paragraphs 2 and 3 of this Order are limited to those that provide in-kind (non-cash) benefits and are open to individuals needing or desiring to participate without regard to income or resources. Programs, services or assistance delivered at the community level, even if they serve purposes of the type described in paragraph 3 above, are not within this specification if they condition (a) the provision of assistance, (b) the amount of assistance provided, or (c) the cost of the assistance provided on the individual recipient's income or resources.

Dated: August 23, 1996.

Janet Reno,

Attorney General.

[FR Doc. 96–22233 Filed 8–29–96; 8:45 am]

BILLING CODE 4410–01–M

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 15, 1995, Celgene Corporation, 7 Powder Horn Drive, Warren, NJ 07059, 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
Amphetamine (1100)	II

The firm plans to manufacture small quantities of 2,5-dimethoxyamphetamine using biocatalysis to develop, manufacture and sell high value added compounds to pharmaceutical and agrochemical industries and amphetamine for distribution of the bulk active substances 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, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, U.S. Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than October 29, 1996.

Dated: August 21, 1996.

Gene R. Haislip,

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

[FR Doc. 96–22218 Filed 8–29–96; 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 July 25, 1996, 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 methylphenidate (1724).

The firm plans to manufacture finished product for distribution to this 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, 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 October 29, 1996.

Dated: August 21, 1996.

Gene R. Haislip,

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

[FR Doc. 96–22219 Filed 8–29–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated May 22, 1996, and published in the Federal Register on May 30, 1996, (61 FR 27099), 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

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 Lonza Riverside to manufacture the listed controlled substances is consistent with the public interest at this time. Therefore, pursuant to 21 U.S.C. § 823 and 28 C.F.R. §§ 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed above is granted.

Dated: August 21, 1996.

Gene R. Haislip,

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

[FR Doc. 96–22148 Filed 8–29–96; 8:45 am]

BILLING CODE 4410–09–M

Manufacturer of Controlled Substances; Notice of Correction

As set forth in the Federal Register (FR Doc. 96–14057) Vol. 61, No. 109 at page 28598, dated June 5, 1996, Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer for certain controlled substances. The listing of controlled substances for which Penick Corporation applied should not have included the basic classes of controlled substances listed below:

Drug	Schedule
Cocoa Leaves (9040)	II
Opium, raw (9600)	II
Opium poppy (9650)	II
Poppy Straw Concentrate (9670)	II
Ethylmorphine (9190)	II

Therefore, Penick Corporation no longer wishes to be registered for the above listed controlled substances and they are hereby deleted from the list of controlled substances for which Penick Corporation made application to manufacture in bulk.