

Metropolitan Statistical Areas (MSAs) that span more than one State should be included in a single region. Furthermore, the conference report suggests that the required market study determine the best configuration of regions to maximize plan participation as well as the availability of plans to beneficiaries.

These statutory requirements and MMA conference report guidelines have several implications for the definition of MA regional areas. Geographic regions must be defined to meet multiple objectives and satisfy multiple constraints. Demographic data on the distribution of the aged population must be considered in conjunction with market factors that would impact insurance-supplier response. Incentives provided for in the legislation have the potential to offset unfavorable factors in the MA region and must also be considered in the analysis of these heterogeneous regions. In addition, the sizes and configuration of regions will themselves impact the entry behavior of plans.

B. Regional Definition for PDPs

Title I of the MMA establishes a prescription drug insurance benefit under a new Part D of Medicare and is intended to provide prescription drug coverage for beneficiaries enrolled in traditional Medicare FFS or MA plans. The law also provides for premium, deductible, and co-payment subsidies for certain low-income beneficiaries. The PDPs are effective in 2006.

To provide access to options for Medicare beneficiaries in all geographic areas, Medicare PDPs are intended to be regional in scope. Since private companies (with a public subsidy) will operate the PDPs, offering a plan in a region will be voluntary on the part of the plan operators. A plan must offer the same benefits and charge the same premiums and co-payments to all eligible beneficiaries in its region regardless of how the plan's costs vary within a region. If less than two full-risk plans are offered in a region (one of which must be a PDP), then we will approve any reduced risk plans that have applied to serve the region. In any regions or parts of regions that still lack two plans, we will arrange for a non-risk-bearing fallback plan to be offered.

The success of the Part D benefit will depend on the willingness of private plan operators to offer plans in the various regions and therefore, at least in part, on the region definitions selected by CMS. Implications for regional definition for PDPs include the trade-off of conforming to existing markets versus

encouraging plan choice in areas projected to be underserved.

The MMA mandates that there be between 10 and 50 PDP regions. In addition, we will establish regions for the territories as required in section 1860D–11(a)(2)(C) of the Act. We must define these regions by January 1, 2005. The legislative guidelines for the definition of regions are the same for regional MA plans. The MMA requires that PDP regions be the same as with MA regions “to the extent practicable.” However, the PDP regions do not necessarily need to be identical to the MA regions if it can be shown that a different configuration of regions for PDPs improves beneficiaries’ access to prescription drugs.

II. Meeting Topics and Format

The meeting will address the following topics:

- A presentation of proposed regional definitions for MA Regional Plans, followed by public comments and a question and answer period.
- A presentation of proposed regional definitions for PDPs, followed by public comments and a question and answer period.

Time for participants to ask questions or offer individual comments will be limited according to the number of registered participants.

The agenda will include presentations by CMS and RTI International (CMS’ contractor) staff. We are interested in an open dialogue on the topic of defining regions for regional MA plans and PDPs under the MMA legislation, and believe that an active discussion will help us more clearly identify the key issues for consideration. In this public meeting, we plan to engage in a discussion of the scenarios for MA regional and PDP region configurations, particularly on regional scenarios where PDP and regional MA definitions may, or may not, overlap.

III. Registration

Registration for this public meeting is required and will be on a first-come, first-served basis, limited to two attendees per organization up to the 1,000 person capacity of the meeting room. A waiting list will be available for additional requests. The registration deadline will be July 14, 2004. Registration can be accomplished through three mechanisms:

1. A special on-line meeting Web site set up specifically for this meeting: <https://register.rti.org/medicaremeeting/>.
2. A specific meeting e-mail address: medicaremeeting@rti.org.

3. By contacting Nathan West, RTI International, at (919) 485–2661.

A confirmation notice will be sent to attendees upon finalization of registration. Information on hotel accommodations will be provided to registered individuals as part of their confirmation notice. General information regarding meeting logistics will also be available on the meeting Web site at <https://register.rti.org/medicaremeeting/>.

Persons who are not registered in advance will not be guaranteed attendance due to space limitations. Attendees will be provided with meeting materials at the time of the meeting.

To submit written questions regarding logistics of the meeting or to requests material before the meeting, see instructions for *Written Statements and Requests* under the **ADDRESSES** section of this notice.

Written public comments are preferred following the meeting and will be accepted until August 5, 2004. See instructions for *Public Comments* under the **ADDRESSES** section of the notice.

(Authority: Sections 1851 through 1859 of the Social Security Act (42 U.S.C. 1395w–21 through 1395w–28)) (Catalog of Federal Domestic Assistance Program No. 93.773 Medicare—Hospital Insurance Program; and No. 93.774, Medicare—Supplementary Medical Insurance Program)

Dated: May 19, 2004.

Mark B. McClellan,

Administrator, Centers for Medicare & Medicaid Services.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CM–3130–N]

Medicare Program; Meeting of the Medicare Coverage Advisory Committee—July 14, 2004

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

ACTION: Notice of meeting.

SUMMARY: This notice announces a public meeting of the Medicare Coverage Advisory Committee (MCAC). This Committee 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. The Committee will discuss and make

recommendations regarding using transmyocardial revascularization (TMR) and percutaneous myocardial revascularization (PMR) to treat severe angina.

Notice is given under the Federal Advisory Committee Act (5 U.S.C. App. 2, section 10(a)(1) and (a)(2)).

DATES: *The Meeting:* The public meeting will be held on Wednesday, July 14, 2004 from 7:30 a.m. until 3:30 p.m. e.d.t., at the Holiday Inn Inner Harbor, 301 West Lombard Street, Baltimore, MD 21201.

Special Accommodations: For anyone attending the meeting who is hearing or visually impaired, or who requires special assistance or accommodations, please notify the Executive Secretary by June 25, 2004 (see **FOR FURTHER INFORMATION CONTACT**).

Presentations and Comments: Interested persons may present data, information, or views orally or in writing on issues pending before the Committee. Please submit written comments to Michelle Atkinson, by email at matkinson@cms.hhs.gov or by mail to the Executive Secretary, Office of Clinical Standards and Quality, Centers for Medicare & Medicaid Services, 7500 Security Boulevard, Mail Stop C1-09-06, Baltimore, MD 21244.

Deadline for Presentations and Comments: Written comments must be received by June 25, 2004, 5 p.m., e.d.t.

Web site: You may access up-to-date information on this meeting at www.cms.gov/coverage.

FOR FURTHER INFORMATION CONTACT: Michelle Atkinson, Executive Secretary, by telephone at 410-786-2881 or by e-mail at matkinson@cms.hhs.gov.

SUPPLEMENTARY INFORMATION: On December 14, 1998, we published a notice in the **Federal Register** (63 FR 68780) to describe the Medicare Coverage Advisory Committee, which provides advice and recommendations to us about clinical issues. This notice announces a public meeting of the Committee.

Meeting Topic: The Committee will discuss the evidence, hear presentations and public comment, and make recommendations regarding the use of transmyocardial revascularization (TMR) and percutaneous myocardial revascularization (PMR) for treatment of severe angina. TMR is a surgical technique that uses a laser to bore holes through the myocardium of the heart in an attempt to restore perfusion to areas of the heart not being reached due to diseased or clogged arteries; PMR is a subset of this technique which is less invasive and is used as a late or last resort to relieve symptoms of severe

angina in patients suffering ischemic heart disease who are not amenable to direct coronary revascularization interventions such as angioplasty, stenting, or open coronary bypass. Background information about this topic, including panel materials, is available on the Internet at <http://www.cms.gov/coverage>.

Procedure: 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 named in the **FOR FURTHER INFORMATION CONTACT** section of this notice, and submit the following by June 25, 2004, 5 p.m., e.d.t.: A brief statement of the general nature of the evidence or arguments you wish to present, and the names and addresses of proposed participants. A written copy of your presentation must be provided to each Committee member before offering your public comments. Your presentation must address the questions asked by CMS to the Committee. If the specific questions are not addressed your presentation will not be accepted. The questions will be available on the CMS Web site at <http://www.cms.gov/coverage/mcac>. We request that you declare at the meeting whether or not 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.

Authority: 5 U.S.C. App. 2, section 10(a)(1) and (a)(2).

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

Dated: May 6, 2004.

Sean R. Tunis,

Director, Office of Clinical Standards and Quality, Chief Medical Officer, Centers for Medicare & Medicaid Services.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Assets for Independence Demonstration Program

Agency: Administration for Children and Families (ACF), Office of Community Services (OCS).

Funding Opportunity Title: Assets for Independence Demonstration Program.

Announcement Type: Competitive Grant-Initial.

Funding Opportunity Number: HHS-2004-ACF-OCS-EI-0027.

CFDA Number: 93.602.

Due Dates for Applications: July 27, 2004.

I. Funding Opportunity Description

The Administration for Children and Families, Office of Community Services (OCS) will accept applications from organizations seeking financial assistance to establish and administer Assets for Independence (AFI) Projects. These projects are designed to assist low-income people in becoming economically self-sufficient. They do so by helping clients learn about economic and consumer issues and establish matched savings accounts called Individual Development Accounts (IDA) in order to save for a first home, a business or higher education. Grant recipient organizations (grantees) will be required to use a portion of the Federal financial assistance to support information collection and other activities related to an on-going national evaluation of the impact of AFI Projects and IDAs.

Grantees must comply with requirements in this program's authorizing legislation, the Assets for Independence Act (AFIA) (Title IV of the Community Opportunities, Accountability, and Training and Educational Services Act of 1998, as amended, Pub. L. 105-285, 42 U.S.C. 604 note). A copy of the Act is available at <http://www.acf.hhs.gov/assetbuilding/>

Program Purpose and Scope

The purpose of the Assets for Independence Program is to demonstrate and evaluate the effectiveness of asset-building projects that teach low-income families about financial issues and enable them to save earned income over the long-term in special matched savings accounts called Individual Development Accounts (IDA). The program is designed specifically to demonstrate and evaluate the effects of IDAs generally and AFI