

Drug	Schedule	Drug	Schedule
Cathinone (1235)	I	Alpha-Methylfentanyl (9814)	I
Methcathinone (1237)	I	Acetyl-alpha-methylfentanyl (9815)	I
N-Ethylamphetamine (1475)	I	Beta-hydroxyfentanyl (9830)	I
N,N-Dimethylamphetamine (1480)	I	Beta-hydroxy-3-methylfentanyl (9831)	I
Aminorex (1585)	I	Alpha-Methylthiofentanyl (9832)	I
4-7-Methylaminorex (cis isomer) (1590)	I	3-Methylthiofentanyl (9833)	I
Gamma hydroxybutyric acid (2010)	I	Thiofentanyl (9835)	I
Methaqualone (2565)	I	Amphetamine (1100)	II
Alpha-Ethyltryptamine (7249)	I	Methamphetamine (1105)	II
Lysergic acid diethylamide (7315)	I	Phenmetrazine (1631)	II
Tetrahydrocannabinols (7370)	I	Methylphenidate (1724)	II
Mescaline (7381)	I	Ambobarbital (2125)	II
3,4,5-Trimethoxyamphetamine (7390)	I	Pentobarbital (2270)	II
4-Bromo-2,5-dimethoxyamphetamine (7391)	I	Secobarbital (2315)	II
4-Bromo-2,5-dimethoxyphenethylamine (7392)	I	Glutethimide (2550)	II
4-Methyl-2,5-dimethoxyamphetamine (7395)	I	Nabilone (7379)	II-1
2,5-Dimethoxyamphetamine (7396)	I	Phenylcyclohexylamine (7460)	II
2,5-Dimethoxy-4-ethylamphetamine (7399)	I	Phencyclidine (7471)	II-1
3,4-Methylenedioxyamphetamine (7400)	I	Piperidinocyclohexanecarbonitrile (8603)	II
5-Methoxy-3,4-methylenedioxyamphetamine (7401)	I	Alphaprodine (9010)	II
N-Hydroxy-3,4-methylenedioxyamphetamine (7402)	I	Cocaine (9041)	II
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I	Codeine (9050)	II
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I	Dihydrocodeine (9120)	II
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I	Oxycodone (9143)	II
4-Methoxyamphetamine (7411)	I	Hydromorphone (9150)	II
Bufotenine (7433)	I	Diphenoxylate (9170)	II
Diethyltryptamine (7434)	I	Benzoylcegonine (9180)	II
Dimethyltryptamine (7435)	I	Ethylmorphine (9190)	II
Psilocybin (7437)	I	Hydrocodone (9193)	II
Psilocyn (7438)	I	Levomethorphan (9210)	II
Acetyldihydrocodeine (9051)	I	Levorphanol (9220)	II
Benzylmorphine (9052)	I	Isomethadone (9226)	II
Codeine-N-oxide (9053)	I	Meperidine (9230)	II
Dihydromorphone (9145)	I	Methadone (9250)	II
Heroin (9200)	I	Methadone-intermediate (9254)	II
Hydromorphinol (9301)	I	Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Methyldihydromorphone (9304)	I	Morphine (9300)	II
Morphine-N-oxide (9307)	I	Thebaine (9333)	II
Normorphine (9313)	I	Levo-alphaacetylmethadol (9648)	II
Pholcodine (9314)	I	Oxymorphone (9652)	II
Acetylmethadol (9601)	I	Noroxymorphone (9668)	II
Allylprodine (9602)	I	Racemethorphan (9732)	II
Alphacetylmethadol except levo-alphaacetylmethadol (9603)	I	Alfentanil (9737)	II
Alphameprodine (9604)	I	Sufentanil (9740)	II
Alphamethadol (9605)	I	Fentanyl (9801)	II
Betacetylmethadol (9607)	I		
Betameprodine (9608)	I		
Betamethadol (9609)	I		
Betaprodine (9611)	I		
Hydroxypethidine (9627)	I		
Noracymethadol (9633)	I		
Norlevorphanol (9634)	I		
Normethadone (9635)	I		
Trimeperidine (9646)	I		
Phenomorphan (9647)	I		
Para-Fluorofentanyl (9812)	I		
3-Methylfentanyl (9813)	I		

The company plans to manufacture small quantities of the listed controlled substances to make reference standards for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Cerilliant Corporation to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Cerilliant Corporation to ensure that the company's registration is consistent with the public interest. The 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 22, 2005.

William J. Walker,

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

[FR Doc. 05-4205 Filed 3-3-05; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated September 28, 2004 and published in the **Federal Register** on October 14, 2004, (69 FR 61044), Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78664, 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 Schedules I and II:

Drug	Schedule
Cathinone (1235)	I
Methcathinone (1237)	I
N-Ethylamphetamine (1475)	I
Gamma hydroxybutyric acid (2010)	I
Ibogaine (7260)	I
Tetrahydrocannabinols (7370)	I
Mescaline (7381)	I
4-Bromo-2,5-dimethoxyamphetamine (7391)	I
4-Bromo-2,5-dimethoxyphenethylamine (7392)	I
4-Methyl-2,5-dimethoxyamphetamine (7395)	I
2,5-Dimethoxyamphetamine (7396)	I
3,4-Methylenedioxyamphetamine (7400)	I
3,4-Methylenedioxy-N-ethylamphetamine (7404)	I
4-Methoxyamphetamine (7411)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
Etorphine (9056)	I
Heroin (9200)	I
Pholcodine (9314)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Ambobarbital (2125)	II
Pentobarbital (2270)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II

Drug	Schedule
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoyllecgonine (9180)	II
Ethylmorphine (9190)	II
Meperidine (9230)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648) ..	II
Oxymorphone (9652)	II

The company plans to import small quantities of the listed controlled substances for the manufacture of analytical reference standards. No comments or objections have been received. DEA has considered the factors in 21 U.S.C. sections 823(a) and 952(a) and determined that the registration of Cerilliant Corporation to import the basic classes of 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 Cerilliant Corporation to ensure that the company's registration is consistent with the public interest. The 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. sections 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: February 22, 2005.
William J. Walker,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. 05-4234 Filed 3-3-05; 8:45 am]
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DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances; Notice of Registration

By notice dated September 28, 2004, and published in the **Federal Register** on October 13, 2004, (69 FR 60898), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes

of controlled substances listed in Schedules I and II:

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 company plans to manufacture the listed controlled substances in bulk for distribution to its customers. No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Chattem Chemicals, Inc. to manufacture the listed basic classes of 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. The 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 23, 2005.
William J. Walker,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. 05-4230 Filed 3-3-05; 8:45 am]
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DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 8, 2004, and published in the **Federal Register** on November 22, 2004, (69 FR 67962-

67963), ISP, Freetown Fine Chemicals, Inc., 238 South Main Street, Assonet, Massachusetts 02702, 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 Schedules I and II:

Drug	Schedule
2,5-Dimethoxyamphetamine (7396).	I
Amphetamine (1100)	II
Phenylacetone (8501)	II

The company plans to manufacture phenylacetone to be used in the manufacture of the amphetamine. The bulk 2, 5-dimethoxyamphetamine will be used for conversion into non-controlled substances. No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of ISP, Freetown Fine Chemicals, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated ISP, Freetown Fine Chemicals, Inc. to ensure that the company's registration is consistent with the public interest. The 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 in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 22, 2005.
William J. Walker,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
 [FR Doc. 05-4197 Filed 3-3-05; 8:45 am]
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DEPARTMENT OF JUSTICE
Drug Enforcement Administration
Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 1, 2004, and published in the **Federal Register** on November 10, 2004, (69 FR 65229), Guilford Pharmaceuticals, Inc., 6611 Tributary Street, Baltimore, Maryland 21224, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk