Dated: January 8, 1997.

Carolyn J. Russell,

Director, Management Analysis and Services Office, Centers of Disease Control and Prevention (CDC).

[FR Doc. 97–965 Filed 1–14–97; 8:45 am]

BILLING CODE 4160–19–P

NIOSH Meeting; The National Institute for Occupational Safety and Health (NIOSH) of the Centers for Disease Control and Prevention (CDC) Announces the Following Meeting

Name: ''Postural Stability and Motor Response Times During Scaffold End Frame Handling'' study protocol peer review.

Time and Date: 1–4 P.M., February 13, 1997.

Location: Suncrest Facility, Large Conference Room, NIOSH, CDC, 3040 University Avenue, Morgantown, West Virginia 26505.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: Participants will provide NIOSH with their individual advice and comments regarding technical and scientific aspects of the NIOSH protocol ''Postural Stability and Motor Response Times During Scaffold End Frame Handling.'' Peer review panelists will review the study protocol and provide individual advice on the conduct of the study. Viewpoints and suggestions from industry, labor, academia, other governmental agencies, and the public are invited.

Agenda items are subject to change, as priorities dictate.

For Further Information Contact: Brian E. Moyer, M/S 119, 1095 Willowdale Road, Morgantown, West Virginia 26505, telephone (304) 285–5969.

Dated: January 8, 1997.

Carolyn J. Russell,

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

[FR Doc. 97–964 Filed 1–14–97; 8:45 am]

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Administration for Children and Families

Proposed Information Collection Activity; Comment Request

Proposed Projects:

Title: Child Care Quarterly Unit Report

OMB No.: New collection

Description: This legislatively-mandated report collects program and participants data on children and families receiving direct CCDF services. Disaggregate data will be collected and will be used to determine the participants and program characteristics as well as cost and level of child care services. The data will be used to provide a report to Congress. Form ACF 801 represents the data elements to be collected and reported to ACF.

Respondents (States and Territories) will be asked to sample the population of families receiving benefits on a monthly basis and submit the three most current monthly samples to ACF quarterly. Each monthly sample is drawn independent of the other samples and retained for submission within a quarterly report. ACF is not issuing specifications on how respondents compile overall database(s) from which samples are drawn. ACF will provide to the respondents a sampling plan which will specify minimum sample size. It is expected to be a monthly sample of approximately 150 cases for large States with smaller samples based on population size adjustments for smaller respondents.

Respondents: States, D.C., Guam, Virgin Islands and Puerto Rico

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF–801	54	4	20	4,320

Estimated Total Annual Burden Hours: 4,320.

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, Division of Information Resource Management Services, 370 L'Enfant Promenade, SW., Washington, DC 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 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: January 9, 1997.

Douglas J. Godesky,

Reports Clearance Officer.

[FR Doc. 97–940 Filed 1–14–97; 8:45 am]

BILLING CODE 4184–01–M

Food and Drug Administration

[Docket No. 96N–0488]

Use of Clorsulon Drench in Goats; Availability of Data

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the availability of target animal safety and effectiveness data, human food safety data, and environmental data to be used in support of a new animal drug application (NADA) or supplemental NADA for use of a suspension containing 8.5 percent clorsulon as a drench in goats for the treatment of adult liver fluke infestation. The data, contained in Public Master File (PMF) 5440, were compiled under National Research Support Project No. 7 (NRSP–7), a national agricultural program for obtaining clearances for use of new drugs in minor animal species or in any animal species for the control of diseases that occur infrequently or in limited geographical areas.

ADDRESSES: Submit NADA's or supplemental NADA's to the Document Control Unit (HFV–199), Center for Veterinary Medicine, Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855.