

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 Administrator, 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), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than April 21, 2008.

Dated: February 13, 2008.

**Joseph T. Rannazzisi,**

*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 § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 10, 2008, Roche Diagnostics Operations, Inc., Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (THC) (7370).	I
Alphamethadol (9605) .....	I
Hydromorphone (9150) .....	II
Benzoyllecgonine (9180) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II

The company plans to manufacture small quantities of the listed controlled substances for use in diagnostic products.

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 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), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than April 21, 2008.

Dated: February 12, 2008.

**Joseph T. Rannazzisi,**

*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 Registration**

By Notice dated September 21, 2007, and published in the **Federal Register** on September 27, 2007, (72 FR 54930), ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, 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
2,5-Dimethoxyamphetamine (7396).	I
Amphetamine (1100) .....	II
Phenylacetone (8501) .....	II

The company plans to manufacture Phenylacetone to be used in the manufacture of Amphetamine for distribution to its customers. 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 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 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 12, 2008.

**Joseph T. Rannazzisi,**

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

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**DEPARTMENT OF LABOR**

**Employment and Training Administration**

[TA-W-62,422]

**Curtain & Drapery Fashions Including On-Site Leased Workers From Paychex Business Solutions, Lowell, NC; Amended Certification Regarding Eligibility To Apply for Worker Adjustment Assistance and Alternative Trade Adjustment Assistance**

In accordance with section 223 of the Trade Act of 1974 (19 U.S.C. 2273), and section 246 of the Trade Act of 1974 (26 U.S.C. 2813), as amended, the Department of Labor issued a Certification of Eligibility to Apply for Worker Adjustment Assistance and Alternative Trade Adjustment Assistance on December 19, 2007, applicable to workers of Curtain & Drapery Fashions, Lowell, North Carolina. The notice was published in the **Federal Register** on January 16, 2008 (72 FR 2943).

At the request of the State agency, the Department reviewed the certification for workers of the subject firm. The workers were engaged in the production of curtains, bedspreads and comforters.

New information shows that leased workers of Paychex Business Solutions were employed on-site at the Lowell, North Carolina location of Curtain & Drapery Fashions. The Department has determined that these workers were sufficiently under the control of Curtain & Drapery Fashions to be considered leased workers.

Based on these findings, the Department is amending this certification to include leased workers of Paychex Business Solutions working on-site at the Lowell, North Carolina location of the subject firm.

The intent of the Department's certification is to include all workers employed at Curtain & Drapery Fashions, Lowell, North Carolina who were adversely impacted by increased customer imports of curtains, bedspreads and comforters.