

in 42 CFR 484.1–484.52; *Form No.*: HCFA–1515/1572 (OMB# 0938–0355); *Use*: In order to participate in the Medicare program as a Home Health Agency (HHA) provider, the HHA must meet Federal Standards. These forms are used to record information about patients' health and provider compliance with requirements; *Frequency*: Annually; *Affected Public*: Business or other for-profit, Not-for-profit institutions; *Number of Respondents*: 6,997; *Total Annual Responses*: 13,994; *Total Annual Hours*: 13,994.

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: Allison Eydt, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: December 6, 2001.

John P. Burke, III,

CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–2449 Filed 1–31–02; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Center for Medicare and Medicaid Services

[Document Identifier: CMS–R–306]

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

AGENCY: Center 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 Center 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.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* Restraint and Seclusion Standards for Psychiatric Residential Treatment Facilities; *Form No.:* CMS–R–305 (OMB# 0938–0786); *Use:* Psychiatric residential treatment facilities are required to report deaths, serious injuries and attempted suicides to State Medicaid Agency and Protection and Advocacy Organization. They are also required to provide residents restraint and seclusion policy in writing and to document resident record of all activities involving use of restraint and seclusion; *Frequency:* On occasion; *Affected Public:* Business or other for-profit, Not for profit institutions; *Number of Respondents:* 500; *Total Annual Responses:* 2,600,000; *Total Annual Hours:* 877,750.

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.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: December 6, 2001.

John P. Burke, III,

CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Center for Medicare and Medicaid Services

[Document Identifier: CMS–R–200]

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

AGENCY: Center 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 Center 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.

Type of Information Collection Request: Reinstatement, with change, of a previously approved collection for which approval has expired; *Title of Information Collection:* Health Plan Employer Data and Information Set (HEDIS) and Health Outcome Survey (HOS) and Supporting Regulations in 42 CFR 422.152; *Form No.:* CMS–R–200 (OMB# 0938–0701); *Use:* The Centers for Medicare & Medicaid Services (formerly HCFA) collects quality performance measures in order to hold the Medicare managed care industry accountable for the care being delivered, to enable quality improvement, and to provide quality information to Medicare beneficiaries in order to promote an informed choice. It is critical to CMS's mission that we collect and disseminate information that will help beneficiaries choose among health plans, contribute to improved quality of care through identification of improvement opportunities, and assist CMS in carrying out its oversight and purchasing responsibilities; *Frequency:* Annually; *Affected Public:* Business or other for-profit, Not-for-profit institutions, Individuals or households;

Number of Respondents: 313,825; *Total Annual Responses:* 313,825; *Total Annual Hours:* 571,488.

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: Allison Eydt, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: January 23, 2002.

John P. Burke, III,

CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Center for Medicare and Medicaid Services

[Document Identifier: CMS-10044]

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

AGENCY: Center 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 Center 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.

Type of Information Collection Request: New Collection; *Title of Information Collection:* Medicare Lifestyle Modification Program Demonstration; *Form No.:* CMS-10044 (OMB# 0938-NEW); *Use:* This demonstration will focus on two Medicare sponsored, lifestyle modification programs designed to reverse, reduce or ameliorate the progression of coronary artery disease (CAD) at risk for significant morbidity and mortality. This demonstration will test the cost-effectiveness and feasibility of providing payment for cardiovascular lifestyle modification program services to Medicare beneficiaries.; *Frequency:* Baseline Enrollment, 12 and 24 months; *Affected Public:* Individuals or Households; *Number of Respondents:* 2,240; *Total Annual Responses:* 1,680; *Total Annual Hours:* 1,106.

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: Allison Eydt, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: January 24, 2002.

John P. Burke, III,

CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Best Practices for Reducing Transfusion Errors; Public Workshop

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public workshop.

The Food and Drug Administration (FDA) is announcing the following public workshop: Best Practices for Reducing Transfusion Errors. The

purpose of the public workshop is to discuss practices and techniques that may decrease transfusion errors, including systems and technology that can be applied to reducing transfusion errors.

Date and Time: The public workshop will be held on February 14, 2002, from 8:30 a.m. to 5 p.m., and February 15, 2002, from 8:30 a.m. to 12:30 p.m.

Location: The public workshop will be held at the Natcher Conference Center, National Institutes of Health, Bldg. 45, 45 Center Dr., 8600 Rockville Pike, Bethesda, MD.

Contact: Joseph Wilczek, Center for Biologics Evaluation and Research (HFM-302), Food and Drug Administration, 1401 Rockville Pike, Rockville, MD 20852-1448, 301-827-6129, FAX 301-827-2843.

Registration: Send registration information (including name, title, firm name, address, telephone, and fax number) to the contact person by February 5, 2002. Onsite registration on a space available basis will begin at 7:30 a.m. on the days of the workshop. There is no registration fee. If you need special accommodations due to a disability, please contact Joseph Wilczek at least 7 days in advance.

Transcripts: Transcripts of the public workshop may be requested in writing from the Freedom of Information Office (HFI-35), Food and Drug Administration, 5600 Fishers Lane, rm. 12A-16, Rockville, MD 20857, approximately 15 working days after the meeting at a cost of 10 cents per page. The public workshop transcript will also be available on the Internet at <http://www.fda.gov/cber/minutes/workshop-min.htm>.

SUPPLEMENTARY INFORMATION: FDA and the Agency for Healthcare Research and Quality, Department of Health and Human Services, are cosponsoring a public workshop on avoiding errors in transfusion medicine. On the first day of the workshop, topics to be discussed include: Patient and medication identification, errors in manufacturing and testing of blood and blood components, system errors and cultural factors, and the role of product deviation reporting in reducing transfusion errors. The second day of the workshop will address current as well as future technology trends that should help prevent transfusion errors. The public workshop agenda is posted on the Internet at <http://www.fda.gov/cber/meetings/trnfsnerr021402.htm>.