be kept in an active file for two years. New hire information will then be stored for an additional three years before being destroyed.

Tax refund and administrative offset information will be maintained for six years in an active master file for purposes of collection and adjustment. After this time, records of cases for which there was no collection will be destroyed. Records of cases with a collection will be stored on-line in an inactive master file.

Records pertaining to passport denial will be updated and/or deleted as obligors meet satisfactory restitution or other State approved arrangements.

Records of information provided by the FPLS to authorized users will be maintained only long enough to communicate the information to the appropriate State or Federal agent. Thereafter, the information provided will be destroyed. However, records pertaining to the disclosures, which include information provided by States, Federal agencies contacted, and an indication of the type(s) of information returned, will be stored on a history tape and in hard copy for five years and then destroyed.

SYSTEM MANAGER(S) AND ADDRESS:

Director, Division of Program Operations Office of Child Support Enforcement Administration for Children and Families 370 L'Enfant Promenade, SW., 4th Floor East Washington, DC. 20447.

NOTIFICATION PROCEDURES:

To determine if a record exists, write to the System Manager listed above. The requester must provide his or her full name and address. Additional information, such as your Social Security Number, date of birth or mother's maiden name, may be requested by the system manager in order to distinguish between individuals having the same or similar names.

RECORD ACCESS PROCEDURES:

Write to the System Manager specified above to attain access to records. Requesters should also reasonably specify the record contents they are seeking.

CONTESTING RECORD PROCEDURE:

Contact the official at the address specified under system manager above, and reasonably identify the record and specify the information to be contested and corrective action sought with supporting justification to show how the record is inaccurate, incomplete, untimely or irrelevant.

RECORD SOURCE CATEGORIES:

Information is obtained from departments, agencies, or instrumentalities of the United States or any State.

SYSTEMS EXEMPTED FROM CERTAIN PROVISIONS OF THE ACT:

None.

[FR Doc. 97–26049 Filed 10–1–97; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

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: Circulatory System Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on October 24, 1997, 9:30 a.m. to 6 p.m.

Location: Gaithersburg Marriott Washingtonian Center, Salons C and D, 9751 Washingtonian Blvd., Gaithersburg, MD.

Contact Person: John E. Stuhlmuller, Center for Devices and Radiological Health (HFZ–450), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–443–8243, ext. 157, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12625. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee is being asked to provide input to the agency regarding the design of clinical trials to support premarket approval applications for devices intended to treat atrial septal defects, patent foramen ovale, and patent ductus arteriosus. Of particular concern are the following issues: What are the appropriate controls to be used in such trials? What are the appropriate safety and efficacy measures? When should assessments of these measures be made?

Procedure: On October 24, 1997, from 12:30 p.m. to 6 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by October 14, 1997. Oral presentations from the public will be scheduled between approximately 12:30 p.m. and 1:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before October 14, 1997, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On October 24, 1997, from 9:30 a.m. to 12:30 p.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential information (5 U.S.C. 552b(c)(4)). FDA staff will present trade secret and/or confidential information regarding pending and future circulatory system device submissions.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: September 25, 1997.

Michael A. Friedman,

Deputy Commissioner for Operations.

[FR Doc. 97–26113 Filed 10–1–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Care Financing Administration

[HCFA–382]

Agency Information Collection Activities: Submission for OMB Review; 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