significantly affect the quality of the human environment in the sense of NEPA 102(2)(C). A FONSI is prepared in those instances where the MMS finds that approval will not result in significant effects on the quality of the human environment. The FONSI briefly presents the basis for that findings and includes a summary or copy of the EPA.

This notice constitutes the public Notice of Availability of environmental documents required under the NEPA regulations.

Dated: September 25, 1995.

J. Lisle Reed,

Regional Director, Pacific OCS Region.

[FR Doc. 95–25132 Filed 10–10–95; 8:45 am]

BILLING CODE 4310–MR–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated July 24, 1995, and published in the **Federal Register** on August 1, 1995, (60 FR 39185), Arenol Chemical Corporation, 189 Meister Avenue, Somerville, New Jersey 08876, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

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

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: September 29, 1995.

Gene R. Haislip,

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

[FR Doc. 95–25078 Filed 10–10–95; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

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 July 17, 1995, Ciba-Geigy Corporation, Pharmaceuticals Division Regulatory Compliance, 556 Morris Avenue, Summit, New Jersey 07901, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the Schedule II controlled substance Methylphenidate (1724).

The firm plans to manufacture the finished product for distribution to its customers.

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 to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Departments of Justice, Washington, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 11, 1995.

Dated: September 29, 1995.

Gene R. Haislip,

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

[FR Doc. 95–25076 Filed 10–10–95; 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 August 22, 1995, Lonza Riverside, 900 River Road, Conchohocken, Pennsylvania 19428, 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
4-Methoxyamphetamine (7411)	I
Amphetamine (1100)	II
Phenylacetone (8501)	II

The firm plans to manufacture the listed controlled substances as bulk product for distribution to its customers.

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 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 December 11, 1995.

Dated: September 29, 1995.

Gene R. Haislip,

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

[FR Doc. 95–25077 Filed 10–10–95; 8:45 am]

BILLING CODE 4410–09–M

Importer of Controlled Substances; Notice of Registration

By Notice dated May 30, 1995, and published in the Federal Register on June 8, 1995, (60 FR 30319), Radian Corporation, 8501 Mopac Blvd., P.O. Box 201088, Austin, Texas 78720, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Ibogaine (7260)	I
Etorphine (except HC1) (9056)	I
Heroin (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Oxycodone (9143)	II
Dextropropoxyphene, bulk (non-dosasge forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II

No comments or objections have been received. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1311.42, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: September 29, 1995.

Gene R. Haislip,

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

[FR Doc. 95–25079 Filed 10–10–95; 8:45 am]

BILLING CODE 4410–09–M