proposed collection of information; (c) ways to enhance 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. Send comments to Sandi Gambescia, CDC Assistant Reports Clearance Officer, 1600 Clifton Road, MS–E11, Atlanta, GA 30333. Written comments should be received within 14 days of this notice.

Proposed Project

Assessment of Occupational Electric and Magnetic Field (EMF) Exposures—Validation of Interview Procedures used in a Brain Tumor Study against Measurements of Biologically-based Exposure Metrics—New—National Institute for Occupational Safety and Health (NIOSH), Centers for Disease Control and Prevention (CDC).

This study to assess occupational exposures to electric and magnetic fields (EMF) has the following objectives: (1) Validate an interview-based EMF exposure assessment algorithm against measurements of the time-weighted average (TWA) magnetic field magnitude used in previous epidemiologic studies, (2) calibrate the parameters in the algorithm in order to improve the exposure estimates, and (3) determine the correlation between the EMF exposures from the algorithm and biologically-based metrics measured by new instrumentation. These biologically-based metrics consist of either characteristics of the magnetic field that have produced biological effects in laboratory studies or currents in the body resulting from contact with charged surfaces. For the higher correlations with the TWA magnetic field magnitude, these data will be used to determine whether the exposure algorithm can be modified to accurately assess exposures to the biologically-based metrics.

This is a one-time study of workers of an electric utility in Canada and a federal research laboratory in the U.S. There will be no cost to respondents.

Respondents	Number of respondents	Number of responses/respondent	Average burden/response (in hrs)	Total burden (in hrs)
Workers	108	1	15/60	27
Total	108			27

Dated: August 12, 2004.

Alvin Hall,

Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 04–18922 Filed 8–17–04; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title:

Improper Payments Best Practices Survey for the TANF Program

Improper Payments Best Practices Survey for the CCDF Program

OMB No.: New collection.

Description: These surveys for the Temporary Assistance for Needy Families (TANF) and the Child Care and Development Fund (CCDF) programs will request that states voluntarily provide information, including how they fefine improper payments in their state, the process used to identify such payments and what actions are taken in the state to reduce or eliminate improper payments. The Administration for Children and Families (ACF) within the U.S. Department of Health and Human Services (HHS) intends to establish a repository for the state submissions which will be available to all states for viewing on an HHS/ACF website. This website will provide information that will help states improve their program integrity system(s) so that improper payments in the programs can be reduced.

Respondents: The 50 States of the United States, the District of Columbia, and the Territories of Guam, Puerto Rico, and the Virgin Islands.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Improper Payments Best Practices Survey for the TANF Program	54	1	24	1,296
Improper Payments Best Practices Survey for the CCDF Program	54	1	24	1,296
Estimated Total Annual Burden Hours				2,592

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 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.* 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 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