registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methamphetamine (1105)	II
Phenylacetone (8501)	II

The firm plans to import the listed controlled substances to manufacture pharmaceutical products.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import a basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.

Dated: June 18, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 96–16195 Filed 6–25–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on May 9, 1996, Arenol Chemical Corporation, 189 Meister Avenue, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396).	I
3,4-Methylenedioxyamphetamine (7400).	I
Difenoxin (9168)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II

The firms plans to manufacture difenoxin, amphetamine, methamphetamine and methylphenidate to produce pharmaceutical products for distribution to its customers; and 2,5-dimethoxyamphetamine and 3,4-methylenedioxyamphetamine as intermediates for the development of other pharmaceutical products.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the above application.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 26, 1996.

Dated: June 18, 1996.

Gene R. Haislip,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 96–16196 Filed 6–25–96; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Labor Advisory Committee for Trade Negotiations and Trade Policy

Pursuant to the provisions of the Federal Advisory Committee Act (P.L. 92–463 as amended), notice is hereby given of a meeting of the Steering Subcommittee of the Labor Advisory Committee for Trade Negotiations and Trade Policy.

Date, time and place: July 10, 1996, 10:00 am–12:00 noon, U.S. Department of Labor, Room S–1011, 200 Constitution Ave. NW., Washington, DC 20210.

Purpose: The meeting will include a review and discussion of current issues which influence U.S. trade policy. Potential U.S. negotiating objectives and bargaining positions in current and anticipated trade negotiations will be discussed. Pursuant to section 9(B) of the Government in the Sunshine Act, 5 U.S.C. 552b(c)(9)(B) it has been determined that the meeting will be concerned with matters the disclosure of which would seriously compromise the Government's negotiating objectives or bargaining positions. Accordingly, the meeting will be closed to the public.

For further information contact: Fernand Lavallee, Director, Trade Advisory Group or Jorge Perez-Lopez, Director, Office of International Economics Affairs, Phone: (202) 219–4752.

Signed at Washington, DC this 18th day of June, 1996.

Joaquin Otero,

Deputy Under Secretary, International Affairs.

[FR Doc. 96–16255 Filed 6–25–96; 8:45 am]

BILLING CODE 4510–28–M

Mine Safety and Health Administration

Petitions for Modification

The following parties have filed petitions to modify the application of mandatory safety standards under section 101(c) of the Federal Mine Safety and Health Act of 1977.

1. Kade Coal Company, Inc.

[Docket No. M–96–36–C]

Kade Coal Company, Inc., Route 1, Box 513, Grundy, Virginia 24614 has filed a petition to modify the application of 30 CFR 77.214(a) (refuse piles; general) to its Mine No. 2 (I.D. No. 44–06483) located in Buchanan County, Virginia. The petitioner proposes to cover several entries at each abandoned mine opening with coarse refuse material during construction of a refuse fill. Presently, the petitioner is depositing coarse refuse material on the existing Red Ash seam bench (Refuse Disposal No: 1211–VA5–0297). The petitioner requests this modification of the standard, to allow four drift openings to be filled with refuse where drift entries to old mine workings at Mine No. 3 (I.D. No. 44–06310) exist, during construction of refuse fill. The petitioner asserts that the proposed alternative method would provide at least the same measure of protection as would the mandatory standard.

2. Monterey Coal Company

[Docket No. M–96–37–C]

Monterey Coal Company, Rural Route 4, Box 235, Carlinville, Illinois 62626 has filed a petition to modify the application of 30 CFR 75.380(d)(4)