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.

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

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 contact person. 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 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.

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 the sections 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

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]
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This notice is to inform interested parties of the opportunity to apply for funds for a cooperative agreement from HCFA’s Office of Research and Demonstrations for the “End-Stage Renal Disease (ESRD) Managed Care Demonstration.”

FOR FURTHER INFORMATION, CONTACT: Bonnie Edington (410) 786–6617.

SUPPLEMENTARY INFORMATION:

I. Background

Section 2355 of the Deficit Reduction Act of 1984 (Pub. L. 98–369) required the Secretary to grant demonstration waivers for social health maintenance organization (SHMO) projects that provide for the integration of health and social services at a fixed annual prepaid capitation rate.

Section 4207(b)(4)(B) of the Omnibus Budget Reconciliation Act of 1990 (Pub. L. 101–508) amended section 2355 of Pub. L. 98–369 to include a requirement that the Secretary conduct up to four additional SHMO projects to demonstrate the effectiveness and feasibility of innovative approaches to refining current targeting and financing methodologies and benefit design for SHMOs.

Section 13567(b) of the Omnibus Budget Reconciliation Act (OBRA) of 1993 (Pub. L. 103–66) further amended section 2355 of Pub. L. 98–369, requiring the Secretary to include the integration of acute and chronic care management for patients with end-stage renal disease through expanded community care case management services in at least one of the four additional SHMO projects.

II. Provisions of this Notice

This notice is to inform interested parties of the opportunity to apply for funds for a cooperative agreement to operate an “End-Stage Renal Disease (ESRD) Managed Care Demonstration,” involving the treatment of Medicare-eligible ESRD patients in a managed care setting as required by OBRA 1993. Interested parties are required to submit an official application for consideration for grant funding and commencement of site development activities. Subject to funds availability, a one-time award of approximately $175,000 is expected to be given to each selected awardee. HCFA expects to award one or more demonstrations through the application process. Any organizational entity may apply. However, the applicant must be capable of assuring that the service delivery system under the demonstration will integrate acute and chronic care services, through expanded community care case management services, for ESRD patients. In addition, the applicant must meet all applicable State requirements for bearing financial risk. Awardees are expected to have a 9 to 12-month development period subsequent to award and prior to service delivery. In the three-year service delivery phase of the demonstration, awardees will be paid on a capitation basis in which the capitation amount will be adjusted to reflect treatment status (that is, maintenance dialysis, transplant, or functioning graft).

Potential applicants who wish to request the full solicitation and application packet should call Ms. Edington at the above telephone number, send an e-mail message to BEDINGTON@HCFA.GOV, or write to the following address: Bonnie Edington, Health Care Financing Administration, Office of Research and Demonstrations, Mail Stop C3–24–07, 7500 Security Boulevard, Baltimore, MD 21244–1850. These packets will be mailed to all requestors within approximately 10 days following the date of this notice. Completed applications, including full proposals, will be due approximately 70 days following the date of this notice. The exact due date for applications will be specified in the application packet. Awards are expected to be made in 1996.


In accordance with the provisions of Executive Order 12866, this notice was reviewed by the Office of Management and Budget. (Catalog of Federal Domestic Assistance Program No. 93.779 Health Financing Research, Demonstrations and Experiments)


Bruce C. Vladeck, Administrator, Health Care Financing Administration.

[FR Doc. 96–1261 Filed 1–25–96; 8:45 am]
BILLING CODE 4120–01–P

BPO–134–NC

Medicare Program; Revised Criteria and Standards for Evaluating Durable Medical Equipment, Orthotics, and Supplies Regional Carriers’ Performance Beginning February 1, 1996

AGENCY: Health Care Financing Administration (HCFA), HHS.

[FR Doc. 96–1321 Filed 1–25–96; 8:45 am]
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