

**ADDRESSES:** Comments should be submitted to the GSA Privacy Act Officer (CI), Office of the Chief People Officer, General Services Administration, 1800 F Street NW, Washington DC 20405.

**FOR FURTHER INFORMATION CONTACT:** The GSA Privacy Act Officer at the above address, or call 202-501-1452.

Dated: August 27, 2004.

**June V. Huber,**

*Director, Office of Information Management,  
Office of the Chief People Officer.*

### **GSA/HRO-9**

*System name:* Emergency Management Records (GSA/HRO-9).

*System location:* The system is the responsibility of the GSA Office of Emergency Management, located at 1800 F Street NW, Washington DC 20405. System records are located in the GSA Central Office and regional offices with assigned emergency management responsibilities.

*Categories of individuals covered by the system:* All GSA associates, contractor employees, and other key governmental and non-governmental persons essential to carrying out emergency activities or with a need to know of actions taken by GSA in an emergency.

*Categories of records in the system:* The records, composed of emergency notification rosters and files, may consist of paper records and/or electronic databases, including the Emergency Management Information Database (EMID), the Quick Notify database, and continuity of operations (COOP) files. The data may be consolidated into a centralized emergency contact database to expedite communication. Personal information in the system records includes name; office, cell, and home telephone numbers; out-of-area contact telephone numbers; home address; home e-mail address; and home fax number. System records also may include special needs information such as medical, mobility, and transportation requirements by individuals. Additional information may include official titles and emergency assignments for individuals in the system.

*Authority for maintaining the system:* The Federal Property and Administrative Services Act of 1949, as amended 40 U.S.C. §§ 101 *et seq.*; E.O. Order 12565, Assignment of Emergency Preparedness Responsibilities; and Presidential Decision Directive 67, Ensuring Constitutional Government and Continuity of Government Operations.

*Purpose:* To maintain current information on GSA associates and other persons covered by this system for use by persons with emergency management responsibilities to notify officials, employees, and other affected individuals of conditions that require their urgent attention during a public or personal emergency.

### **Routine uses of records maintained in the system, including types of users and purposes of such uses:**

System information may be used by authorized individuals in the performance of duties associated with their emergency management responsibilities. Routine uses are:

a. To disclose needed information to a Federal, State, or local agency investigating, prosecuting, or enforcing a statute, rule, regulation, or order, where GSA becomes aware of a possible violation of civil or criminal law or regulation.

b. To disclose information to a Member of Congress or a congressional staff member at the request of the individual who is the subject of the record.

c. To disclose information to another Federal agency or to a court where the Government is a party to a judicial proceeding before the court.

d. To disclose information to a Federal agency, in response to its request, in connection with hiring or retaining an associate, issuing a security clearance, conducting a security or suitability investigation, classifying a job, letting a contract, or issuing a license, grant, or other benefit by the requesting agency, to the extent that the information is necessary to the agency's decision on the matter.

e. To disclose information to an appeal, grievance, or formal complaints examiner; equal employment opportunity investigator; arbitrator; exclusive representative; or other official engaged in investigating, or settling a grievance, complaint, or appeal filed by an employee.

f. To disclose information to the Office of Personnel Management (OPM) and the Government Accountability Office (GAO) when the information is required for evaluation of program activities.

g. To disclose information to the National Archives and Records Administration (NARA) for records management purposes.

h. To disclose information to an expert, consultant, or contractor in the performance of a Federal government duty to which the information is relevant.

### **Policies and practices for storing, accessing, retrieving, retaining, and disposing of records in the system:**

*Storage:* System records may be stored on paper or electronically in secure locations or computer systems.

*Retrievability:* Records may be retrieved by name, organization, location, teleworking capability, or special medical or other health or safety need of an individual.

*Safeguards:* When not in use by an authorized person, the records are secured from unauthorized access. Paper records are placed in lockable file cabinets or in secured areas. Electronic records are protected by passwords, access codes, and other appropriate technical security measures.

*Retention and disposal:* Disposal of system records is according to the Handbook, GSA Records Maintenance and Disposition System (OAD P 1820.2A), and the requirements of the National Archives and Records Administration.

*System manager(s) and address:* The official with overall responsibility for the system of records is the Director, Office of Emergency Management (ACE), 1800 F Street NW, Washington DC 20405. GSA Services, Staff Offices, and regions are responsible for the integrity of data within their jurisdictions.

*Notification procedure:* Individuals may determine whether the system contains their records by submitting a request to the System Manager or the appropriate Service, Staff Office, or regional official.

*Record access procedures:* An individual may obtain information on the procedures for gaining access to their records from the System Manager or the appropriate Service, Staff Office, or regional official.

*Procedures for contesting records:* Individuals wishing to request amendment of their records should contact the System Manager or the appropriate Service, Staff Office, or regional official.

*Record sources:* The records contain information provided by the individuals themselves, their supervisors, or their Service, Staff Office, or region.

[FR Doc. 04-20563 Filed 9-10-04; 8:45 am]

BILLING CODE 6820-34-S

## **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

### **Chronic Fatigue Syndrome Advisory Committee**

**AGENCY:** Office of the Secretary, HHS.

**ACTION:** Notice of meeting.

**SUMMARY:** As stipulated by the Federal Advisory Committee Act, the U.S. Department of Health and Human Services is hereby giving notice that the Chronic Fatigue Syndrome Advisory Committee (CFSAC) will hold a meeting. The meeting will be open to the public.

**DATES:** The meeting will be held on Monday, September 27, 2004, from 9 a.m. to 5 p.m.

**ADDRESSES:** Department of Health and Human Services, Hubert H. Humphrey Building, 200 Independence Avenue, SW., Room 705A, Washington, DC 20201.

**FOR FURTHER INFORMATION, CONTACT:** Dr. Larry E. Fields, Executive Secretary, Chronic Fatigue Syndrome Advisory Committee, U.S. Department of Health and Human Services, Hubert H. Humphrey Building, 200 Independence Avenue, SW., Room 719H, Washington, DC 20201; (202) 690-7694.

**SUPPLEMENTARY INFORMATION:** CFSAC was established on September 5, 2002, to replace the Chronic Fatigue Syndrome Coordinating Committee. CFSAC was established to advise, consult with, and make recommendations to the Secretary, through the Assistant Secretary for Health, on a broad range of topics including (1) the current state of knowledge and research about the epidemiology and risk factors relating to chronic fatigue syndrome, and identifying potential opportunities in these areas; (2) current and proposed diagnosis and treatment methods for chronic fatigue syndrome; and (3) development and implementation of programs to inform the public, health care professionals, and the biomedical, academic, and research communities about chronic fatigue syndrome advances.

The tentative agenda for this meeting is as follows:

- 9 a.m. Chairperson
  - Call to Order
  - Request for Roll Call
  - Introductions and Opening Remarks
  - Approval of the Minutes of June 21st, 2004
  - Discussion
- 9:20 a.m. Executive Secretary
  - Roll Call
  - Summary of Public Comments
  - Operational Matters
  - Discussion
- 9:30 a.m. Invited Organizational Updates
  - K. Kimberly McCleary
  - CFIDS Association of America
  - Research Funding

## Discussion

Jill McLaughlin

National CFIDS Foundation, Inc.

Patient Issues

Discussion

10:30 a.m. Break

10:45 a.m. Ex Officio Members

Requested follow-ups

Other Updates

Discussion

11:30 a.m. Public Comment

12 noon Lunch Break

1 p.m. Subcommittee Updates

Disabilities: Lyle Lieberman, Chair

Education: Dr. Roberto Patarca, Chair

Research: Dr. Nahid Mohagheghpour, Chair

2:24 p.m. Break

3:15 p.m. Planning: Future Directions

Recommendations

Invited Organizational Updates

Other Matters

Discussion

4 p.m. Public Comment

4:30 p.m. Summary

Action Steps

Timelines

5 p.m. Adjournment

Public attendance at the meeting is limited to space available. Individuals must provide a photo ID for entry into the meeting. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person. Members of the public will have the opportunity to provide comments at the meeting. Pre-registration is required for public comment by September 17, 2004. Any individual who wishes to participate in the public comment session should call the telephone number listed in the contact information to register. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed material distributed to CFSAC members should submit materials to the Executive Secretary, CFSAC, whose contact information is listed above prior to close of business September 17, 2004.

Dated: September 8, 2004.

**Dr. Larry E. Fields,**

Executive Secretary, Chronic Fatigue Syndrome Advisory Committee.

[FR Doc. 04-20635 Filed 9-9-04; 10:03 am]

**BILLING CODE 4150-28-P****DEPARTMENT OF HEALTH AND HUMAN SERVICES****Food and Drug Administration**

[Docket No. 2004D-0377]

**International Conference on Harmonisation; Draft Guidance on E14 Clinical Evaluation of QT/QTc Interval Prolongation and Proarrhythmic Potential for Non-Antiarrhythmic Drugs; Availability**

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

**ACTION:** Notice.

**SUMMARY:** The Food and Drug Administration (FDA) is announcing the availability of a draft guidance entitled "E14 Clinical Evaluation of QT/QTc Interval Prolongation and Proarrhythmic Potential for Non-Antiarrhythmic Drugs." The draft guidance was prepared under the auspices of the International Conference on Harmonisation of Technical Requirements for Registration of Pharmaceuticals for Human Use (ICH). The draft guidance provides recommendations to sponsors concerning clinical studies to assess the potential of a new drug to cause cardiac arrhythmias, focusing on the assessment of changes in the QT/QTc interval on the electrocardiogram as a predictor of risk. The draft guidance is intended to encourage the assessment of drug effects on the QT/QTc interval as a standard part of drug development and to encourage the early discussion of this assessment with FDA.

**DATES:** Submit written or electronic comments on the draft guidance by December 13, 2004.

**ADDRESSES:** Submit written comments on the draft guidance to the Division of Dockets Management (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to <http://www.fda.gov/dockets/ecomments>. Submit written requests for single copies of the draft guidance to the Division of Drug Information (HFD-240), Center for Drug Evaluation and Research, Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857; or the Office of Communication, Training and Manufacturers Assistance (HFM-40), Center for Biologics Evaluation and Research (CBER), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448. The guidance may also be obtained by mail by calling the CBER Voice Information System at 1-800-835-4709 or 301-827-