

The Department specifically requests comments on: (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated November 3, 1999.
Bob Sargis,
Acting Reports Clearance Officer.
 [FR Doc. 99-29240 Filed 11-8-99; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Title: 45 CFR Part 95, Supart F—Automatic Data Processing Equipment and Services—Conditions for Federal Financial Participation (FFP).
OMB No.: 0992-0005.
Description: The advance planning document (APD) process, established in the rules at 45 CFR Part 95, Subpart F, is the procedure by which States request

and obtain approval for Federal financial participation in their cost of acquiring automatic data processing equipment and services. The State Agency submitted APD, provides the Department of Health and Human Services (DHHS) with the following information necessary to determine the State's need to acquire the requested ADP equipment and/or services:

1. A statement of need;
2. A requirements analysis and feasibility study;
3. A cost benefits analysis;
4. A proposed activity schedule; and,
5. A proposed budget.

DHHS' determination, of a State agency's need to acquire requested ADP equipment or services, is authorized at sections 402(a)(5), 452(a)(1), 1902(a)(4) and 1102 of the Social Security Act.

Respondents: State, Local or Tribal Governments.

Annual Burden Estimates:

Instrument	Number of respondents	Number of respondents per respondent	Average burden hours per response	Total burden hours
Advance Planning Document	50	1.84	60	5,520
RFP and Contract	50	1.54	1.5	115.5
Emergency Funding Request	27	1	1	27
Service Agreement	14	1	1	14
Biennial Reports	50	1	1.5	75

Estimated Total Annual Burden Hours: 5,751.5.

In Compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above.

Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Gastroenterology and Urology Devices Panel of the Medical Devices 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: Gastroenterology and Urology Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and

recommendations to the agency on FDA's regulatory issues.

Date and Time: The meeting will be held on November 19, 1999, 8 a.m. to 5 p.m.

Location: Corporate Bldg., conference room 020B, 9200 Corporate Blvd., Rockville, MD.

Contact Person: Mary J. Cornelius, Center for Devices and Radiological Health (HFZ-470), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301-594-2194, ext. 118, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area), code 12523. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss, make recommendations, and vote on a premarket approval application for an optical device intended to assist in the evaluation of colonic polyps using laser-induced autofluorescence.

Procedure: On November 19, 1999, from 8:30 a.m. to 1:30 p.m., and from 2 p.m. to 5 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