

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Application**

Pursuant to 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 21 U.S.C. 952(a)(2)(B) 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 21 CFR 1301.34(a), this is notice that on March 22, 2006, Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in Schedule II:

Drug	Schedule
Coca Leaves (9040) .....	II
Raw Opium (9600) .....	II
Poppy Straw (9650) .....	II
Concentrate of Poppy Straw (9670).	II

The company plans to import the listed controlled substances to manufacture bulk controlled substance intermediates for sale to its customers.

Any manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than September 14, 2006.

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 published in the

Federal Register on September 23, 1975, (40 FR 43745-46), all applicants for registration to import a basic class of any controlled substance listed 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(b), (c), (d), (e) and (f) are satisfied.

Dated: August 7, 2006.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E6-13329 Filed 8-14-06; 8:45 am]  
BILLING CODE 4410-09-P

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Application**

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 21, 2006, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island, 02816, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed in Schedule I and II:

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Methylphenidate (1724) .....	II
Codeine (9050) .....	II
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Thebaine (9333) .....	II
Noroxymorphone (9668) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to dosage form manufacturers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration,

Washington, DC 20537, Attention: DEA Federal Register Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than October 16, 2006.

Dated: August 7, 2006.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. E6-13326 Filed 8-14-06; 8:45 am]  
BILLING CODE 4410-09-P

**DEPARTMENT OF JUSTICE**

**Office of Justice Programs**

[OMB Number 1121-0271]

**Bureau of Justice Assistance; Agency Information Collection Activities; Proposed Collection; Comments Requested**

**ACTION:** 60-Day Notice of Information Collection Under Review—Extension of currently approved collection.

Bureau of Justice Assistance Application Form: *Project Safe Neighborhood Semi-Annual Researcher Reporting Form*.

The Department of Justice, Office of Justice Programs, Bureau of Justice Assistance, has submitted the following information collection request for review and clearance in accordance with the Paperwork Reduction Act of 1995. This proposed information collection is published to obtain comments from the public and affected agencies.

Comments are encouraged and will be accepted for "sixty days" until October 16, 2006. If you have additional comments, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact M. Pressley at 202-353-8643 or 1-866-859-2687, Bureau of Justice Assistance, Office of Justice Programs, U.S. Department of Justice, 810 7th Street, NW., Washington, DC 20531.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility;