Elmo Avenue, Building 18 Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substance listed below:

Drug	Schedule
N-Ethylamphetamine (1475)	I
4-Methoxyamphetamine (7411)	I
2,5-Dimethoxyamphetamine (7396).	I
Difenoxin (9168)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Methylphenidate (1724)	II
Secobarbital (2315)	II
Meperidine (9230)	II
Codeine (9050)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II
Dextropropoxyphene (9273)	II

The firm plans to bulk manufacture the listed controlled substances to produce products for distribution to its customers.

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

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 (CCD), and must be filed no later than 60 days from publication.

Dated: April 3, 2003.

Laura M. Nagel,

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

[FR Doc. 03–9228 Filed 4–14–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on April 25, 2002, Cody Laboratories, Inc., 331 33rd Street, Cody, Wyoming 82414, made application by renewal, and on March 5, 2003, by letter to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of Schedule II of controlled substances listed below:

Drug	Schedule
Dihydromorphine (9145)	II
Methamphetamine (1105)	II
Amphetamine (1100)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phenylacetone (8501)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Fentanyl (9801)	II

The firm plans to produce bulk products 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 proposed registration.

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: Drug Operations Section, Domestic Drug Unit (ODOD) and must be filed no later than June 16, 2003.

Dated: April 3, 2003.

Laura M. Nagel,

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

[FR Doc. 03–9227 Filed 4–14–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated June 7, 2002, and published in the **Federal Register** on June 20, 2002 (67 FR 42060), Penick Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Coca Leaves (9040) and Poppy Straw (9650) basic classes of controlled substances listed in Schedule II.

The firm plans to import the controlled substances to manufacture bulk pharmaceutical controlled substances and non-controlled substance flavor extract.

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 Penick corporation to import these controlled substances 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. DEA has investigated Penick Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest.

This investigation included inspection and testing of the company's physical security system, verification of the company's compliance with state and local laws, and a review of the company's background and history. 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 1301.34 the above firm is granted registration as an importer of the basic classes of controlled substances listed.

Dated: April 3, 2003.

Laura M. Nagel,

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

[FR Doc. 03–9230 Filed 4–14–03; 8:45 am]

BILLING CODE 4410–09–M

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 registration 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 20, 2002, Tocris Cookson, Inc., 16144 Westwoods Business Park, Ellisville, MO 63021–4500, made application to the Drug Enforcement Administration to be registered as an importer of Tetrahydrocannabinols (7370), a basic class of controlled. Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substances may file