Information Collection: Application for Participation in the Medicare Care Management Performance Demonstration; *Form Number:* CMS–10165 (OMB#: 0938–0965); *Use:* The Medicare Care Management Performance (MCMP) Demonstration and its corresponding Report to Congress are mandated by the section 649 of the Medicare Prescription Drug, Improvement, and Modernization Act of 2003 (MMA). Section 649 of the MMA provides for the implementation of a ''pay for performance'' demonstration under which Medicare would pay incentive payments to physicians who (1) adopt and use health information technology; and (2) meet established standards on clinical performance measures. This demonstration will be held in four states, Arkansas, California, Massachusetts, and Utah. Providers that are enrolled in the Doctors' Office Quality—Information Technology (DOQ–IT) project are eligible to participate in the demonstration. To enroll in the MCMP Demonstration, a physician/provider must submit an application form. The information collected will be used to assess eligibility for the demonstration; *Frequency:* Reporting—One-time only; *Affected Public:* Business or other for-profit; *Number of Respondents:* 800; *Total Annual Responses:* 800; Total Annual Hours: 133.

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/regulations/pra/*, 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 December 13, 2005. CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Melissa Musotto, Room C4–26–05, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: October 6, 2005.

Michelle Shortt,

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

[FR Doc. 05–20517 Filed 10–13–05; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10064]

Agency Information Collection Activities: Submission for OMB Review; 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), 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 function; (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:* Minimum Data Set (MDS) for Swing Bed Hospitals and Supporting Regulations in 42 CFR 483.20 and 413.337; *Form No.:* CMS–10064 (OMB # 0938–0872); *Use:* As required under Section 1888(e)(7) of the Social Security Act, swing bed hospitals must be reimbursed under the skilled nursing facility prospective payment system. CMS uses the MDS data to reimburse swing bed hospitals for SNF-level care furnished to Medicare beneficiaries. The MDS3.0 is currently being developed with plans for field testing to begin in 2006 with the expectation of completion in 2007. At that time, CMS will analyze the data derived from the study, including the implementation of the new version of the MDS for swing bed hospitals. Since we do not have the MDS3.0 version available, we are requesting an extension for the current SB–MDS.; *Frequency:* Reporting—Other (days 5, 14, 30, 60, and 90 of stay); *Affected Public:* Not-for-profit institutions, and State, Local, and Tribal governments; *Number of Respondents:* 820; *Total Annual Responses:* 92,789; *Total Annual Hours:* 51,314.

To obtain copies of the supporting statement and any related forms for these paperwork collections referenced above, access CMS Web site address at *http://www.cms.hhs.gov/regulations/pra/*, 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 by the OMB Desk Officer at the address below, no later than 5 p.m. on November 14, 2005. OMB Human Resources and Housing Branch, Attention: CMS Desk Officer, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: October 6, 2005.

Martique S. Jones,

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

[FR Doc. 05–20521 Filed 10–13–05; 8:45 am]

BILLING CODE 4120–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Blood Products 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: Blood Products 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 November 3, 2005, from 8 a.m. to 5:30 p.m., and on November 4, 2005, from 8 a.m. to 3:30 p.m.

Location: Holiday Inn Gaithersburg, 2 Montgomery Village Ave., Gaithersburg, MD 20879.

Contact Person: Donald W. Jehn or Pearline K. Muckelvene, Center for Biologics Evaluation and Research, Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852, 301–827–0314, or FDA Advisory Committee Information Line, 1–800–