of the district be redrawn to exclude lands located within the Borough of Chambersburg, based upon a claimed loss of historic integrity of the area. Documentation relative to the historic integrity of this portion of the district was submitted to the National Register. Copies of this documentation are available from the National Register at the address below. In order to accommodate those who wish to provide new information concerning the boundary of the Eastern Greene Township Rural Historic District, the National Park Service is providing a 60 day comment period. A written statement on the determination of eligibility will be issued by the National Park Service within 30 days of the close of the comment period.

The determination of eligibility remains in effect pending review of responses submitted during the comment period. In order to revise the boundary the National Park Service must receive authoritative information, which evaluated in conjunction with documentation already on file, results in a finding that the determined eligible boundary does not accurately delineate the historic district in accordance with established National Register standards.

Comments should be addressed to the National Register of Historic Places, National Park Service, 1849 C St., N.W., Room NC400, Washington, D.C. 200240.

Carol D. Shull,

Keeper of the National Register of Historic Places, National Register, History and Education.

[FR Doc. 98–21015 Filed 8–5–98; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to Section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on May 1, 1998, applied Science Labs, Inc., A Division of Altech Associates, Inc., 2701 Carolean Industrial Drive, P.O. Box 440, State College, Pennsylvania 16801, made application to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Heroin (9200)	I
Morphine (9300)	II

The firm plans to import these controlled substances for the manufacture of reference standards.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances 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 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 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 1301.34(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 the basic classes of any controlled substances 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 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 17, 1998.

John H. King,

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

[FR Doc. 98–20972 Filed 8–5–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated April 17, 1998, and published in the **Federal Register** on April 30, 1998, (63 FR 23796), Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of phenylacetone (8501), a basic class of controlled substance listed in Schedule II.

The phenylacetone will be imported for conversion to amphetamine base, isomers and salts thereof for sale in bulk form to customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code, Section 823(a) and determined that the registration of Johnson Matthey, Inc. to import phenylacetone is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971, at this time. 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: July 17, 1998.

John H. King,

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

[FR Doc. 98–20973 Filed 8–5–98; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 8, 1998, and published in the **Federal Register** on February 12, 1998 (63 FR 7182), Nycomed, Inc., 33 Riverside Avenue, Renssalaer, New York 12144 made application to the Drug Enforcement Administration (DEA) by letter to be registered as a bulk manufacturer of methylphenidate (1724).

A registered bulk manufacturer of methylphenidate filed written comments and an objection in response to the notice of application. Review of the APA's definitions of license and