Drug	Schedule
Amphetamine (1100)	II
Phenylacetone (8501)	II
Methypenidate (1724)	II

The firm plans to produce bulk products for finished dosage units 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 Lonza Riverside to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Lonza Riverside 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: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22324 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Notice of Registration

By Notice dated April 29, 2003, and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Organichem Corporation, 33 Riverside Avenue, Rensselaer, New York 12144, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic class of Schedule II of controlled substance listed below:

Drug	Schedule
Dextropoxyphene (9273)	II

The firm plans to manufacture bulk 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 Organichem Corporation to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Organichem Corporation 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 class of controlled substance listed is granted.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22325 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated May 2, 2003 and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Schedules I & II, for the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic Acid Diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Benzoylecogonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to import the listed controlled substances to manufacture diagnostic 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 Roche Diagnostics Corporation to import the listed 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 Roche Diagnostics Corporation on a regular basis to ensure that the company's continued registration is consistent with the public interest. This investigation 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 Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1301.34, the above firm is granted registration as an importer of the basic classes of controlled substances listed above.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22322 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated may 2, 2003, and published in the **Federal Register** on May 29, 2003, (68 FR 32089), Roche Diagnostics corporation, Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic Acid Diethylamide (7515)	I
Tetrahydrocannabinol (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Benzoylecogonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

The firm plans to manufacture small quantities of controlled substances for use in diagnostic products.

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 Rocke Diagnostics

Corporation to manufacture the listed controlled substances is consistent with the public interested at this time. DEA has investigated Roche Diagnostics Corporation to insure 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 in granted.

Dated: August 19, 2003.

Laura M. Nagel,

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

[FR Doc. 03–22323 Filed 8–29–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Employment and Training Administration

Notice of Determinations Regarding Eligibility To Apply for Worker Adjustment Assistance

In accordance with section 223 of the Trade Act of 1974, as amended, (19 U.S.C. 2273), the Department of Labor herein presents summaries of determinations regarding eligibility to apply for trade adjustment assistance for workers (TA–W) issued during the period of August 2003.

In order for an affirmative determination to be made and a certification of eligibility to apply for directly-impacted (primary) worker adjustment assistance to be issued, each of the group eligibility requirements of Section 222(a) of the Act must be met.

I. Section (a)(2)(A) all of the following must be satisfied:

A. A significant number or proportion of the workers in such workers' firm, or an appropriate subdivision of the firm, have become totally or partially separated, or are threatened to become totally or partially separated;

B. The sales or production, or both, of such firm or subdivision have decreased absolutely; and

C. Increased imports of articles like or directly competitive with articles produced by such firm or subdivision have contributed importantly to such workers' separation or threat of separation and to the decline in sales or production of such firm or subdivision; or

II. Section (a)(2)(B) both of the following must be satisfied:

A. A significant number or proportion of the workers in such workers' firm, or an appropriate subdivision of the firm, have become totally or partially separated, or are threatened to become totally or partially separated;

B. There has been a shift in production by such workers' firm or subdivision to a foreign country of articles like or directly competitive with articles which are produced by such firm or subdivision; and

C. One of the following must be satisfied:

1. The country to which the workers' firm has shifted production of the articles is a party to a free trade agreement with the United States;

2. The country to which the workers' firm has shifted production of the articles to a beneficiary country under the Andean Trade Preference Act, African Growth and Opportunity Act, or the Caribbean Basin Economic Recovery Act; or

3. There has been or is likely to be an increase in imports of articles that are like or directly competitive with articles which are or were produced by such firm or subdivision.

Also, in order for an affirmative determination to be made and a certification of eligibility to apply for worker adjustment assistance as an adversely affected secondary group to be issued, each of the group eligibility requirements of section 222(b) of the Act must be met.

(1) Significant number or proportion of the workers in the workers' firm or an appropriate subdivision of the firm have become totally or partially separated, or are threatened to become totally or partially separated;

(2) The workers' firm (or subdivision) is a supplier or downstream producer to a firm (or subdivision) that employed a group of workers who received a certification of eligibility to apply for trade adjustment assistance benefits and such supply or production is related to the article that was the basis for such certification; and

(3) Either—(A) The workers' firm is a supplier and the component parts it supplied for the firm (or subdivision) described in paragraph (2) accounted for at least 20 percent of the production or sales of the workers' firm; or

(B) A loss or business by the workers' firm with the firm (or subdivision) described in paragraph (2) contributed importantly to the workers' separation or threat of separation.

Negative Determinations for Worker Adjustment Assistance

In the following case, the investigation revealed that the criteria for eligibility have not been met for the reasons specified.

The investigation revealed that criterion (a)(2)(A)(I.C.) (Increased imports) and (a)(2)(B)(II.B.) (No shift in production to a foreign country) have not been met.

TA–W–51,958; UTI Star Guide, Arvada, CO

TA–W–52,093; Mendocino Forest Products, Fort Bragg, CA

TA–W–52,149; OEC Medical Systems, Inc., d/b/a GE OEC Medical Systems, Inc., a subsidiary of The General Electric Co., Warsaw, IN

TA–W–52,215; Cooper B-Line, Portland, OR

TA–W–51,918; Alstom T&D Industries, High Voltage Switchgear Div., Charleroi, PA

TA–W–51,995; Occidental Chemical Corp., New Owner—Elementis Chromium LP, a subsidiary of Elementis PLC, Castle Hayne, NC

TA–W–52,267; Ken-Bar Manufacturing Co., Baldwin, GA

TA–W–52,292; Manning Lighting, Sheboygan, WI

TA–W–52,297; Intermet, Radford Foundry, Radford, VA

TA–W–52,327; NIBCO, Inc., Central Tooling Services Center, Elkhart, IN

TA–W–51,800; Meridian Automotive Systems, Exterior Composites Div., Centralia, IL

TA–W–51,838; Rio Grande Forest Products, Inc., Espanola, NM

TA–W–52,128 & A, B; Control Engineering Co., Pellston, MI, Harbor Springs, MI and Boyne City, MI

TA–W–52,171; Read-Rite Corp., Fremont, CA

TA–W–52,293; Hilti North America, a Div. of Hilti Corp., Plant 5, Tulsa, OK

The workers firm does not produce an article as required for certification under section 222 of the Trade Act of 1974.

TA–W–52,003; Menlo Logistics, Inc., d/b/a Menlo Worldwide Logistics, a wholly owned subsidiary of Menlo Worldwide, LLC, Edison, NJ

TA–W–52,357; Motorola, Inc., Personal Communications Sector (PCS), Libertyville, IL

TA–W–52,255; Solectron Technology, Inc., Charlotte, NC

TA–W–52,403; Jones Equipment, Inc., Missoula, MT

TA–W–51,970; Northwest Airlines, Inc., St. Paul, MN