the relevant product markets. Pursuant to the Consent Agreement, Sun is required to divest all of its rights and assets related to the Products to a Commission-approved acquirer no later than the earlier of ten (10) days after the acquisition occurs or ten (10) days after the Commission's Order becomes final. Specifically, the proposed Consent Agreement requires that Sun divest its assets in the Products to Torrent Pharmaceutical Limited (''Torrent'').

The acquirer of the divested assets must receive the prior approval of the Commission. The Commission's goal in evaluating a possible purchaser of divested assets is to maintain the competitive environment that existed prior to the acquisition. A proposed acquirer of divested assets must not itself present competitive problems.

Torrent, a growing generic manufacturer, headquartered in India, is particularly well-positioned to manufacture and market its acquired products and compete effectively in those markets. Currently, Torrent sells generic pharmaceuticals in the United States but none of the relevant products, and therefore its acquisition of the relevant products would not raise independent competitive concerns. Torrent has numerous Abbreviated New Drug Applications (ANDAs'') pending approval at the FDA, and has the resources, capabilities, reputation, and experience in marketing generic products, as well as a central focus on rapidly growing its U.S. generic drugs business, necessary to expeditiously replicate the competition that would be lost with the proposed acquisition.

If the Commission determines that Torrent is not an acceptable acquirer of the assets to be divested, or that the manner of the divestitures to Torrent is not acceptable, Sun must unwind the sale and divest the assets within six (6) months of the date the Order becomes final to another Commission-approved acquirer. If the parties fail to divest within six (6) months, the Commission may appoint a trustee to divest the Products.

The proposed remedy contains several provisions to ensure that the divestitures are successful. The Order requires Sun to provide transitional services to enable the Commission-approved acquirer to obtain all of the necessary approvals from the FDA. These transitional services include technology transfer assistance to manufacture the Products in substantially the same manner and quality employed or achieved by Sun.

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

By direction of the Commission.

Donald S. Clark

Secretary

[FR Doc. E8–19213 Filed 8–19–08; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee, National Institute for Occupational Safety and Health (NIOSH)

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 meeting for the aforementioned committee:

Time and Date: 8 a.m.—5:30 p.m., September 25, 2008.

Place: Hilton Garden Inn, 7830 S. Las Vegas Boulevard, Las Vegas, Nevada 89123, telephone (702) 453–7830, fax (702) 453–7850.

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

Purpose: This committee is charged with providing advice to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NIOSH, on priorities in mine safety and health research, including grants and contracts for such research, 30 U.S.C. 812(b)(2), Section 102(b)(2).

Matters To Be Discussed: The meeting will focus on diesel monitoring and control research, communications and tracking developments, training research, chemical hazards in mining, and dust monitoring and control research. The agenda will also include an update from the Associate Director for Mining, NIOSH.

Agenda items are subject to change as priorities dictate.

CONTACT PERSON FOR MORE INFORMATION: Jeffery L. Kohler, PhD, Executive Secretary, Mine Safety and Health Research Advisory Committee, NIOSH, CDC, 626 Cochrans Mill Road, Pittsburgh, Pennsylvania 15236, *telephone:* (412) 386–5301, Fax (412) 386–5300.

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 the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Dated: August 13, 2008.

Elaine L. Baker,

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

[FR Doc. E8–19240 Filed 8–19–08; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Comment Request

Proposed Project:

Title: Early Head Start Family and Child Experiences Survey (Baby FACES).

OMB No. New Collection

Description: The Administration for Children and Families (ACF), U.S. Department of Health and Human Services, is conducting a descriptive study of Early Head Start Programs (Early Head Start Family and Child Experiences Survey, or Baby FACES). Baby FACES is a longitudinal study of a nationally representative sample of programs and children in two cohorts (perinatal and age 1) that will collect information about programs, services, families, and children. Data for Baby FACES will be annually collected through interviews with parents, teachers, home visitors, and program directors/managers, as well as direct child assessments, videotaped parent child interactions, and observations of the home environment when children are two and three years old. Data collection will also include quality observations of child care center classrooms and home visits conducted by program staff.

Data will be collected on a sample of approximately 2,000 children and families selected at random from 90 Early Head Start programs. Over the life of the project, Baby FACES will involve four waves of data collection, ending when the second cohort of children (perinatal cohort) reaches 36 months of age. This information collection request covers the first three years of data collection. All waves of data collection will acquire program level information through an hour-long program director interview. Additionally, staff from all programs will complete a simple service