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: May 22, 2000.

John H. King,

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

[FR Doc. 00–13785 Filed 6–1–00; 8:45 am]

BILLING CODE 4410–09–M

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 December 22, 1999, Lonza Riverside, 900 River Road, Conshohocken, Pennsylvania 19428, made application by renewal which was received for processing April 12, 2000, to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Amphetamine (1100)	II
Phenylacetone (8501)	II

The firm 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 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than August 1, 2000.

Dated: May 24, 2000.

John H. King,

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

[FR Doc. 00–13786 Filed 6–1–00; 8:45 am]

BILLING CODE 4410–09–M

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 April 14, 2000, Sigma Aldrich Research Biochemicals, Inc., Attn: Richard Milius, 1–3 Strathmore Road, Natick, Massachusetts 01760, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

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
3, 4-Methylenedioxy-N-ethylamphetamine (7404).	I
3, 4-Methylenedioxymethamphetamine (7405).	I
Psilocybin (7437)	I
1-[1-(2-Thienyl) cyclohexyl] piperidine (7470).	I
Heroin (9200)	I
Normorphine (9313)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Pentobarbital (2270)	II
Phenylcyclohexylamine (7460)	II
Phencyclidine (7471)	II
Cocaine (9041)	II
Codeine (9050)	II
Diprenorphine (9058)	II
Benzoylecgonine (9180)	II
Levomethorphan (9210)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Metazocine (9240)	II
Methadone (9250)	II
Morphine (9300)	II
Thebaine (9333)	II
Levo-alphacetylmethadol (LAAM) (9648).	II
Fentanyl (9801)	II

The firm plans to manufacture the listed controlled substances for laboratory reference standards and neurochemicals.

Any other such applicant and any person who is presently registered with DEA to manfuacture 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be file no later than August 1, 2000.

Dated: May 12, 2000.

John H. King,

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

[FR Doc. 00–13787 Filed 6–1–00; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Employment and Training Administration

[TA–W–37,303 and NAFTA–3688]

Motor Coils Manufacturing, Emporium, Pennsylvania; Notice of Revised Determination on Reconsideration

By application dated March 24, 2000, petitioners requested administrative reconsideration of the Department's denial of Trade Adjustment Assistance (TAA) petition number TA–W–37,303 and North American Free Trade Agreement-Transitional Adjustment Assistance (NAFTA–TAA) petition number NAFTA–3688, for workers and former workers of Motor Coils Manufacturing, Emporium, Pennsylvania. The notices of negative determination were issued March 8, 2000 and published in the **Federal Register** on March 31, 2000 (65 FR 17312) and (65 FR 17313), respectively.

The March 8, 2000 denial of TAA and NAFTA–TAA for workers of the subject firm engaged in employment related to the refurbishment of traction motors for locomotives at Motor Coils Manufacturing, Emporium, Pennsylvania, was based on the finding that the workers provided a service and did not produce an article within the meaning of the group eligibility requirements of section 222 and 250(a) of the Trade Act of 1974, as amended.

The Department, on reconsideration, has learned that the workers produce an article, and all the worker group eligibility requirements of section 222 of the Trade Act of 1974 have been met. There were declines in sales or