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]
BILLING CODE 4184–01–M

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 bur- den hours per response	Total burden hours
Advance Planning Document RFP and Contract Emergency Funding Request Service Agreement Biennial Reports	50 50 27 14 50	1.84 1.54 1 1	60 1.5 1 1 1.5	5,520 115.5 27 14 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.

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 clarify 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. 99–29241 Filed 11–8–99; 8:45 am]
BILLING CODE 4184–01–M

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

person by November 15, 1999. Oral presentations from the public will be scheduled between approximately 9:15 a.m. and 9:45 a.m. and between approximately 3 p.m. and 3:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before November 15, 1999, 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 November 19, 1999, from 8 a.m. to 8:30 a.m., the meeting will be closed to permit discussion and review of trade secret and/or confidential commercial information regarding pending and future device issues (5 U.S.C. 552b(c)(4)). This portion of the meeting will be closed to permit discussion of this information.

FDA regrets that it was unable to publish this notice 15 days prior to the November 19, 1999, Gastroenterology and Urology Devices Panel of the Medical Devices Advisory Committee meeting. Because the agency believes there is some urgency to bring this issue to public discussion and qualified members of the Gastroenterology and Urology Devices Panel of the Medical Devices Advisory Committee were available at this time, the Commissioner concluded that it was in the public interest to hold this meeting even if there was not sufficient time for the customary 15-day public notice.

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

Dated: October 29, 1999.

Dennis E. Baker,

Associate Commissioner for Regulatory Affairs.

[FR Doc. 99–29223 Filed 11–8–99; 8:45 am] BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Substance Abuse Prevention; Notice of Meeting

Pursuant to Pub. L. 92–463, notice is hereby given of the meeting of the Drug Testing Advisory Board of the Center for Substance Abuse Prevention in December 1999.

The Drug Testing Advisory Board meeting will be open from 8:30 a.m.

until 10:15 a.m. on December 8, 1999. The open session will include a roll call, an HHS update, a DOT update, a brief review of the on-site testing meeting held on October 5 and 6, and a brief review of the working group meeting held on November 3, 4, and 5 to discuss alternative specimen testing. A public comment period will be scheduled during the open session. If anyone needs special accommodations for persons with disabilities please notify the Contact listed below.

The Drug Testing Advisory Board meeting will be closed from 10:15 a.m. on December 8, 1999, until 3:30 p.m. on December 9, 1999, because it involves the review of sensitive National Laboratory Certification Program (NLCP) internal operating procedures and program development issues. Therefore, this portion of the meeting will be closed to the public as determined by the Administrator, SAMHSA, in accordance with 5 U.S.C. 552b(c)(2), (4), and (6) and 5 U.S.C. App.2, § 10(d).

An agenda for the open session of this meeting and a roster of board members may be obtained from: Mrs. Giselle Hersh, Division of Workplace Programs, 5600 Fishers Lane, Rockwall II, Suite 815, Rockville, MD 20857, Telephone: (301) 443–6014.

Substantive program information may be obtained from the contact whose name and telephone number is listed below.

Committee Name: Drug Testing Advisory Board.

Meeting Date: December 8-9, 1999.

Place: Holiday Inn, 5520 Wisconsin Avenue, Chevy Chase, Maryland 20815.

Open: December 8, 1999; 8:30 a.m.–10:15 a.m.

Closed: December 8, 1999; 10:15 a.m.– 3:30 p.m.

Closed: December 9, 1999; 8:30 a.m.–3:30 p.m.

Contact: Donna M. Bush, Ph.D., Executive Secretary, Telephone: (301) 443–6014 and FAX: (301) 443–3031.

Dated: November 3, 1999.

Sandra E. Stephens,

Committee Management Officer, Substance Abuse and Mental Health Services Administration.

[FR Doc. 99-29225 Filed 11-8-99; 8:45 am] BILLING CODE 4162-20-P

DEPARTMENT OF THE INTERIOR

Bureau of Land Management [ID-933-1430-AG; IDI-017246]

Termination of Recreation and Public Purpose Act Classification and

Opening Order, Idaho
AGENCY: Bureau of Land Management,

Interior.

ACTION: Notice

SUMMARY: This notice terminates a Recreation and Public Purpose Act Classification on 43.81 acres as this classification is no longer needed. **EFFECTIVE DATE:** November 9, 1999.

FOR FURTHER INFORMATION CONTACT: Catherine D. Foster, BLM Idaho State Office, 1387 S. Vinnell Way, Boise, Idaho 83709, 208–373–3863.

SUPPLEMENTARY INFORMATION: On June 3, 1971, 43.81 acres were classified as suitable for Recreation and Public Purposes. The classification is hereby terminated and the segregation for the following described lands is hereby terminated:

T. 2 N., R. 3 E., B.M.

Section 5: Lot 4.

The area described above aggregates 43.81 acres in Ada County.

At 9 a.m. on November 9, 1999, the Recreation and Public Purposes Act classification identified above will be terminated. The lands will remain closed to location and entry under the public land laws and the general land laws, as the lands are currently segregated for exchange.

Dated: November 3, 1999.

Jimmie Buxton,

Branch Chief, Lands and Minerals. [FR Doc. 99–29333 Filed 11–8–99; 8:45 am] BILLING CODE 4310–GG–P

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[NM-010-00-1430-HN/NM-101521 and NM-101522/G010-G0-0251]

Notice of Exchange Proposal/Notice of Intent To Prepare Two Environmental Impact Statements for the Proposed Santo Domingo and San Felipe Land Exchanges

AGENCY: Bureau of Land Management, Interior.

ACTION: Notice of exchange proposal; notice of intent.

SUMMARY: Notice is hereby given that the Bureau of Land Management (BLM)