

addition, CMS has established policies to limit payment abuse that will be based on FIs tracking patient movement among these co-located providers. *Form Number:* CMS-10088 (OMB#: 0938-0897; *Frequency:* Reporting—as needed; *Affected Public:* Business or other for profit and Not-for-profit institutions; *Number of Respondents:* 200; *Total Annual Responses:* 200; *Total Annual Hours:* 50.

2. *Type of Information Collection Request:* Revision of a currently approved collection; *Title of Information Collection:* Conditions of Coverage for Organ Procurement Organizations (OPOs) and Supporting Regulations in 42 CFR 486.301-348; *Use:* Organ Procurement Organizations are required to submit accurate data to CMS through the Organ Procurement and Transplantation Network (OPTN). The data concerns the organ procurement activities, as well as various OPO business activities, including information on its designated service area; structure; various policies, procedures, and protocols; and its quality assessment and performance improvement (QAPI) program. This information is necessary to assure maximum effectiveness in the procurement and distribution of organs. *Form Number:* CMS-R-13 (OMB#: 0938-0688; *Frequency:* Reporting—Every 4 years and as needed; *Affected Public:* Not-for-profit institutions; *Number of Respondents:* 58; *Total Annual Responses:* 58; *Total Annual Hours:* 21,427.

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.cms.hhs.gov/PaperworkReductionActof1995>, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786-1326.

Written comments and recommendations for the proposed information collections must be mailed or faxed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Carolyn Lovett, New Executive Office Building, Room 10235, Washington, DC 20503, Fax Number: (202) 395-6974.

Dated: November 7, 2006.

Michelle Shortt,

*Director, Regulations Development Group,
Office of Strategic Operations and Regulatory Affairs.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS-R-235]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Centers for Medicare & 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 & Medicaid Services (CMS) 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:* Data Use Agreement Information Collection Requirements, Model Language and Supporting Regulations in 45 CFR part 5b. *Use:* The Data Use Agreement (DUA) is needed as part of the review of each CMS data request to ensure compliance with the requirements of the Privacy Act for disclosure of data that contain individually-identifiable information. In addition, the DUA is used to maintain appropriate accounting and tracking of disclosures of records from Privacy Act systems of records. *Form Number:* CMS-R-235 (OMB#: 0938-0734); *Frequency:* Reporting-On occasion; *Affected Public:* Not-for-profit institutions; *Number of Respondents:* 1,500; *Total Annual Responses:* 1,500; *Total Annual Hours:* 750.

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.cms.hhs.gov/PaperworkReductionActof1995>, or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to Paperwork@cms.hhs.gov, or call the Reports Clearance Office on (410) 786-1326.

To be assured consideration, comments and recommendations for the proposed information collections must be received at the address below, no later than 5 p.m. on *January 16, 2007*.

CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development—B, Attention: William N. Parham, III, Room C4-26-05, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: November 7, 2006.

Michelle Shortt,

*Director, Regulations Development Group,
Office of Strategic Operations and Regulatory Affairs.*

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Clinical Chemistry and Clinical Toxicology 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). The meeting will be open to the public.

Name of Committee: Clinical Chemistry and Clinical Toxicology 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 December 6, 2006, from 8 a.m. to 4:30 p.m.

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

Contact Person: Veronica J. Calvin, Center for Devices and Radiological Health (HFZ-440), Food and Drug Administration, 2098 Gaither Rd.,