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: August 11, 2004.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 04–18839 Filed 8–17–04; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Title: 45 CFR Part 95, Section F.

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's submitted APD provides the Department of Health and Human Services (HHS) 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 benefit analysis;

(4) A proposed activity schedule; and,

(5) A proposed budget.

HHS' 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: States.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses 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 Agreements	14	1	1	14
Biennial Reports	50	1	1.5	75

Estimated Total Annual Burden Hours: 5,751.5

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, Office of Information Services, 370 L'Enfant Promenade, SW., Washington, DC 20447, Attn: ACF Reports Clearance Officer. e-mail address: *grjohnson@acf.hhs.gov*.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following: Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW., Washington, DC 20503, Attn: Desk Officer for ACF, e-mail address: *katherine_t._astrich@omb.eop.gov*.

Dated: August 8, 2004.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 04–18840 Filed 8–17–04; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Joint Meeting of the Ear, Nose, and Throat Devices Panel and the Dental Products 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: Joint meeting of the Ear, Nose, and Throat Devices Panel and the Dental Products 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 October 6, 2004, from 8 a.m. to 5:30 p.m.

Location: Hilton Washington DC North/Gaithersburg, Salons A, B, and C, 620 Perry Pkwy., Gaithersburg, MD.

Contact Person: Sara M. Thornton, Center for Devices and Radiological Health (HFZ–460), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–2053, ext. 127, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512522. Please call the Information Line for up-to-date information on this meeting.

Agenda: The committee will discuss general issues surrounding the prescription use versus over the counter (OTC) use of devices intended to treat snoring or mild to severe obstructive sleep apnea (OSA). The discussion will include the role of the medical/dental provider in the diagnosis, treatment, and followup of snoring and OSA; the ability of the patient to self diagnose and treat OSA; the types of clinical data that would be needed to support an OTC intended use; and the components of adequate device labeling. The discussion will not include continuous positive airway pressure (CPAP) devices and surgical treatments for OSA. Background information, including the attendee list, agenda, and questions for the committee, will be available to the public 1 business day before the meeting, on the Internet at *http://www.fda.gov/cdrh/panelmtg.html*.

Procedure: On October 6, 2004, from 8:30 a.m. to 5:30 p.m., the meeting will be 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 September 17, 2004. Oral presentations from the public will be scheduled between approximately 9:15 a.m. and 9:45 a.m. Near the end of the committee discussion, a second 30-minute open public session will be conducted for interested persons to comment further on the discussion topic. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before September 17, 2004, 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 6, 2004, from 8 a.m. to 8:30 a.m., the meeting will be closed to permit FDA staff to present to the committee trade secret and/or confidential commercial information relevant to pending and future device submissions for ear, nose, and throat devices. This portion of the meeting will be closed to permit discussion of this information (5 U.S.C. 552b(c)(4)).

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact AnnMarie Williams, Conference Management Staff, at 301–594–1283, ext. 113, at least 7 days in advance of the meeting.

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

Dated: August 11, 2004.

William K. Hubbard,

Associate Commissioner Policy and Planning.

[FR Doc. 04–18849 Filed 8–17–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Bureau of Customs and Border Protection

Proposed Collection; Comment Request: Alien Crewman Landing Permit

ACTION: Notice and request for comments.

SUMMARY: As part of its continuing effort to reduce paperwork and respondent burden, Bureau of Customs and Border Protection (CBP) invites the general public and other Federal agencies to comment on an information collection requirement concerning the Alien Crewman Landing Permit. This request for comment is being made pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)).

DATES: Written comments should be received on or before October 18, 2004, to be assured of consideration.

ADDRESSES: Direct all written comments to Bureau of Customs and Border Protection, Information Services Group, Room 3.2.C, 1300 Pennsylvania Avenue, NW., Washington, DC 20229.

FOR FURTHER INFORMATION CONTACT: Requests for additional information should be directed to Bureau of Customs and Border Protection, Attn.: Tracey Denning, Room 3.2.C, 1300 Pennsylvania Avenue NW., Washington, DC 20229, Tel. (202) 344–1429.

SUPPLEMENTARY INFORMATION: CBP invites the general public and other Federal agencies to comment on proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (Pub. L. 104–13; 44 U.S.C. 3505(c)(2)). The comments should address: (1) Whether the 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 estimates of the burden of the collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden including the use of automated collection techniques or the use of other forms of information technology; and (e) estimates of capital or start-up costs and costs of operations, maintenance, and purchase of services to provide information. The comments that are submitted will be summarized and included in the CBP request for Office of Management and Budget (OMB) approval. All comments will become a matter of public record. In this document CBP is soliciting comments concerning the following information collection:

Title: Alien Crewman Landing Permit.

OMB Number: 1651–0114.

Form Number: Form CBP–95A and 95B.

Abstract: This collection of information is used by CBP to document conditions and limitations imposed upon an alien crewman applying for benefits under Section 251 of the Immigration and Nationality Act.

Current Actions: There are no changes to the information collection. This submission is being submitted to extend the expiration date.

Type of Review: Extension.

Affected Public: Individuals.

Estimated Number of Respondents: 433,000.

Estimated Time Per Respondent: 5 minutes.

Estimated Total Annual Burden Hours: 35,939.

Estimated Total Annualized Cost on the Public: $359,390.

Dated: August 11, 2004.

Tracey Denning,

Agency Clearance Officer, Information Services Group.

[FR Doc. 04–18886 Filed 8–17–04; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR–4903–N–64]

Notice of Submission of Proposed Information Collection to OMB; Fair Housing Literacy Survey

AGENCY: Office of the Chief Information Officer.

ACTION: Notice.

SUMMARY: The proposed information collection requirement described below has been submitted to the Office of Management and Budget (OMB) for review, as required by the Paperwork Reduction Act. The Department is soliciting public comments on the subject proposal.

Survey to determine the extent of public awareness of the nation's fair housing laws.

DATES: *Comments Due Date:* September 17, 2004.

ADDRESSES: Interested persons are invited to submit comments regarding this proposal. Comments should refer to the proposal by name and/or OMB approval Number (2528–0212) and should be sent to: HUD Desk Officer, Office of Management and Budget, New Executive Office Building, Washington, DC 20503; fax: 202–395–6974.

FOR FURTHER INFORMATION CONTACT: Wayne Eddins, Reports Management Officer, AYO, Department of Housing and Urban Development, 451 Seventh Street, SW., Washington, DC 20410; e-mail *Wayne_Eddins@HUD.gov*; telephone (202) 708–2374. This is not a toll-free number. Copies of available documents submitted to OMB may be