DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services

[Document Identifier: CMS–10029 and CMS–R–254]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare and Medicaid Services, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections 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.

1. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* Medicare Program Integrity Customer Service Project; *Form No.:* CMS–10029 (OMB# 0938–0837); *Use:* Medicare's Integrity Program seeks to improve customer service provided to beneficiaries and providers. The study's purpose is to identify baseline satisfaction with Program Integrity efforts, to prioritize improvement areas, and to identify potential service delivery changes that can be implemented by CMS or its contractors. Respondents include beneficiaries whose billing questions were transferred to Fraud, and providers who have been through enrollment, medical review, or cost report audit; *Frequency:* Annually; *Affected Public:* Individuals or households, Business or other for-profit, Not-for-profit institutions; *Number of Respondents:* 5,250; *Total Annual Responses:* 5,250; *Total Annual Hours:* 782.

2. *Type of Information Collection Request:* Extension of a currently approved collection; *Title of Information Collection:* National Medicare Education Program (NMEP) Community Survey of Medicare Beneficiaries; *Form No.:* CMS–R–254 (OMB# 0938–0738); *Use:* A survey of Medicare beneficiaries in six communities will be conducted in January and February, 2003. A random, representative sample of Medicare beneficiaries will be selected using CMS administrative data. This approach will gather information on changes in: awareness of Medicare+Choice expansion and options; knowledge about Medicare and Medicare+Choice options; where beneficiaries go to find more information; and whether they are aware of the many information resources available to them; and satisfaction with their information/knowledge; *Frequency:* On occasion; *Affected Public:* Individuals or households; *Number of Respondents:* 2400; *Total Annual Responses:* 2400; *Total Annual Hours:* 600.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web site address at *http://www.hcfa.gov/regs/prdact95.htm,* or e-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov,* or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: September 12, 2002.

John P. Burke III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development and Issuances.

[FR Doc. 02–23977 Filed 9–19–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Radiological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Radiological Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on October 16, 2002, from 8:30 a.m. to 5 p.m.

Location: Holiday Inn, Two Montgomery Village Ave., Gaithersburg, MD.

Contact Person: Robert J. Doyle, Center for Devices and Radiological Health (HFZ–470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–1212, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12526. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss, make recommendations, and vote on a premarket approval application for a device that produces a computerized thermal image of the breast of women recommended for biopsy. Background information, including the agenda and questions for the committee, will be available to the public 1 business day before the meeting, on the Internet at http://www.fda.gov/cdrh/panelmtg.html. Material for the October 16, 2002, meeting will be posted on October 15, 2002.

Procedure: On October 16, 2002, from 8:30 a.m. to 1 p.m., and from 1:30 p.m. to 5 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by October 7, 2002. Oral presentations from the public will be scheduled between approximately 9:15 a.m. and 9:45 a.m., and for an additional 30 minutes near the end of the committee deliberations. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before October 9, 2002, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On October 16, 2002, from 1 p.m. to 1:30 p.m., the meeting will be closed to the public to permit FDA to present to the