

DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated April 3, 2003, and published in the **Federal Register** on April 15, 2003, (68 FR 18261), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration 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
Difenoxin (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
Dextropropoxyphene, bulk, (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances to produce products for distribution to its 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 Chattem Chemicals, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Chattem Chemicals, Inc. to ensure that the company's registration is consistent with the public interest. This investigation has included inspection and testing of the company's physical security systems, 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 classes of controlled substances listed is granted.

Dated: July 22, 2003.

Laura M. Nagel,

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

[FR Doc. 03-19815 Filed 8-04-03; 8:45 am]

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DEPARTMENT OF JUSTICE**Drug Enforcement Administration****Manufacturer of Controlled Substances; Notice of Registration**

By Notice dated March 14, 2003, and published in the **Federal Register** on April 2, 2003, (68 FR 16089), Mallinckrodt, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Codeine-N-oxide (9053)	I
Dihydromorphine (9145)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Heroin (9200)	I
Morphine-N-oxide (9307)	I
Nicomorphine (9312)	I
Normorphine (9313)	I
Norlevorphanol (9634)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Etorphine HCL (9059)	II
Dihydrocodeine (9120)	II
Hydromorphone (9150)	II
Oxycodone (9143)	II
Diphenoxylate (9170)	II
Benzoylcegonine (9180)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II
Metopon (9260)	II
Dextropropoxyphene, bulk (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Opium extracts (9610)	II
Opium fluid extract (9620)	II
Opium fluid extract (9620)	II
Opium tincture (9630)	II
Opium, powdered (9639)	II
Opium, granulated (9640)	II
Levo-alphaacetylmethadol (9648)	II
Oxymorphone (9652)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The above cited Notice of Application contained an error in that the drug code

for Oxycodone was listed as 9160 rather than 9143.

The firm plans to manufacture the listed controlled substances for internal use and for sale to other companies.

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 Mallinckrodt, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Mallinckrodt, Inc. to ensure that the company's registration is consistent with the public interest. This investigation has included inspection and testing of the company's physical security systems, 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 classes of controlled substances listed is granted.

Dated: July 22, 2003.

Laura M. Nagel,

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

[FR Doc. 03-19814 Filed 8-4-03; 8:45 am]

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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 April 2, 2003, Penick, Corporation, 158 Mount Olivet Avenue, Newark, New Jersey 07114, 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
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Diphenoxylate (9170)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II

The firm plans to manufacture bulk controlled substances and non-controlled flavor extracts.

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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than October 6, 2003.

Dated: July 22, 2003.

Laura M. Nagel,

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

[FR Doc. 03-19809 Filed 8-4-03; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturers of Controlled Substances; Notice of Registration

By Notice dated March 14, 2003, and published in the **Federal Register** on April 2, 2003, (68 FR 16091) Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of Schedule I and II controlled substances listed below:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Methylphenidate (1724)	II
Codeine (9050)	II
Dihydrocodeine (9102)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Thebaine (9333)	II
Noroxymorphone (9668)	II
Fentanyl (9801)	II

The firm plans to produce bulk products for conversion and distribution to its 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 Rhodes Technologies to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Rhodes Technologies to

ensure that the company's registration is consistent with the public interest. This investigation has included inspection and testing of the company's physical security systems, 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 classes of controlled substances listed is granted.

Dated: July 22, 2003.

Laura M. Nagel,

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

[FR Doc. 03-19812 Filed 8-4-03; 8:45 am]

BILLING CODE 4410-09-M

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 May 23, 2003, Wildlife Laboratories, Inc., 1401 Duff Drive, Suite 600, Fort Collins, Colorado 80524, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Carfentanyl (9743), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture the listed controlled substance 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 Assistance Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than October 6, 2003.

Dated: July 22, 2003.

Laura M. Nagel,

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

[FR Doc. 03-19810 Filed 8-4-03; 8:45 am]

BILLING CODE 4410-09-M

DEPARTMENT OF JUSTICE

Federal Bureau of Investigations

Agency Information Collection Activities: Proposed Collection; Comments Requested

ACTION: 60-Day Notice of Information Collection Under Review: FBI Questionnaire for National Security Positions.

The Department of Justice (DOJ), Federal Bureau of Investigations (FBI), has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The 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 6, 2003. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Ms. Gari L. Carter, Supervisory Applicant Case Specialist, Bureau Applicant Employment Unit, Applicant Processing Section, Administrative Services Division, Federal Bureau of Investigation, Washington, DC 20535, (202) 324-0801.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments 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 functions of the agency, including whether the information will have practical utility;

(2) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of this information collection:

(1) *Type of Information Collection:* New Collection.