7156). A ListServ is available to provide unmoderated discussion at bdgddd@listserv.cdc.gov

The BDGDDD Work Group will work to develop a set of objectives that represent a common ground for health professionals, persons with related health conditions and their family members, and interested others. ''Issues-of-interest'' will be identified that are barriers to common agreement and discussion is invited to help revise objectives to better reflect the common views of the Work Group.

Dated: February 10, 1998.

Joseph R. Carter,

Acting Associate Director for Management, and Operations Centers for Disease Control and Prevention (CDC).

[FR Doc. 98–3824 Filed 2–13–98; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 91N–0396]

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 ''Reports of Corrections and Removals for Manufacturers, Importers, and Distributors of Medical Devices (21 CFR 806.10 and 806.20)'' 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 November 26, 1997 (62 FR 63182), 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 0910–0359. The approval expires on January 31, 2001.

Dated: February 9, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–3777 Filed 2–13–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 94P–0240]

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 ''Food Labeling; Serving Sizes; Reference Amount for Baking Powder, Baking Soda, Pectin'' 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 November 18, 1997 (62 FR 61476), 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 0910–0357. The approval expires on January 31, 2001.

Dated: February 4, 1998.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 98–3901 Filed 2–13–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[Document Identifier: HCFA–R–108]

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

AGENCY: Health Care Financing Administration.

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:* Criteria for Medicare Coverage of Liver Transplants; Form No.: HCFA-R–108 (OMB# 0938–0580); *Use:* Medicare participating hospitals must file an application to be approved for coverage and payment of liver transplants performed on Medicare beneficiaries; *Frequency:* Annually; *Affected Public:* Business or other for-profit; *Number of Respondents:* 12; *Total Annual Responses:* 12; *Total Annual Hours:* 1,880.

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 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, D.C. 20503.

Dated: February 4, 1998.

John P. Burke III,

HCFA Reports Clearance Officer, HCFA Office of Information Services, Information Technology Investment Management Group, Division of HCFA Enterprise Standards.

[FR Doc. 98–3818 Filed 2–13–98; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Correction

AGENCY: Health Resources and Services Administration, HHS.

ACTION: Notice; correction.

SUMMARY: In the **Federal Register** issue of Thursday, October 9, 1997, make the following correction:

Correction

In FR Doc. 97–26645, on page 52908, in the third column under the heading ''Sudden Infant Death Syndrome (SIDS)/ Other Infant Death (OID) Program,'' the program is being withdrawn from competition due to financial and programmatic concerns.

Dated: February 10, 1998.

Claude Earl Fox,

Acting Administrator.

[FR Doc. 98–3832 Filed 2–13–98; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

National Practitioner Data Bank for Adverse Information on Physicians and other Health Care Practitioners: Availability of and Fee for Public Use Data File

The Health Resources and Services Administration (HRSA), Department of Health and Human Services (DHHS), is announcing a fee of $195 for the recently available public use data file which includes selected information from approximately 168,000 reports submitted to the National Practitioner Data Bank (Data Bank) between September 1, 1990 and December 31, 1997. HRSA plans to make updated versions of the complete file available every 4 months. A separate $195 fee will be charged for each updated copy of the file. The file contains information concerning: (1) Malpractice payments made for the benefit of physicians, dentists, and other health care practitioners; and (2) adverse licensure, clinical privileges, and professional society membership actions concerning physicians and dentists.

The file does not contain information which would allow identification of individual physicians, dentists, or other health care practitioners. It also does not contain information identifying either entities which filed reports with the Data Bank or patients. This information is being made available for research purposes in conformance with 42 USC 11137(b). Hospitals cannot fulfill their obligation under 42 USC 11135 to query the Data Bank by obtaining this data file. Other health care entities cannot fulfill obligations to query the Data Bank imposed by accreditation agencies by obtaining this file.

Information in the file includes type of practitioner, type of reporting entity, and the practitioner's State. For malpractice payment reports, information includes malpractice payment amount, reasons for malpractice payment, date of payment, and whether payment is a result of judgment or settlement. For adverse action reports, the file includes information on the reason for the licensure or clinical privileges adverse action, the type of action taken, and the duration of such action.

The public use file is in ASCII format and is approximately 20 megabytes in size. It is available in compressed form on IBM–PC compatible high density 3.5 inch diskettes and may also be made available in CD–ROM format. In addition to the data themselves, a complete file description in ASCII text format is included. For informaton on how to order the file, call Data Bank ''Help Line'' at 1–800–767–6732.

The Data Bank is authorized by the Health Care Quality Improvement Act of 1986 (the Act), title IV of Public Law 99–660, as amended (42 U.S.C. 11101 *et seq.*). Section 427(b)(4) of the Act authorizes the establishment of fees for the costs of processing requests for disclosure and of providing such information.

Final regulations at 45 CFR part 60 set forth the criteria and procedures for information to be reported to and disclosed by the Data Bank. Section 60.3 of these regulations defines the terms used in this announcement.

In determining any changes in the amount of the user fee, the Department uses the criteria set forth in § 60.12(b) of the regulations, as well as allowable costs pursuant to the DHHS Appropriations Act of 1998, Pub. L. 105–78, enacted November 13, 1997. This Act requires that the Department recover the full costs of operating the Data Bank through user fees. Section 60.12(b) of the regulations states:

''The amount of each fee will be determined based on the following criteria:

(1) Use of electronic data processing equipment to obtain information—the actual cost for the service, including computer search time, runs, printouts, and time of computer programmers and operators, or other employees,

(2) Photocopying or other forms of reproduction, such as magnetic tapes—actual cost of the operator's time, plus the cost of the machine time and the materials used,

(3) Postage—actual cost, and

(4) Sending information by special methods requested by the applicant, such as express mail or electronic transfer—the actual cost of the special service.''

Additionally, in establishing this charge, the Agency used guidance issued in the Office of Management and Budget (OMB) Circular A–25, applicable to the imposition of user fees. This circular authorizes agencies to collect user fees for the ''full cost'' of providing a service. These allowable costs include research. All other Data Bank user fees remain the same.

The Department will review this charge periodically, and will revise it as necessary. Any changes in the fee and their effective dates will be announced in the **Federal Register**.

Dated: February 10, 1998.

Claude Earl Fox,

Acting Administrator.

[FR Doc. 98–3833 Filed 2–13–98; 8:45 am]

BILLING CODE 4160–15–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; Notice of Closed Meeting

Pursuant to Section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting:

Name of Committee: National Human Genome Research Institute, Special Emphasis, Panel ZHG1 HGR P M1.

Agenda/Purpose: To review and evaluate grant applications and/or contract proposals.

Date: February 26, 1998.

Time: 8:30 am to 5 p.m.

Place: The Sheraton Washington Hotel, Washington, D.C.

Contact Person: Rudy Pozzatti, Ph.D., Office of Scientific Review, National Human Genome Research Institute, National Institutes of Health, Building 38A, Room 604, Bethesda, Maryland 20892, (301) 402–0838.