

form would allow applicants for employment with the Department of Justice who do not have access to the Internet to provide the required personal and experience information and job specific criteria in a format that can be scanned into the electronic recruitment module that automatically rates and ranks applicants.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* 1000 responses are estimated annually with an average of thirty minutes per response.

(6) *An estimate of the total public burden (in hours) associated with the collection:* 500 hours annually.

If additional information is required contact: Robert B. Briggs, Department Clearance Officer, Information Management and Security Staff, Justice Management Division, United States Department of Justice, Patrick Henry Building, Suite 1600, 601 D Street, NW, Washington, D.C. 20004.

Dated: August 28, 2001.

Robert B. Briggs,

Department Clearance Officer, United States Department of Justice.

[FR Doc. 01-22310 Filed 9-5-01; 8:45 am]

BILLING CODE 4410-AR-M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 6, 2001, and published in the **Federal Register** on April 17, 2001, (66 FR 19796), Novartis Pharmaceutical Corporation, 59 Route 10, East Hanover, New Jersey 07936, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of methylphenidate (1724), a basic class of controlled substance listed in Schedule II.

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

DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Novartis Pharmaceutical Corporation to manufacture methylphenidate is consistent with the public interest at this time. DEA has investigated Novartis Pharmaceutical Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's

records, verification of the company's compliance with state and local laws, and a review of the company's background and history.

Therefore, pursuant to 21 U.S.C. 823 and 28 CFR 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 class of controlled substance listed above is granted.

Dated: August 23, 2001.

Laura M. Nagel,

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

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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 February 9, 2001, Chattam Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal and by letter dated June 11, 2001, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
N-Ethylamphetamine (1475)	I
4-Methoxyamphetamine (7411)	I
2,5-Dimethoxyamphetamine (7396)	I
Difenoxim (9168)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Codeine (9050)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Morphine (9300)	II
Thebaine (9333)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	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 (CCR), and must be filed no later than November 5, 2001.

Dated: August 24, 2001.

Laura M. Nagel,

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

[FR Doc. 01-22326 Filed 9-5-01; 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 January 31, 2001, Houba Inc., P.O. Box 190, 16235 State Road 17, Culver, Indiana 46511, made application by to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Opium raw (9600)	II
Opium poppy (9650)	II
Poppy straw concentrate (9670) ..	II

The firm plans to import the controlled substances to use in the manufacture of active pharmaceutical ingredients.

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