (2) a brief statement of the requesting person's interest in the Regional Administrator's determination and of information that the requesting person intends to submit at such hearing; and (3) the signature of the individual making the request, or, if the request is made on behalf of an organization or other entity, the signature of the responsible official of the organization or other entity.

Notice of any hearing shall be given not less than fifteen (15) days prior to the time scheduled for the hearing. Such notice will be made by the Regional Administrator in the Federal Register and in newspapers of general circulation in the State of South Dakota. A notice will also be sent to the person(s) requesting the hearing as well as to the State of South Dakota. The hearing notice will include a statement of purpose, information regarding time and location, and the address and telephone number where interested persons may obtain further information. A final determination will be made upon review of the hearing record.

Frivolous or insubstantial requests for a hearing may be denied by the Regional Administrator. However, if a substantial request is made within thirty (30) days after this notice, a public hearing will be held.

Please bring this notice to the attention of any persons known by you to have an interest in this determination.

Dated: November 7, 2000.

## Jack W. McGraw,

Acting Regional Administrator, EPA, Region 8.

[FR Doc. 00–29646 Filed 11–17–00; 8:45 am]

# GENERAL SERVICES ADMINISTRATION

### Office of Communications; Cancellation of a Standard Form

**AGENCY:** General Services

Administration.

ACTION: Notice.

**SUMMARY:** The Department of State is cancelling the following Optional Form because of low usage:

OF 129, Individual Property Record Card

**DATES:** Effective November 20, 2000. **FOR FURTHER INFORMATION CONTACT:** Ms. Barbara Williams, General Services Administration, (202) 501–0581.

Dated: October 20, 2000.

### Barbara M. Williams,

Deputy Standard and Optional Forms Management Officer.

[FR Doc. 00–29616 Filed 11–17–00; 8:45 am] BILLING CODE 6820–34–M

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

## Advisory Committee for Injury Prevention and Control (ACIPC), Family and Intimate Violence Prevention Subcommittee; Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), the Centers for Disease Control and Prevention (CDC) announces the following subcommittee meeting.

Name: ACIPC Family and Intimate Violence Prevention Subcommittee.

Time and Date: 12 p.m.-4 p.m., November 28, 2000.

Place: The Hyatt Regency Atlanta, 265 Peachtree Street, N.E., Atlanta, Georgia 30303 Status: Open to the public, limited only by the space available.

Purpose: To advise and make recommendations to ACIPC and the Director, National Center for Injury Prevention and Control (NCIPC), regarding feasible goals for prevention and control of family and intimate violence and sexual assault. The Subcommittee will make recommendations regarding policies, strategies, objectives and priorities.

Matters to be Discussed: The Subcommittee will review current and planned family and intimate violence prevention program initiatives and activities.

Agenda items are subject to change as priorities dictate.

This notice is published less than 15 days prior to the meeting due to administrative delay.

Contact Person for More Information: Ileana Arias, Ph.D., Team Leader, Family and Intimate Violence Prevention Team, DVP, NCIPC, CDC, 4770 Buford Highway, NE, M/ S K60, Atlanta, Georgia 30341–3724, telephone 770/488–4410.

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 15, 2000.

#### Carolyn J. Russell,

Management Analysis and Services Office, Centers for Disease Control and Prevention. [FR Doc. 00–29719 Filed 11–16–00; 12:15 pm]

BILLING CODE 4163-18-P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

# Centers for Disease Control and Prevention

### Advisory Committee for Injury Prevention and Control (ACIPC), Science and Program Review Subcommittee Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Public Law 92–463), the Centers for Disease Control and Prevention (CDC) announces the following Federal Advisory committee meetings.

*Name*: ACIPC Science and Program Review Subcommittee.

*Time and Date:* 1:30 p.m.–5 p.m., November 28, 2000.

Place: The Hyatt Regency Atlanta, 26 Peachtree Street, N.E., Atlanta, Georgia 30303

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

*Purpose:* The Subcommittee provides advice on the needs, structure, progress and performance of NCIPC programs. The Subcommittee provides secondlevel scientific and programmatic review for applications for research grants, cooperative agreements, and training grants related to injury control and violence prevention, and recommends approval of projects that merit further consideration for funding support. The Subcommittee also advises on priorities for research to be supported by contracts, grants, and cooperative agreements and provides concept review of program proposals and announcements.

Matters to be Discussed: The Subcommittee will discuss the national violent death reporting system, NCIPC programmatic reviews, cooperative agreement funding for fiscal year 2000, and the external peer review of intramural research.

Agenda items are subject to change as priorities dictate.

Name: Advisory Committee for Injury Prevention and Control (ACIPC).

Time and Date: 8:30 a.m.–3:30 p.m., November 29, 2000.

Place: The Hyatt Regency Atlanta, 26 Peachtree Street, N.E., Atlanta, Georgia 30303. *Status:* Open to the public, limited only by the space available.

Purpose: The Committee advises and makes recommendations to the Secretary, the Assistant Secretary for Health, the Director, CDC, and Director, National Center for Injury Prevention and Control (NCIPC) regarding feasible goals for the prevention and control of injury. The Committee makes recommendations regarding policies, strategies, objectives, and priorities, and reviews progress toward injury prevention and control. The Committee provides advice on the appropriate balance of intramural and extramural research, and also provides guidance on the needs, structure, progress and performance of intramural programs, and on extramural scientific program matters. The Committee provides second-level scientific and programmatic review for applications for research grants, cooperative agreements, and training grants related to injury control and violence prevention, and recommends approval of projects that merit further consideration for funding support. The Committee also recommends areas of research to be supported by contracts and cooperative agreements and provides concept review of program proposals and announcements.

Matters to be Discussed: Following the Acting Director's update, which will include an introduction of the new NCIPC Director, NCIPC's Division of Violence Prevention will give an overview of violence against women programs. The Committee will also discuss reports from the November 28, 2000, meetings of the Subcommittee on Family and Intimate Violence Prevention and Subcommittee on Science and Program Review.

This notice is published less than 15 days prior to the meeting due to administrative delay.

Contact Person for More Information: Mr. Thomas E. Blakeney, Acting Executive Secretary, ACIPC, NCIPC, CDC, 4770 Buford Highway, NE, M/S K61, Atlanta, Georgia 30341–3724, telephone 770/488–1481.

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 15, 2000.

#### Carolyn J. Russell,

Management Analysis and Services Office, Centers for Disease Control and Prevention. [FR Doc. 00–29720 Filed 11–16–00; 12:15 pm]

BILLING CODE 4163-18-P

# DEPARTMENT OF HEALTH AND HUMAN SERVICES

### **Food and Drug Administration**

### Request for Nominations for Voting Members on Public Advisory Committees

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.

The Food and Drug Administration (FDA) is requesting nominations for voting members to serve on the Allergenic Products Advisory Committee, Biological Response Modifiers Advisory Committee, Blood Products Advisory Committee, Transmissible Spongiform Encephalopathies Advisory Committee, and the Vaccines and Related Biological Products Advisory Committee in the Center for Biologics Evaluation and Research (CBER). Nominations will be accepted for vacancies that will or may occur through January 31, 2002.

FDA has a special interest in ensuring that women, minority groups, and individuals with disabilities are adequately represented on advisory committees and, therefore, encourages nominations of qualified candidates from these groups.

**DATES:** Because scheduled vacancies occur on various dates throughout each year, no cutoff date is established for the receipt of nominations. However, when possible, nominations should be received at least 6 months before the date of scheduled vacancies for each year, as indicated in this notice.

**ADDRESSES:** All nominations and curricula vitae should be sent to the addresses below.

## FOR FURTHER INFORMATION CONTACT:

Regarding nominations, except for consumer representatives: Nancy T. Cherry, Scientific Advisors and Consultants Staff, Center for Biologics Evaluation and Research (HFM–71), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852–1448, 301–827–0314.

Regarding nominations for consumer representatives: Maureen A. Hess, Office of Consumer Affairs (HFE–50), Food and Drug Administration, 5600 Fishers

Lane, Rockville, MD 20857, 301–827–4421.

**SUPPLEMENTARY INFORMATION:** FDA is requesting nominations of voting members with appropriate expertise for vacancies listed below.

- 1. Allergenic Products Advisory Committee: Three vacancies occurring August 31, 2001; immunology, pediatrics, internal medicine, biochemistry, statistics, and related scientific fields.
- 2. Biological Response Modifiers Advisory Committee: Five vacancies occurring March 31, 2001; biological response modifiers, immunology, virology, molecular biology, rDNA technology, infectious diseases, viral oncology, statistics, and cellular kinetics.
- 3. Blood Products Advisory Committee: Nine vacancies occurring September 30, 2001; clinical and administrative medicine, hematology, immunology, blood banking, surgery, internal medicine, biochemistry, engineering, statistics, biological and physical sciences, and other related scientific fields.
- 4. Transmissible Spongiform Encephalopathies Advisory Committee: Three vacancies occurring January 31, 2002; clinical administrative medicine, hematology, virology, neurology, infectious diseases, immunology, blood banking, surgery, internal medicine, biochemistry, biostatistics, epidemiology, biological and physical sciences, sociology/ethics, and other related professions.
- 5. Vaccines and Related Biological Products Advisory Committee: Three vacancies occurring January 31, 2002; immunology, molecular biology, rDNA, virology, bacteriology, epidemiology, biostatistics, allergy, preventive medicine, infectious diseases, pediatrics, microbiology, and biochemistry.

#### **Functions**

Allergenic Products Advisory Committee

Reviews and evaluates available data concerning the safety, effectiveness, and adequacy of labeling of marketed and investigational allergenic biological products or materials that are administered to humans for the diagnosis, prevention, or treatment of allergies and allergic diseases.

Biological Response Modifiers Advisory Committee

Reviews and evaluates available data relating to the safety, effectiveness, and appropriate use of biological response modifiers which are intended for use in