

Monitor has the requisite capability and applicable business knowledge to supervise the proper transfer of divested assets and monitor the critical manufacturing and supply activities of Metso and Svedala. Thus, the transitional manufacturing agreement, in conjunction with the Interim Monitor, provides a guarantee to Sandvik that its production of jaw crushers will be seamless and uninterrupted after the divestiture.

In order to ensure that the Commission remains informed about the status of the crushing businesses and the grinding mill business pending divestiture, and about the efforts being made to accomplish the divestitures, the Consent Agreement requires Metso and Svedala to file reports with the Commission within thirty (30) days of the date they sign the Consent Agreement, and periodically thereafter, until the divestitures are accomplished.

The purpose of this analysis is to facilitate public comment on the Consent Agreement, and it is not intended to constitute an official interpretation of the Consent Agreement or to modify in any way its terms.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 01-23234 Filed 9-17-01; 8:45 am]

BILLING CODE 6750-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Citizens Advisory Committee on Public Health Service Activities and Research at Department of Energy (DOE) Sites: Idaho National Engineering and Environmental Laboratory Health Effects Subcommittee (INEELHES)

In accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463), the Agency for Toxic Substances and Disease Registry (ATSDR) and the Centers for Disease Control and Prevention (CDC) announce the following meeting.

Name: Citizens Advisory Committee on Public Health Service Activities and Research at Department of Energy (DOE) Sites: Idaho National Engineering and Environmental Laboratory Health Effects Subcommittee (INEELHES).

Times and Dates: 8:30 a.m.—4:45 p.m., October 16, 2001; 8:30 a.m.—3:45 p.m., October 17, 2001.

Place: WestCoast Pocatello Hotel, 1555 Pocatello Creek Road, Pocatello,

Idaho 83201, telephone, (208) 233-2200, fax (208) 234-4524.

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

Background: Under a Memorandum of Understanding (MOU) signed in December 1990 with DOE, and replaced by MOUs signed in 1996 and 2000, the Department of Health and Human Services (HHS) was given the responsibility and resources for conducting analytic epidemiologic investigations of residents of communities in the vicinity of DOE facilities, workers at DOE facilities, and other persons potentially exposed to radiation or to potential hazards from non-nuclear energy production use. HHS delegated program responsibility to CDC.

In addition, a memo was signed in October 1990 and renewed in November 1992, 1996, and in 2000, between ATSDR and DOE. The MOU delineates the responsibilities and procedures for ATSDR's public health activities at DOE sites 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: This subcommittee is charged with providing advice and recommendations to the Director, CDC, and the Administrator ATSDR, regarding community concerns pertaining to CDC's and ATSDR's public health activities and research at this DOE site. The purpose of this meeting is to provide a forum for community interaction and serve as a vehicle for community concerns to be expressed as advice and recommendations to CDC and ATSDR.

Matters to be Discussed: Agenda items include an update regarding progress of current studies; a review of the COSMOS evaluation report; strategies to develop INEELHES' internal evaluation; an overview of Idaho National Engineering and Environmental Laboratory; and a presentation on Health Consult by ATSDR.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Paul G. Renard, Executive Secretary, Radiation Studies Branch, Division of Environmental Hazards and Health Effects, NCEH, CDC, 1600 Clifton Road, NE (E-39), Atlanta, GA 30333, telephone (404) 498-1800, fax (404) 498-1811.

The Director, Management Analysis and Services Office, has been delegated the authority to sign **Federal Register** notices pertaining to announcements of meetings and other committee management activities for both CDC and ATSDR.

Dated: September 7, 2001.

Carolyn J. Russell,

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

[FR Doc. 01-23246 Filed 9-17-01; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare & Medicaid Services

[CMS-4026-PN]

RIN 0938-ZA21

Medicare Program; Medicare+Choice Organizations—Application by the Joint Commission on Accreditation of Healthcare Organizations (JCAHO) for Approval of Deeming Authority for Medicare+Choice Organizations That Are Licensed as Health Maintenance Organizations (HMOs) or Preferred Provider Organizations (PPOs)

AGENCY: Centers for Medicare & Medicaid Services (CMS), HHS.

ACTION: Proposed notice.

SUMMARY: This proposed notice announces the receipt of an application from the Joint Commission on Accreditation of Healthcare Organizations (JCAHO) for recognition as a national accreditation program for health maintenance organizations (HMOs) and preferred provider organizations (PPOs) that wish to participate in the Medicare+Choice program. Regulations set forth at 42 CFR 422.157(b)(1) specify that a **Federal Register** notice will announce our receipt of the accreditation organization's application for approval, describe the criteria we will use in evaluating the application, and provide at least a 30-day public comment period.

DATES: We will consider comments if we receive them at the appropriate