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:* Conditions of Participation for Portable X-ray Suppliers and Supporting Regulations in 42 CFR, Sections 486.104, 486.106, and 486.110; *Use:* This information collection request contains the recordkeeping requirements contained in the above noted regulation sections. These requirements are designed to ensure that each supplier has a properly trained staff to provide the appropriate type and level of care, as well as a safe physical environment for patients. CMS uses these conditions to certify portable X-ray Suppliers wishing to participate in the Medicare program.; *Form Number:* CMS–R–43 (OMB#: 0938–0338); *Frequency:* Recordkeeping; *Affected Public:* Business or other for-profit and Not-for-profit institutions; *Number of Respondents:* 602; *Total Annual Responses:* 602; *Total Annual Hours:* 1,505.

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.

Written comments and recommendations for the proposed information collections must be mailed within 60 days of this notice directly to the CMS Paperwork Reduction Act Reports Clearance Officer designated at the address below: CMS, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development, Attention: Melissa Musotto, Room C5–14–03, 7500 Security Boulevard, Baltimore, Maryland 21244–1850.

Dated: February 3, 2005.

John P. Burke, III,

CMS Paperwork Reduction Act Reports Clearance Officer, Office of Strategic Operations and Regulatory Affairs, Regulations Development Group.

[FR Doc. 05–2658 Filed 2–10–05; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[Document Identifier: CMS–10131]

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.

Type of Information Collection Request: New Collection; *Title of Information Collection:* Evaluation of Medicare Disease Management Demonstrations; *Form No.:* CMS–10131 (OMB# 0938–NEW); *Use:* CMS contracted with Mathematica Policy Research, Inc. for the evaluation of programs and disease management. The purpose of the patient survey is to assess the impact of disease management and prescription drug benefits (the latter in 3 of the sites) on patient's health and functioning status, care satisfaction, health behaviors and knowledge of condition. Data from the physician survey will be used to assess physician satisfaction with disease management services, their perceptions of the impact of disease management on patient outcomes, education, and service use, and on their own practice and office workload.; *Frequency:* On Occasion; *Affected Public:* Individuals or households, Business or other for-profit; *Number of Respondents:* 5000; *Total Annual Responses:* 2500; *Total Annual Hours:* 1625.

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.

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: Christopher Martin, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: February 3, 2005.

John P. Burke, III,

CMS Paperwork Reduction Act Reports Clearance Officer, Office of Strategic Operations and Regulatory Affairs, Regulations Development Group.

[FR Doc. 05–2659 Filed 2–10–05; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 1998N–0046]

Annual Comprehensive List of Guidance Documents at the Food and Drug Administration; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of January 5, 2005 (70 FR 824). The document provided the agency's annual comprehensive list of guidance documents. The list provided information on current guidance documents and those that have been withdrawn. The document was published with some inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Joyce Strong, Office of Policy (HF–27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–7010.

SUPPLEMENTARY INFORMATION: In FR Doc. 05–155, appearing on page 824 in the **Federal Register** of Wednesday, January 5, 2005, the following corrections are made:

1. On page 867, in the list, under the heading ''Guidance Documents Issued by CDRH—Continued,'' the entire entry is removed for the document entitled ''Review of 510(k)s for Computer Controlled Medical Devices (blue book