epidemiologic study of almost 1,000,000 nuclear workers in 14 nations.

5. Bringing together investigators from different countries is a well-developed practice at IARC. Most of their research on occupational, environmental or lifestyle hazards has been, or is presently being, conducted on an international basis. This provides the opportunity for a wide selection of populations suitable for epidemiologic investigation and facilitates the accumulation of large study populations, thus making the identification and quantification of cancer risk easier.

Executive Order 12372—Review

The application is not subject to review under Executive Order 12372.

Public Health System Reporting Requirements

This program is not subject to the Public Health System Reporting Requirements.

Catalog of Federal Domestic Assistance Number

The Catalog of Federal Domestic Assistance Number for this program is 93.283.

Other Requirements

Paperwork Reduction Act

Projects that involve the collection of information from 10 or more individuals and funded by the cooperative agreement will be subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act.

Human Subjects

If the proposed project involves research on human subjects, the applicant must comply with the Department of Health and Human Services Regulations, 45 CFR part 46, regarding the protection of human subjects. Assurance must be provided to demonstrate that the project will be subject to initial and continuing review by an appropriate institutional review committee. The applicant will be responsible for providing assurance in accordance with the appropriate guidelines and forms provided in the application kit.

Where To Obtain Additional Information

If you are interested in obtaining additional information regarding this project, please refer to Announcement 520 and contact Oppie M. Byrd, Grants Management Specialist, Grants Management Branch, Procurement and Grants Office, Centers for Disease Control and Prevention (CDC), 255 East Paces Ferry Road, NE., Room 300, Mailstop E–13, Atlanta, GA 30305, telephone (404) 842–6546.

A copy of Healthy People 2000 (Full Report, Stock No. 017–001–00474–0) or Healthy People 2000 (Summary Report, Stock No. 017–001–00473–1) referenced in the ''Summary'' may be obtained through the Superintendent of Documents, Government Printing Office, Washington, DC 20402–9325, telephone (202) 783–3238.

Dated: January 19, 1995.

Richard A. Lemen,

Acting Director, National Institute for Occupational Safety and Health, Centers for Disease Control and Prevention (CDC).

[FR Doc. 95–1808 Filed 1–24–95; 8:45 am]

BILLING CODE 4163–19–P

Advisory Committee on Immunization Practices; 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 committee meeting:

Name: Advisory Committee on Immunization Practices.

Times and Dates: 8:30 a.m.–6:30 p.m., February 9, 1995. 8:15 a.m.–4:45 p.m., February 10, 1995.

Place: CDC, Auditorium A, Building 2, 1600 Clifton Road, NE, Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available.

Purpose: The committee is charged with advising the Director, CDC, on the appropriate uses of immunizing agents.

Matters to be Discussed: The committee will discuss recommendations for prevention of hepatitis A: Hepatitis A vaccine and immune globulin; revised recommendations for hepatitis B vaccination; update on varicella vaccine; status on principles and guidelines for combination products; vaccines for children (VFC): influenza vaccine in VFC, hepatitis B for adolescents in VFC, hepatitis A, MMR2—expanded use in VFC; pneumococcal polysaccharide vaccine; adolescent vaccination; poliomyelitis prevention; influenza: 1995–96 influenza vaccine strain selection, 1995–96 influenza vaccine and antiviral recommendations, influenza-associated morbidity during pregnancy, assessment of BBS risk associated with 1993–94 and 1994–95 influenza vaccination, optimal needle length for intramuscular injection into the deltoid, national estimates of influenza vaccination rates; update on meningococcal recommendation; report of a meeting regarding conflicting immunization guidelines and harmonization of the Advisory Committee on Immunization Practices/American Academy of Pediatrics recommendations with the Food and Drug Administration labeling; update on simplification; progress towards 1996 disease reduction goals; recommendations for immunization linkage with the women's, infants, and children program; vaccine safety; an update on the Injury Compensation Program; an update on the National Vaccine Program; and a presentation on acellular pertussis. Other matters of relevance among the committee's objectives may be discussed.

Agenda items are subject to change as priorities dictate.

For Further Information Contact: Gloria A. Kovach, Committee Management Specialist, CDC (1–B72), 1600 Clifton Road, NE., Mailstop A20, Atlanta, Georgia 30333, telephone 404/639–3851.

Dated: January 18, 1995.

William H. Gimson,

Acting Associate Director for Policy Coordination, Centers for Disease Control and Prevention (CDC).

[FR Doc. 95–1805 Filed 1–24–95; 8:45 am]

BILLING CODE 4163–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 94F–0454]

Lyondell-Citgo Refining Co., Ltd.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Lyondell-Citgo Refining Co., Ltd., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of white mineral oil as a dust control agent for rough rice.

DATES: Written comments on the petitioner's environmental assessment by February 24, 1995.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Blondell Anderson, Center for Food Safety and Applied Nutrition (HFS–207), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3106.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 5A4440) has been filed by Lyondell-Citgo Refining Co., Ltd., P.O. Box 2451, Houston, TX 77252–2451. The petition proposes to amend the food additive regulations in § 172.878 *White mineral oil* (21 CFR 172.878) to provide for the safe use of white mineral oil as

a dust control agent for rough rice at an application rate of 800 parts per million.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (address above) for public review and comment. Interested persons may, on or before February 24, 1995, submit to the Dockets Management Branch (address above) written comments. Two copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental assessment without further announcement in the **Federal Register**. If, based on its review, the agency finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding of no significant impact and the evidence supporting that finding will be published with the final regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: January 13, 1995.

Alan M. Rulis,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–1766 Filed 1–24–95; 8:45 am]

BILLING CODE 4160–01–F

Food and Drug Administration

[Docket No. 94D–0300]

International Harmonization; Draft Policy on Standards; Availability; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a notice that appeared in the **Federal Register** of November 28, 1994 (59 FR 60870). The document provided a draft policy statement of the agency's development and use of standards with respect to international harmonization of regulatory requirements and guidelines. Specifically, the draft policy addressed the conditions under which FDA participates with standards bodies outside of FDA, domestic or international, in the development of standards applicable to products regulated by FDA. The document was published with some typographical and inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Linda Horton, International Policy Staff (HF-23), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–2831.

In FR Doc. 94–29116, appearing on page 60870, in the **Federal Register** of November 28, 1994, the following corrections are made:

1. On page 60872, in the second column, in the eighth line from the bottom, the word ''Biologic'' is corrected to read ''Biologics''; in the third column, in the first full paragraph, in the 21st line, the acronym ''(PhMA)'' is corrected to read ''(PhRMA)''; and in the same column, beginning in the second line from the bottom, the words ''standardizing the safety-related terminology used in adverse experience reporting'' are corrected to read ''standardizing medical definitions and adverse experience reporting''.

2. On page 60873, in the first column, in the first full paragraph, in the fourth line from the bottom of the paragraph, the word ''Device'' is corrected to read ''Devices''.

Dated: January 18, 1995.

William K. Hubbard,

Interim Deputy Commissioner for Policy.

[FR Doc. 95–1767 Filed 1–24–95; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF THE INTERIOR

Bureau of Land Management

[OR–030–03–1220–04; G5–051]

Notice of Prohibited Acts in Owyhee National Wild and Scenic River Area; Correction

AGENCY: Vale District, Bureau of Land Management, Interior.

ACTION: Correction.

SUMMARY: In notice document 94–30765 beginning on page 64693 in the issue of Thursday, December 15, 1994, make the following correction.

On page 64693 in the third column the **SUMMARY** Section previously stated paragraph 1. Fire a. Building or maintaining any open campfires except those contained in a firepan or similar metal container. This should be changed to read 1. Fire a. Building or maintaining any open campfires except those contained in a firepan or similar container.

James E. May,

District Manager.

[FR Doc. 95–1788 Filed 1–24–95; 8:45 am]

BILLING CODE 4310–33–M

National Park Service

Pea Ridge National Military Park Advisory Team; Notice of Meeting

Notice is hereby given in accordance with the Federal Advisory Committee Act that a meeting of the Pea Ridge National Military Park Advisory Team will be held at 6 p.m., on Thursday, February 16, 1995, in the park visitor center auditorium, 15930 Highway 62, Garfield, Arkansas.

The Pea Ridge National Military Park Advisory Team was established under authority of section 3 of Public Law 91–383 (16 U.S.C. 1a–2(c)) to provide a forum for dialogue between community representatives and the Pea Ridge National Military Park on management issues affecting the park and the community.

The matter to be discussed at this meeting includes:

—Boundary Study

The meeting will be open to the public. However, facilities and space for accommodating members of the public are limited, and persons will be accommodated on a first-come-first serve basis. Any member of the public may file a written statement concerning the matters to be discussed with the Superintendent, Pea Ridge National Military Park.

Persons wishing further information concerning this meeting, or who wish to submit written statements may contact Steve Adams, Superintendent, Pea Ridge National Military Park, P.O. Box 700, Pea Ridge, AR 72751–0700, Telephone 501/451–8122.

Minutes of the meeting will be available for public inspection four weeks after the meeting at the office of Pea Ridge National Military Park.

Dated: January 12, 1995.

John D. Linahan,

Acting Regional Director, Southwest Region.

[FR Doc. 95–1769 Filed 1–24–95; 8:45 am]

BILLING CODE 4310–70–M