

Drug	Schedule
Methadone (9250)	II
Dextropropoxyphene, bulk, (9273)	II
Morphine (9300)	II

The company plans to manufacture the listed controlled substances as bulk reagents for use in drug abuse testing.

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 Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than September 4, 2007.

Dated: June 26, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-12973 Filed 7-3-07; 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 February 14, 2007, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, 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 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
Oxymorphone (9652)	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 Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than September 4, 2007.

Dated: June 26, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-12974 Filed 7-3-07; 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 9, 2007, Siegfried (USA), Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 schedule I and II:

Drug	Schedule
Dihydromorphine (9145)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Methadone intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Oxymorphone (9652)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

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 Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than September 4, 2007.

Dated: June 26, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-12975 Filed 7-3-07; 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 March 29, 2007, Sigma Aldrich Research Biochemicals, Inc., 1-3 Strathmore Road, Natick, Massachusetts 01760, 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 schedule I and II:

Drug	Schedule
Cathinone (1235)	I
Methcathinone (1237)	I
Aminorex (1585)	I
Alpha-ethyltryptamine (7249)	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
4-Bromo-2,5-dimethoxyamphetamine (7391)	I
4-Bromo-2,5-dimethoxyphenethylamine (7392)	I
2,5-Dimethoxyamphetamine (7396)	I
3,4-Methylenedioxyamphetamine (7400)	I
N-Hydroxy-3,4-methylenedioxyamphetamine (7402)	I

Drug	Schedule
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxy methamphetamine (MDMA) (7405).	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (TCP) (7470).	I
1-Benzylpiperazine (BZP) (7493)	I
Heroin (9200)	I
Normorphine (9313)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Nabilone (7379)	II
1-Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Ecgonine (9180)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Metazocine (9240)	II
Methadone (9250)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphaacetylmethadol (9648) ..	II
Carfentanil (9743)	II
Fentanyl (9801)	II

The company plans to manufacture reference standards.

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 Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), Washington, DC 20537, or any being sent via express mail should be sent to Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 2401 Jefferson Davis Highway, Alexandria, Virginia 22301; and must be filed no later than September 4, 2007.

Dated: June 26, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7-12971 Filed 7-3-07; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-61,298]

American Manufacturing International, Inc. Patterson, NJ; Notice of Negative Determination Regarding Application for Reconsideration; Correction

ACTION: Notice; correction.

SUMMARY: The Department of Labor, Employment and Training Administration published a document in the **Federal Register** on May 30, 2007, concerning a Notice of Negative Determination Regarding Application for Reconsideration. The office restructured a paragraph for clarification purposes.

Correction

In the **Federal Register** of May 30, 2007, in FR Doc. E7-10310, on page 30029, in the first and second column, correct the "text" caption to read:

Since the total number of separated workers was one during the relevant period, workers of American Manufacturing International, Inc., Patterson, New Jersey do not meet the group eligibility requirements for trade adjustment assistance.

Signed in Washington, DC, this 27th day of June 2007.

Elliott S. Kushner,

Certifying Officer, Division of Trade Adjustment Assistance.

[FR Doc. E7-12914 Filed 7-3-07; 8:45 am]

BILLING CODE 4510-FN-P

DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-60,876 and TA-W-60,876B]

Armstrong Wood Products, Inc., Armstrong Hardwood Flooring Company, Parquet Flooring Department and Pattern Plus Flooring Department, Oneida, TN; Notice of Revised Determination on Reconsideration; Correction

ACTION: Notice; correction.

SUMMARY: The Department of Labor, Employment and Training Administration published a document in the **Federal Register** on June 14, 2007, concerning a Notice of Revised Determination on Reconsideration. The document contained an incorrect date.

Correction.

In the **Federal Register** of June 14, 2007, in FR Doc. E7-11479, on page 32914, in the third column, correct the "text" caption to read:

All workers of Armstrong Wood Products, Inc., Armstrong Hardwood Flooring Company, Parquet Flooring Department, Oneida, Tennessee [TA-W-60,876], who became totally or partially separated from employment on or after March 17, 2007 through two years from the date of this certification, are eligible to apply for adjustment assistance under Section 223 of the Trade Act of 1974, are also eligible to apply for alternative trade adjustment assistance under Section 246 of the Trade Act of 1974.

and

All workers of Armstrong Wood Products, Inc., Armstrong Hardwood Flooring Company, Pattern Plus Flooring Department, Oneida, Tennessee [TA-W-60,876B], who became totally or partially separated from employment on or after January 31, 2006 through two years from the date of this certification, are eligible to apply for adjustment assistance under Section 223 of the Trade Act of 1974, and are also eligible to apply for alternative trade adjustment assistance under Section 246 of the Trade Act of 1974.

Signed in Washington DC this 27th day of June 2007.

Elliott S. Kushner,

Certifying Officer, Division of Trade Adjustment Assistance.

[FR Doc. E7-12911 Filed 7-3-07; 8:45 am]

BILLING CODE 4510-FN-P

DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-61,281]

Form Tech Industries, LLC, Canal Fulton, OH; Notice of Affirmative Determination Regarding Application for Reconsideration

By application dated June 12, 2007, the United Automobile, Aerospace & Agricultural Implement Workers of America, Local 1112 requested administrative reconsideration of the Department of Labor's Notice of Negative Determination Regarding Eligibility to Apply for Worker Adjustment Assistance, applicable to workers and former workers of the subject firm. The determination was signed on May 9, 2007 and published in the **Federal Register** on May 24, 2007 (72 FR 29182).

The initial investigation resulted in a negative determination based on the finding that imports of machine parts, such as shafts and sheaves for CVT