to comments and suggestions submitted within 60 days of this publication.

Dated: January 22, 1996.

Roberta Katson,

Director, Division of Information, Resource Management Services.

[FR Doc. 96–1341 Filed 1–25–96; 8:45 am]

BILLING CODE 4184–01–M

Proposed Information Collection Activity; Comment Request

Proposed Project(s)

Title: Welfare Reform Demonstration: Special Application Form.

OMB No.: 0970–0134.

Description: The form will be used by State welfare agencies to apply for federal waivers for certain welfare reform demonstrations under section 115(a) of the Social Security Act. Requests for waivers of federal law for demonstration projects falling within any of 5 broad policy areas outlined by the President in his 7/31/95 speech to the National Governors' association and submitted with the information requested on this form will be approved by the federal government within 30 days of receipt of the request.

Respondents: State governments

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Special Form	54	1	0.75	40.5.

Estimated Total Annual Burden Hours: 40.5

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 below. 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, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by title. Electronic comments must be submitted as an ASCII file without special characters or encryption.

Comments are invited 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: January 22, 1996.

Roberta Katson,

Director, Division of Information, Resource Management Services.

[FR Doc. 96–1340 Filed 1–25–96; 8:45 am]

BILLING CODE 4184–01–M

Proposed Information Collection Activity; Comment Request

Proposed Project(s)

Title: Statistical Report on Recipients Under Public Assistance.

OMB No.: 0970–0003.

Description: This report is basic to the proper administration and monitoring of the AFDC program. It provides quarterly information on applications, disposition of applications and reasons for discontinuances. The information is aggregated, analyzed and published in ''Quarterly Public Assistance Statistics'' by the Office of Family Assistance.

Respondents: State governments.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
AFC–3800	54	4	4	864

Estimated Total Annual Burden Hours: 864.

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 below. 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, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by title. Electronic comments must be submitted as an ASCII file without special characters or encryption.

Comments are invited 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) 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. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: January 22, 1996.

Roberta Katson,

Director, Division of Information Resource Management Services.

[FR Doc. 96–1339 Filed 1–25–96; 8:45 am]

BILLING CODE 4184–01–M

Centers for Disease Control and Prevention

Injury Research Grant Review Committee: Conference Call Meeting

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces the following conference call committee meeting.

Name: Injury Research Grant Review Committee (IRGRC).

Time and Date: 2 p.m.–5 p.m., February 12, 1996.

Place: National Center for Injury Prevention and Control (NCIPC), CDC, Koger Center, Vanderbilt Building, 1st Floor, Conference Room 1006, 2939 Flowers Road, South, Atlanta, Georgia 30341. (Exit Chamblee-Tucker Road off I–85.)

Status: Open: 2 p.m.–2:15 p.m., February 12, 1996. Closed: 2:15 p.m.–5 p.m., February 12, 1996.

Purpose: This committee is charged with advising the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the scientific merit and technical feasibility of grant applications relating to the support of injury control research projects and injury prevention research centers.

Matters To Be Discussed: Agenda items for the meeting will include announcements, discussion of review procedures, future meeting dates, and review of grant applications.

Beginning at 2:15 p.m., through 5 p.m., February 12, the Committee will meet to conduct a review of grant applications. This portion of the meeting will be closed to the public in accordance with provisions set forth in section 552b(c) (4) and (6), title 5 U.S.C., and the Determination of the Associate Director for Management and Operations, CDC, pursuant to Pub. L. 92–463.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Richard W. Sattin, M.D., Executive Secretary, IRGRC, National Center for Injury Prevention and Control, CDC, 4770 Buford Highway, NE, M/S K58, Atlanta, Georgia 30341–3724, telephone (770) 488–4580.

Dated: January 23, 1996.

Julia M. Fuller,

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

[FR Doc. 96–1495 Filed 1–25–96; 8:45 am]

BILLING CODE 4163–18–M

Savannah River Site Environmental Dose Reconstruction Project: Public Workshops

The National Center for Environmental Health (NCEH) of the Centers for Disease Control and Prevention (CDC) announces the following meeting.

Name: Savannah River Site Environmental Dose Reconstruction Project: Public Workshops.

Date: Wednesday, February 14, 1996.

Time: 7 p.m.–9 p.m.

Place: Holiday Inn Express, 1350 Whiskey Road, Aiken, South Carolina 29803.

Date: Thursday, February 15, 1996.

Time: 7 p.m.–9 p.m.

Place: Hilton—The DeSoto, 15 East Liberty Street, Savannah, Georgia 31401.

Status: Open to the public for observation and comment, limited only by the space available. The meeting room will accommodate approximately 50 people.

Background: Under a Memorandum of Understanding (MOU) signed in December 1990 with the Department of Energy (DOE), the Department of Health and Human Services (HHS) has been given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production and use. HHS delegated program responsibility to CDC.

In addition, an MOU was signed in October 1990 and renewed in November 1992 between ATSDR and DOE. The MOU delineates the responsibilities and procedures for ATSDR's public health activities at DOE site required under sections 104, 105, 107, and 120 of the Comprehensive Environmental Response, Compensation, and Liability Act (CERCLA, or ''Superfund''). These activities include health consultations and public health assessments at DOE sites listed on, or proposed for, the Superfund National Priorities List and at sites that are the subject of petitions from the public; and other health-related activities such as epidemiologic studies, health surveillance, exposure and disease registries, health education, substance-specific applied research, emergency response, and preparation of toxicological profiles.

Purpose: The purpose of these meetings is to support research which evaluates past releases of radioactive materials and chemicals from the SRS to the surrounding environment. The Project has already undergone a first phase. Phase I involved searching the site to identify and retrieve important documents to be used for dose reconstruction. Phase II will use this information to calculate chemical and radiological source terms and identify possible intake pathways (eating, drinking, and inhalation) for people who have lived in the SRS area.

Agenda items are identical for each meeting, and subject to change as priorities dictate.

Contact Person for More Information: Paul G. Renard, Project Manager, Radiation Studies Branch, Division of Environmental Hazards and Health Effects, NCEH, CDC, 4770 Buford Highway, NE, (F–35), Atlanta, Georgia 30341–3724, telephone 770/488–7040, FAX 770/488–7044.

Dated: January 22, 1996.

Julia M. Fuller,

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

[FR Doc. 96–1365 Filed 1–25–96; 8:45 am]

BILLING CODE 4163–18–M

Food and Drug Administration

[Docket No. 96N–0012]

Animal Drug Export; ANIPRYL® Tablets

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Deprenyl Animal Health, Inc., has filed an application requesting approval for export of the animal drug ANIPRYL® (l-selegiline hydrochloride, l-deprenyl hydrochloride) tablets to Canada.

ADDRESSES: Relevant information on this application may be directed to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, and to the contact person identified below. Any future inquiries concerning the export of nonfood animal drugs under the Drug Export Amendments of 1986 should also be directed to the contact person.

FOR FURTHER INFORMATION CONTACT: Gregory S. Gates, Center for Veterinary Medicine (HFV–114), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1617.

SUPPLEMENTARY INFORMATION: The drug export provisions in section 802 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 382) provide that FDA may approve applications for the export of drugs that are not currently approved in the United States. Section 802(b)(3)(B) of the act sets forth the requirements that must be met in an application for approval. Section 802(b)(3)(C) of the act requires that the agency review the application within 30 days of its filing to determine whether the requirements of section 802(b)(3)(B) have been satisfied. Section 802(b)(3)(A) of the act requires that the agency publish a notice in the Federal Register within 10 days of the filing of an application for export to facilitate public participation in its review of the