

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: July 28, 2004.

William J. Walker,

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

[FR Doc. 04-18171 Filed 8-9-04; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 5, 2004, and published in the **Federal Register** on March 15, 2004, (69 FR 12182), Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances:

Drug	Schedule
Cocaine (9041)	II
Benzoylcegonine (9180)	II

The company plans to manufacture bulk controlled substances 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 Stepan Company to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Stepan Company 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: July 28, 2004.

Joseph T. Rannazzisi,

Deputy Director, 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 21 CFR 1301.33(a), this is notice that on April 29, 2004, Syva Company, Dade Behring Inc., Regulatory Affairs Dept #1-310, 20400 Mariani Avenue, Cupertino, California 95014, 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, and by letter dated July 6, 2004, to modify its name to Dade Behring, Inc.

Drug	Schedule
Tetrahydrocannabinols (7370) ...	I
Ecgonine (9180)	II
Morphine (9300)	II

The company plans to produce bulk products used for the manufacture of reagents and drug calibrator/controls, DEA exempt products.

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 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, DC 20537, Attention: DEA Federal Register Representative (CCD) and must be filed no later than October 12, 2004.

Dated: July 21, 2004.

William J. Walker,

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

[FR Doc. 04-18181 Filed 8-9-04; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-55,311]

Butler Manufacturing Company/Bluescope Steel, Galesburg, Illinois; Notice of Termination of Investigation

Pursuant to Section 221 of the Trade Act of 1974, as amended, an investigation was initiated on July 23, 2004 in response to a worker petition filed by the United Steelworkers of America, Local 2629 on behalf of workers of Butler Manufacturing Company/Bluescope Steel, Galesburg, Illinois.

The petitioning group of workers is covered by an earlier petition filed on July 22, 2004 (TA-W-55,290) that is the subject of an ongoing investigation for which a determination has not yet been issued. Further investigation in this case would duplicate efforts and serve no purpose; therefore the investigation under this petition has been terminated.

Signed at Washington, DC this 27th day of July 2004.

Linda G. Poole,

Certifying Officer, Division of Trade Adjustment Assistance.

[FR Doc. 04-18228 Filed 8-9-04; 8:45 am]

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DEPARTMENT OF LABOR

Employment and Training Administration

[TA-W-55,172]

Cardinal Health Medical Products & Services Division, El Paso, Texas; Notice of Termination of Investigation

Pursuant to Section 221 of the Trade Act of 1974, as amended, an investigation was initiated on June 30, 2004, in response to a worker petition filed by a company official on behalf of workers at Cardinal Health, Medical Products & Services Division, El Paso, Texas.

The petitioner has requested that the petition be withdrawn. Consequently, further investigation would serve no purpose and the investigation has been terminated.

Signed at Washington, DC, this 22nd day of July, 2004.

Elliott S. Kushner,

Certifying Officer, Division of Trade Adjustment Assistance.

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