restrictions on the extent to which a CDC-funded awardee can participate in or implement environmental changes within their respective communities. (See Section: Use of Funds.)

Potential applicants may obtain a copy of Healthy People 2000 (Full Report, Stock Number 017–001–00474– 0), or Healthy People 2000 (Summary Report, Stock Number 017–001–00473– 1), referenced in the **Introduction** through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325; telephone (202) 512–1800.

Dated: April 17, 1998.

Joseph R. Carter,

Acting Associate Director for Management and Operations, Centers for Disease Control and Prevention (CDC). [FR Doc. 98–10788 Filed 4–22–98; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N-0531]

Agency Information Collection Activities; Announcement of OMB Approval

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Performance Standard for Electrode Lead Wires and Patient Cables: Petitions for Exemptions and Variances" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: Margaret R. Schlosburg, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223. SUPPLEMENTARY INFORMATION: In the Federal Register of January 21, 1998 (63 FR 3141), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under section 3507 of the PRA (44 U.S.C. 3507). An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910-0367. The approval expires on April 30, 2001.

Dated: April 16, 1998. **William K. Hubbard,** *Associate Commissioner for Policy Coordination.* [FR Doc. 98–10778 Filed 4–22–98; 8:45 am] BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 97N-0485]

Agency Information Collection Activities; Announcement of OMB Approval

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Shipment of a Blood Product Prior to Completion of Testing for Hepatitis B Surface Antigen (HbsAg), and Shipment of Blood Products Known Reactive for HbsAg" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (the PRA).

FOR FURTHER INFORMATION CONTACT: JonnaLynn P. Capezzuto, Office of Information Resources Management (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In the Federal Register of December 19, 1997 (62 FR 66633), the agency announced that the proposed information collection had been submitted to OMB for review and clearance under section 3507 of the PRA (44 U.S.C. 3507). An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0168. The approval expires on April 30, 2001.

Dated: April 16, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination. [FR Doc. 98–10781 Filed 4–22–98; 8:45 am] BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98N-0046]

Comprehensive List of Current 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 document that published in the Federal Register of February 26, 1998 (63 FR 9795). The document provided a comprehensive list of all guidance documents currently in use at the agency. FDA committed to publishing this list in its February 1997 "Good Guidance Practices" (GGP's), which set forth the agency policies and procedures for the development, issuance, and use of guidance documents. The document was published with several errors. This document corrects those errors. DATES: General comments on this list and on agency guidance documents are

welcome at any time.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Lisa L. Barclay, Office of Policy (HF–22), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–3360.

In FR Doc. 98–4916, appearing on page 9795, in the **Federal Register** of Thursday, February 26, 1998, the following corrections are made:

1. On page 9795, in the second column, under the "ADDRESSES" caption, "(HFD–305)" is removed and "(HFA–305)" is added in its place.

2. On page 9834, in the fifth entry entitled "Clinical Testing of Infant Formulas with Respect to Nutritional Suitability for Term Infants," in the second column, "1985" is removed and "1988" is added in its place.

3. On page 9834, in the first column, the sixth entry entitled "Guidelines for the Evaluation of the Safety and Suitability of New Infant Formulas for Feeding Infants with Allergic Diseases" is removed and "Evaluation of Safety and Suitability of New Infant Formulas for Feeding Preterm Infants" is added in its place.

4. On page 9834, under the heading, "VI. Guidance Documents Issued by the Center for Veterinary Medicine (CVM)," in the first entry entitled "Citizen Petitions: Policy and Procedures (Guide No. 1240.2030)," in the fourth column, "Do" is removed and "Center for Veterinary Medicine (HFV–12), Communications Staff, Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1755" is added in its place.

5. On page 9842, in the fourth entry entitled "Guide to Inspections of Source Plasma Establishments (PB96–127360)," in the second column, "December 1994" is removed and "June 1997" is added in its place.

Dated: April 16, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination. [FR Doc. 98–10780 Filed 4–22–98; 8:45 am]

BILLING CODE 4160-01-F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA-R-193]

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, 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 the 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:* "An Important Message From Medicare." and Supporting Regulations 42 CFR 466.78, 489.27, .20; *Form No.:* HCFA–R–193, OMB # 0938–0692; *Use:* Hospitals participating in the Medicare program have agreed to distribute "An Important

Message from Medicare" to beneficiaries during each admission. Receiving this information will provide the beneficiary with some ability to participate and/or initiate discussions concerning discussions affecting Medicare coverage or payment and about his or her appeal rights in response to any hospitals notice to the effect that Medicare will no longer cover continued care in the hospital. Frequency: Other, as needed; Affected Public: Individuals or Households, Business or other for-profit, Not-for-profit, Federal Government, State, Local, or Tribal Government; Number of Respondents: 6,700; Total Annual Responses: 11,000,000; Total Annual Hours: 183,000.

To obtain copies of the supporting statement for the proposed paperwork collections referenced above, or any related forms, E-mail your request, including your address and phone number, 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 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards, Attention: John Rudolph, Room C2-26-17, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.

Dated: April 16, 1998.

John P. Burke III,

HCFA Reports Clearance Officer, Division of HCFA Enterprise Standards, Health Care Financing Administration. [FR Doc. 98–10829 Filed 4–22–98; 8:45 am] BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA-576]

Agency Information Collection Activities: Proposed Collection; Comment Request

AGENCY: Health Care Financing Administration, HHS.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, 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: Organ **Procurement Organization (OPO)** Request for Designation and Supporting Regulations in 42 CFR 486.301-486.325; Form No.: HCFA-576 (OMB# 0938-0512); Use: The information provided on this form serves as a basis for certifying OPOs for participation in the Medicare and Medicaid programs and will indicate whether the OPO is meeting the specified performance standards for reimbursement of service. Additionally, the form is used for inputting minimal information into the **Online Survey Certification Reporting** (OSCAR) System; Frequency: Annually; Affected Public: Business or other forprofit, and Not-for-profit institutions; Number of Respondents: 69; Total Annual Responses: 69; Total Annual Hours: 138.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access HCFA's Web Site address at http://www.hcfa.gov/ regs/prdact95.htm, or E-mail your request, including your address, phone number, OMB number, and HCFA 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 60 days of this notice directly to the HCFA Paperwork Clearance Officer designated at the following address: HCFA, Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards, Attention: Louis Blank, Room C2-26-17, 7500 Security Boulevard, Baltimore, Maryland 21244-1850.