Skidmore, Assistant to the Board, Office of Board Members at 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.

Board of Governors of the Federal Reserve System, March 7, 2008.

Robert deV. Frierson,

Deputy Secretary of the Board. [FR Doc. 08–1011 Filed 3–7–08; 2:22 pm] BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Chronic Fatigue Syndrome Advisory Committee

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science. **ACTION:** Notice.

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, May 5, 2008 and Tuesday, May 6, 2008. The meeting will be held from 9 a.m. to approximately 5 p.m. on both days.

ADDRESSES: Department of Health and Human Services; Room 800 Hubert H. Humphrey Building; 200 Independence Avenue, SW.; Washington, DC 20201. FOR FURTHER INFORMATION CONTACT: Dr. Anand K. Parekh, Executive Secretary, Chronic Fatigue Syndrome Advisory Committee; Department of Health and Human Services; 200 Independence Avenue, SW., Room 727H; Washington, DC 20201; (202) 401–7605.

SUPPLEMENTARY INFORMATION: CFSAC was established on September 5, 2002. The Committee 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 agenda for this meeting is being developed. The agenda will be posted on the CFSAC Web site, *http://www.hhs.gov/advcomcfs*, when it is finalized.

Public attendance at the meeting is limited to space available. Individuals must provide a photo ID for entry into the building where the meeting is scheduled to be held. 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. The committee is very interested to hear about experiences that individuals with chronic fatigue syndrome have had interfacing with the medical community. In addition, the committee welcomes specific comments on issues surrounding chronic fatigue syndrome research, provider education, and quality of life. Individuals who wish to address the Committee during the public comment session must preregister by May 1, 2008. 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. Members of the public who wish to have printed material distributed to CFSAC members for discussion should submit, at a minimum, one copy of the material to the Executive Secretary, CFSAC prior to close of business on May 1, 2008. Contact information for the Executive Secretary, CFSAC is listed above.

Dated: March 5, 2008.

Anand K. Parekh,

Executive Secretary, Chronic Fatigue Syndrome Advisory Committee. [FR Doc. E8–4791 Filed 3–10–08; 8:45 am]

BILLING CODE 4150-42-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Secretary's Advisory Committee on Human Research Protections

AGENCY: Department of Health and Human Services, Office of the Secretary, Office of Public Health and Science. **ACTION:** Notice.

SUMMARY: Pursuant to section 10(a) of the Federal Advisory Committee Act, U.S.C. Appendix 2, notice is hereby given that the Secretary's Advisory Committee on Human Research Protections (SACHRP) will hold its fifteenth meeting. The meeting will be open to the public.

DATES: The meeting will be held on Thursday, March 27, 2008 from 8:30 a.m. until 4:30 p.m. and Friday, March 28, 2008 from 8:30 a.m. until 4:30 p.m.

ADDRESSES: The Sheraton National Hotel, 900 South Orme Street, Arlington, Virginia 22204. Phone: 703–521–1900.

FOR FUTHER INFORMATION CONTACT: Ivor Pritchard, Ph.D., Acting Director, Office for Human Research Protections (OHRP), or Kevin Prohaska, D.O., M.P.H., Acting Executive Director, SACHRP; U.S. Department of Health and Human Services, 1101 Wootton Parkway, Suite 200, Rockville, Maryland 20852; 240–453–8231; fax: 240–453–6909; e-mail address: sachrp@osophs.dhhs.gov.

SUPPLEMENTARY INFORMATION: Under the authority of 42 U.S.C. 217a, section 222 of the Public Health Service Act, as amended, SACHRP was established to provide expert advice and recommendations to the Secretary of Health and Human Services and the Assistant Secretary for Health on issues and topics pertaining to or associated with the protection of human research subjects.

On March 27, 2008 SACHRP will receive and discuss a report from the Subcommittee on Inclusion of Individuals with Impaired Decision-Making in Research. The Subcommittee on Inclusion of Individuals with Impaired Decision-Making in Research is charged with developing recommendations for consideration by SACHRP about whether guidance and/ or additional regulations are needed for research involving individuals with impaired decision-making capacity. This Subcommittee was formed as a result of discussions during the July 31-August 1, 2006 SACHRP meeting. In addition, an invited panel will discuss common designs associated with quality improvement, quality assurance, and

health services activities, as well as ethical considerations that may be relevant to such activities.

On March 28, 2008 the Committee will receive and discuss reports from an ad hoc subcommittee and from the Subpart A Subcommittee. The ad hoc subcommittee was established after the October 29-30, 2007 meeting of SACHRP to consider recommendations relative to encouraging diversity in clinical trials and conducting research in the disaster setting. The Subpart A Subcommittee is charged with developing recommendations for consideration by SACHRP about the application of Subpart A of 45 CFR part 46 in the current research environment. This subcommittee was established by SACHRP at its October 4-5, 2006 meeting.

Public attendance at the meeting is limited to space available. Individuals who plan to attend the meeting and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact persons. Members of the public will have the opportunity to provide comments on both days of the meeting. Public comment will be limited to five minutes per speaker. Any members of the public who wish to have printed materials distributed to SACHRP members for this scheduled meeting should submit materials to the Acting Executive Director, SACHRP, prior to the close of business Monday, March 17, 2008. Information about SACHRP and the draft meeting agenda will be posted on the SACHRP Web site at: http:// www.hhs.gov/ohrp/sachrp/index.html.

Dated: March 5, 2008. **Ivor A. Pritchard,** Acting Director, Office for Human Research Protections, Acting Executive Secretary, Secretary's Advisory Committee on Human Research Protections. [FR Doc. E8–4793 Filed 3–10–08; 8:45 am]

BILLING CODE 4150-36-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection Comment Request

In compliance with the requirement for opportunity for public comment on proposed data collection projects (section 3506(c)(2)(A) of Title 44, United States Code, as amended by the Paperwork Reduction Act of 1995, Pub. L. 104–13), the Health Resources and Services Administration (HRSA) will publish periodic summaries of proposed projects being developed for submission to the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995. To request more information on the proposed project or to obtain a copy of the data collection plans, call the HRSA Reports Clearance Officer on (301) 443–1129.

Comments are invited on: (a) Whether the proposed collection of information is necessary for the 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) ways to enhance 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.

Proposed Project: Data Collection Tool for State Offices of Rural Health Grant Program: (New)

The mission of the Office of Rural Health Policy (ORHP) is to sustain and improve access to quality care services for rural communities. In its authorizing language (Sec. 711 of the Social Security Act [42 U.S.C. 912]), Congress charged ORHP with administering grants, cooperative agreements, and contracts to provide technical assistance and other activities as necessary to support activities related to improving health care in rural areas.

In accordance with the Public Health Service Act, section 338J, 42 U.S.C. 254r, HRSA proposes to revise the State Offices of Rural Health Grant Program— Guidance and Forms for the Application. The guidance is used annually by 50 States in writing applications for grants under the State Offices of Rural Health Grant Program (SORH) and in preparing the required report.

ORHP seeks to expand the information gathered from grantees on their efforts to provide technical assistance to clients within their State. SORH grantees would be required to submit a Technical Assistance Report that includes: (1) The total number of technical assistance encounters provided directly by the grantee; and, (2) the total number of clients that received direct technical assistance from the grantee. Submission of the Technical Assistance Report would be done via e-mail to ORHP no later than 30 days after the end of each twelvemonth budget period.

The estimated average annual burden is as follows:

Form	Number of respondents	Responses per respondent	Burden hours per response	Total burden hours
Technical Assistance Report Total	50 50	1	12.5	562.5 562.5