summary will list other comments received on or before 15 days after the meeting, or August 1, 2007. The summary will also display CMS' tentative payment determinations, and interested individuals may submit written comments on the tentative payment determinations by September 21, 2007 to the address specified in the summary.

III. Registration Instructions

We are coordinating the public meeting registration. Beginning June 18, 2007 registration may be completed on-line at *http://www.cms.hhs.gov/ClinicalLabFeeSched.* The following information must be submitted when registering: Name; company name; address; telephone number(s); and E-mail address(es).

When registering, individuals who want to make a presentation must also specify which new clinical laboratory test code(s) they will be presenting. A confirmation will be sent upon receipt of the registration.

Registration Deadline: Individuals must register by July 11, 2007.

IV. Security, Building, and Parking Guidelines

The meeting will be held in a Federal government building; therefore, Federal security measures are applicable. In planning your arrival time, we recommend allowing additional time to clear security. In order to gain access to the building and grounds, participants must bring a government-issued photo identification and a copy of your written meeting registration confirmation. Persons without proper identification may be denied access to the building.

Individuals who are not registered in advance will not be permitted to enter the building and will be unable to attend the meeting. The public may not enter the building earlier than 30 to 45 minutes prior to the convening of the meeting.

Security measures also include inspection of vehicles, inside and out at the entrance to the grounds. In addition, all persons entering the building must pass through a metal detector. All items brought to CMS, whether personal or for the purpose of demonstration or to support a presentation, are subject to inspection.

V. Special Accommodations

Individuals attending a meeting who are hearing or visually impaired and have special requirements, or a condition that requires special assistance, should provide the information upon registering for the meeting.

Authority: Section 1102, 1833(h), and 1871 of the Social Security Act (42 U.S.C. 1302, 42 U.S.C. 13951, and 42 U.S.C. 1395hh)

(Catalog of Federal Domestic Assistance Program No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: March 22, 2007.

Leslie V. Norwalk,

Acting Administrator, Centers for Medicare & Medicaid Services.

[FR Doc. E7–9525 Filed 5–24–07; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS–3172–N]

Medicare Program; Meeting of the Medicare Coverage Advisory Committee—July 18, 2007

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Notice.

SUMMARY: This notice announces a public meeting of the Medicare Evidence Development Coverage Advisory Committee (MedCAC or Committee), formerly the Medicare Coverage Advisory Committee (MCAC). The Committee generally provides advice and recommendations about whether scientific evidence is adequate to determine whether certain medical items and services are reasonable and necessary under the Medicare statute. This meeting concerns percutaneous transluminal angioplasty (PTA) and stenting of the renal arteries.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. App. 2, section 10(a)).

DATES: *Meeting Date*: The public meeting will be held on Wednesday, July 18, 2007 from 7:30 a.m. until 4:30 p.m., eastern daylight time (e.d.t.).

Deadlines for Registration and Request for Special Accommodations: Registration must be completed no later than 5 p.m., e.d.t. on Monday, July 9, 2007. Request for special accommodations must be received by 5 p.m., e.d.t. Tuesday, July 10, 2007.

Deadlines for Written Comments and Presentations: Written comments and presentations must be received by June 18, 2007, 5 p.m., e.d.t. Presentations, once submitted, are final. No further changes to the presentation can be accepted after submission.

ADDRESSES: *Meeting Location*: The meeting will be held in the main auditorium of the Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Baltimore, MD 21244.

Presentation and Comment Submission: Interested persons may present data, information, or views orally or in writing on issues pending before the Committee. Presentation and written comments must be submitted by e-mail to *Michelle.Atkinson@cms.hhs.gov* or by regular mail to Michelle Atkinson, Executive Secretary for MedCAC, Centers for Medicare & Medicaid Services, Office of Clinical Standards and Quality, Coverage and Analysis Group, C1–09–06, 7500 Security Boulevard, Baltimore, MD 21244.

Registration and Special Accommodations: Individuals wishing to participate or who need special accommodations, or both, may register by phone or e-mail by contacting Maria Ellis at 410–786–0309 or *Maria.Ellis@cms.hhs.gov* no later than 5 p.m., e.d.t on Monday, July 9, 2007. Persons attending the meeting who are hearing or visually impaired, or have a condition that requires special assistance or accommodations, are asked to contact Michelle Atkinson, Executive Secretary for MedCAC, no later than July 9, 2007.

Web site: You may access up-to-date information on this meeting at *http://www.cms.hhs.gov/FACA/02_MCAC.asp#TopOfPage.*

FOR FURTHER INFORMATION CONTACT: Michelle Atkinson, Executive Secretary for MedCAC, 410–786–2881; *Michelle.Atkinson@cms.hhs.gov*; Centers for Medicare & Medicaid Services, OCSQ—Coverage and Analysis Group, C1–09–06, 7500 Security Boulevard, Baltimore, MD 21244).

SUPPLEMENTARY INFORMATION:

I. Background

On December 14, 1998, we published a notice in the **Federal Register** (63 FR 68780) to describe the Medicare Coverage Advisory Committee (MCAC), which provides advice and recommendations to CMS about clinical issues.

This notice announces the July 18, 2007 public meeting of the Committee. During this meeting, the Committee will discuss evidence and hear presentations from the public concerning the use of PTA and stenting of the renal arteries for the treatment of atherosclerotic renal artery stenosis (ARAS). The clinical outcomes in the Medicare population will be discussed. MedCAC will review the following kinds of evidence:

- The most informative measures of clinical outcomes.
- Indications.

- Clinical outcomes for the different treatment options.
- Complications.
- Harms and adverse events.
- Persistence of benefits and harms over time.
- Generalizability to the Medicare population in routine practice.

In addition to evaluating the available data, the Committee will identify areas in which the current data are deficient and in which additional research is warranted. Background information about this topic, including panel materials, is available at *http://www.cms.hhs.gov/coverage*.

II. Meeting Procedures

This meeting is open to the public. The Committee will hear oral presentations from the public for approximately 45 minutes. The Committee may limit the number and duration of oral presentations to the time available. If you wish to make formal presentations, you must notify the Executive Secretary for MCAC (see **FOR FURTHER INFORMATION CONTACT**) and submit the following to the address listed in the **ADDRESSES** section of this notice by the date listed in the **DATES** section of this notice: (1) A brief statement of the general nature of the evidence or arguments you wish to present; (2) the names and addresses of proposed participants; and (3) a written copy of your presentation. Your presentation should consider the questions we have posed to the Committee and focus on the issues specific to the topic.

The questions will be available on the following Web site: *http://www.cms.hhs.gov/FACA/02_MCAC.asp#TopOfPage* We require that you declare at the meeting whether you have any financial involvement with manufacturers of any items or services being discussed (or with their competitors).

After the public and CMS presentations, the Committee will deliberate openly on the topic. Interested persons may observe the deliberations, but the Committee will not hear further comments during this time except at the request of the chairperson. The Committee will also allow a 15 minute unscheduled open public session for any attendee to address issues specific to the topic. At the conclusion of the day, the members will vote and the Committee will make its recommendation.

III. Registration Instructions

The Coverage and Analysis Group is coordinating meeting registration. While there is no registration fee, individuals must register to attend. Register by contacting Maria Ellis by phone or e-mail as specified in the **ADDRESSES** section. Please provide your full name (as it appears on your State-issued driver's license), address, organization, telephone, fax number(s), and e-mail address. You will receive a registration confirmation with instructions for your arrival at the CMS complex.

IV. Security, Building, and Parking Guidelines

Because this meeting will be located on Federal property, for security reasons, any persons wishing to attend this meeting must register by the dates specified in the **DATES** section. Individuals who have not registered in advance will not be allowed to enter the building to attend the meeting.

The on-site check-in for visitors will begin at 7 a.m. Please allow sufficient time to go through the security checkpoints at both the entrance to the grounds and the entrance to the building.

Security measures also include a full inspection of vehicles, inside and exterior areas (rear, trunk, and engine) at the entrance to the grounds. In addition, all individuals entering the building must pass through a metal detector. All items brought to CMS, whether personal or for the purpose of or support of a demonstration, are subject to inspection. We cannot assume responsibility for coordinating the receipt, transfer, transport, storage, set-up, safety, or timely arrival of any personal belongings or items used for a demonstration or to support a demonstration.

Parking permits and instructions will be issued upon arrival.

Authority: 5 U.S.C. App. 2, section 10(a).

Dated: May 8, 2007.

Barry M. Straube,

Chief Medical Officer and Director, Office of Clinical Standards and Quality.

[FR Doc. E7–9780 Filed 5–24–07; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0200]

Agency Information Collection Activities; Proposed Collection; Comment Request; Health and Diet Survey

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on FDA's Health and Diet Survey.

DATES: Submit written or electronic comments on the collection of information by July 24, 2007.

ADDRESSES: Submit electronic comments on the collection of information to: *http://www.fda.gov/dockets/ecomments*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's