Reserve Bank indicated. The notices also will be available for inspection at the office of the Board of Governors. Interested persons may express their views in writing to the Reserve Bank indicated for that notice or to the offices of the Board of Governors. Comments must be received not later than November 27, 2007.

A. Federal Reserve Bank of St. Louis (Glenda Wilson, Community Affairs Officer) 411 Locust Street, St. Louis, Missouri 63166-2034:

1. McLane Family Control Group, Poplar Bluff, Missouri, consisting of Joseph T. McLane, Jana McLane Brown, Jerri Ann McLane, the Norma McLane Smith Revocable Trust, Norma McLane Smith as trustee of Trust, and the Midwest Bancorporation, Inc. and Affiliates Employee Stock Ownership Plan Trust Joseph T. McLane as trustee, all of Poplar Bluff, Missouri; to acquire additional voting shares of Midwest Bancorporation, Inc., Poplar Bluff, Missouri, and thereby indirectly acquire additional voting shares of First Midwest Bank of Dexter, Missouri and First Midwest Bank of the Ozarks, Piedmont, Missouri.

Board of Governors of the Federal Reserve System, November 7, 2007.

Robert deV. Frierson,

Deputy Secretary of the Board.

[FR Doc. E7–22105 Filed 11–9–07; 8:45 am]

BILLING CODE 6210–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–New]

Agency Information Collection Request; 30-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, e-mail your request, including your address, phone number, OMB number, and OS document identifier, to *Sherette.funncoleman@hhs.gov,* or call the Reports Clearance Office on (202) 690–6162. Written comments and recommendations for the proposed information collections must be received within 30 days of this notice directly to the OS OMB Desk Officer all comments must be faxed to OMB at 202–395–6974.

Title: Safe Harbor for Federally Qualified Health Centers Arrangements under the Anti-kickback Statute—OMB No. 0990–New—Office of Inspector General (OIG).

Proposed Project: The Office of the Inspector General (OIG), Office of the Secretary (OS), Department of Health and Human Services (HHS) is requesting a 3-year clearance for the data collection under the anti-kickback statute, as described below. In order for an arrangement between a health center and a donor individual or entity to enjoy safe harbor protection, the arrangement: (1) Must be set out in writing (§ 1001.952(w)(1)(i)(A)); (2) the written agreement must be signed by the parties (§ 1001.952(w)(1)(i)(B)); (3) the written agreement must cover, and specify the amount of, all goods, items, services, donations, or loans provided by the individual or entity to the health center (§ 1001.952(w)(1)(i)(C)); (4) the health center must document its basis for its reasonable expectation that the arrangement will benefit a medically underserved population (§ 1001.952(w)(3)); and (5) the health center, at reasonable intervals, must reevaluate the arrangement to ensure that it is expected to continue to benefit a medically underserved population, and must document the re-evaluation contemporaneously (§ 001.952(w)(4)).

OIG may request to see documentation kept pursuant to the safe harbor in order to determine compliance with the terms of the safe harbor and the fraud and abuse laws. Compliance with the safe harbor is voluntary, and no party is ever required to comply with the safe harbor.

The safe harbor does not entail a routine and continuous affirmative collection of data from the regulated community. However, health centers that choose to avail themselves of the safe harbor must have initial documentation and a re-evaluation of the arrangement at least annually. The respondents are businesses and/or other private sector for-profit and not-for-profit institutions.

OIG previously solicited comments on this section of the PRA on July 1, 2005, upon publication of the 60-day notice of proposed rulemaking (70 FR 38081). OIG did not receive any comments specifically addressing the PRA in response to that notice; however, OIG is now providing an additional opportunity for comment on the PRA aspect of the rule only.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of respondent	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Health Centers (Administrative Professionals)	1,873	1	1	1,873

Dated: November 1, 2007.

Alice Bettencourt,

Office of the Secretary, Paperwork Reduction Act Reports Clearance Officer.

[FR Doc. E7–22086 Filed 11–9–07; 8:45 am]

BILLING CODE 4152–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Meeting of the Chronic Fatigue Syndrome Advisory Committee

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

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 Wednesday, November 28, 2007 and Thursday, November 29, 2007. 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) 260–2873.

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. Individuals who wish to address the Committee during the public comment session must pre-register by November 26, 2007. 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 discuss should submit, at a minimum, one copy of the material to the Executive Secretary, CFSAC prior to close of business on November 26, 2007. Contact information for the Executive Secretary, CFSAC is listed above.

Dated: November 6, 2007.

Anand K. Parekh,

Executive Secretary, Chronic Fatigue Syndrome Advisory Committee.

[FR Doc. E7–22100 Filed 11–9–07; 8:45 am]

BILLING CODE 4150–42–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Toxicology Program (NTP); Report on Carcinogens (RoC); Availability of the Draft Background Documents for Aristolochic Acid Related Exposures (Two Candidate Substances: Botanical Products Containing Aristolochic Acid and Aristolochic Acid) and Riddelliine and Request for Public Comment on the Draft Background Documents; Announcement of the Aristolochic Acid Related Exposures and Riddelliine Expert Panel Meeting

AGENCY: National Institute of Environmental Health Sciences (NIEHS); National Institutes of Health (NIH), Department of Health and Human Services (HHS).

ACTION: Request for public comments and meeting announcement.

SUMMARY: The NTP announces the availability of the draft background documents for (1) aristolochic acid related exposures (the background document describes information on two candidate substances: Botanical products containing aristolochic acid and aristolochic acid) and (2) riddelliine on November 13, 2007, on the RoC Web site (*http://ntp.niehs.nih.gov/go/10091* see aristolochic acid related exposures or riddelliine) or in printed text from the RoC (see **FOR FURTHER INFORMATION CONTACT** below). The NTP invites the submission of public comments on the two draft background documents (see **SUPPLEMENTARY INFORMATION** below). The expert panel will meet on January 24–25, 2008, at the Chapel Hill Sheraton Hotel, One Europa Drive, Chapel Hill, North Carolina 27514, to peer review the draft background documents for aristolochic acid related exposures and riddelliine and, once completed, make a recommendation regarding the listing status (i.e., known to be a human carcinogen, reasonably anticipated to be a human carcinogen, or not to list) for botanical products containing aristolochic acid, for aristolochic acid, and for riddelliine in the 12th Edition of the RoC (12th RoC). The RoC expert panel meeting is open to the public with time scheduled for oral public comments. Attendance is limited only by the available meeting room space. Following the expert panel meeting and completion of the expert panel report, the NTP will post the final version of the background documents and the expert panel peer review reports on the RoC Web site.

DATES: The expert panel meeting for aristolochic acid related exposures and riddelliine will be held on January 24–25, 2008. The draft background documents for these substances will be available for public comment on November 13, 2007. The deadline to submit written comments is January 11, 2008, and the deadline for pre-registration to attend the meeting and provide oral comments at the meeting is January 18, 2008. Persons needing special assistance, such as sign language interpretation or other reasonable accommodations in order to attend, should contact 919–541–2475 (voice), 919–541–4644 TTY (text telephone), through the Federal TTY Relay System at 800–877–8339, or by e-mail to *niehsoeeo@niehs.nih.gov*. Requests should be made at least seven business days in advance of the event.

ADDRESSES: The RoC expert panel meeting on aristolochic acid related exposures and riddelliine will be held at the Chapel Hill Sheraton Hotel, One Europa Drive, Chapel Hill, North Carolina 27514. Access to on-line registration and materials for the meeting is available on the RoC Web site (*http://ntp.niehs.nih.gov/go/29679*). Comments on the draft background documents should be sent to Dr. C. W. Jameson, RoC Director, NIEHS, P.O. Box 12233, MD EC–14, Research Triangle Park, NC 27709, Fax: (919) 541–0144, or *jameson@niehs.nih.gov*. Courier address: Report on Carcinogens, 79 T.W. Alexander Drive, Building 4401, Room 3118, Research Triangle Park, NC 27709.

FOR FURTHER INFORMATION CONTACT: Dr. C. W. Jameson, RoC Director, 919–541–4096, *jameson@niehs.nih.gov*.

SUPPLEMENTARY INFORMATION:

Background

On April 16, 2007 (72 FR 18999 available at *http://ntp.niehs.nih.gov/go/9732*), the NTP announced the RoC review process for the 12th RoC. An expert panel meeting is being convened on January 24–25, 2008, to review three candidate substances (botanical products containing aristolochic acid,