Background

In August 1999 the Commission made negative determinations upon reconsideration in its antidumping and countervailing duty investigations concerning ferrosilicon from Brazil, China, Kazakhstan, Russia, Ukraine, and Venezuela. Ferrosilicon from Brazil, China, Kazakhstan, Russia, Ukraine, and Venezuela, Inv. Nos. 303–TA–23, 731–TA–566–570, 731–TA–641 (Final) (Reconsideration), USITC Pub. 3218 (Aug. 1999). The Commission's determinations were appealed to the U.S. Court of International Trade (CIT). On February 21, 2002, the CIT remanded the matter to the Commission for further proceedings. *Elkem Metals Co.* v. *United States,* 193 F. Supp.2d 1314 (Ct. Int'l Trade 2002). On remand, the Commission conducted further proceedings. In September 2002 it reached negative determinations on remand. Ferrosilicon from Brazil, China, Kazakhstan, Russia, Ukraine, and Venezuela, Inv. Nos. 303–TA–23, 731–TA–566–570, and 731–TA–631 (Final) (Reconsideration) (Remand), USITC Pub. 3531 (Sept. 2002). On March 18, 2003, the CIT issued an opinion concerning the Commission's determinations on remand which affirmed the Commission in part and remanded in part for further proceedings. *Elkem Metals Co.* v. *United States,* slip op. 03–66 (Ct. Int'l Trade June 18, 2003) (''Elkem II'').

Written Submissions

The Commission is not reopening the record in the second remand proceeding for submission of new factual information. It will, however, permit the parties to file a written submission addressed to the determination the Commission should reach in its second remand determination in light of Elkem II. Parties should state the factual and legal bases for their position. This submission must be filed with the Commission no later than 14 days after publication of this notice in the **Federal Register**, shall not contain any new factual information, and shall not exceed 20 pages of textual material, double-spaced and single-sided, on stationery measuring 8½ x 11 inches.

All written submissions must conform with the provisions of section 201.8 of the Commission's rules; any submissions that contain business proprietary information (BPI) must also conform with the requirements of sections 201.6, 207.3, and 207.7 of the Commission's rules. The Commission's rules do not authorize filing of submissions with the Secretary by facsimile or electronic means, except to the extent permitted by section 201.8 of the Commission's rules, as amended, 67 FR 68036 (Nov. 8, 2002).

In accordance with sections 201.16(c) and 207.3 of the Commission's rules, each document filed by a party to the investigations must be served on all other parties to the investigations (as identified by either the public or BPI service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

Parties are also advised to consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subpart A (19 CFR part 207) for provisions of general applicability concerning written submissions to the Commission.

Participation in the Proceedings

Only those persons who were parties to the previous reconsideration proceedings (*i.e.*, persons listed on the Commission Secretary's service list) may participate as parties in the second remand proceedings.

Public Vote

The Commission will vote on the remand determinations at a public meeting scheduled to be held on Monday, August 18, 2003. The meeting is tentatively scheduled for 11 a.m.

Authority: This action is taken under the authority of title VII of the Tariff Act of 1930 as amended.

Issued: July 15, 2003.

By order of the Commission.

Marilyn R. Abbott,

Secretary.

[FR Doc. 03–18426 Filed 7–18–03; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated March 21, 2003 and published in the **Federal Register** on April 9, 2003, (68 FR 17403), Boehringer Ingelheim Chemicals, Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Phenylacetone (8501), a basic class of controlled substances listed in Schedule II.

The firm plans to import Phenylacetone for bulk manufacture of amphetamine.

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 Boehringer Ingelheim Chemicals, Inc. 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 Boehringer Ingelheim Chemicals, Inc. 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: July 2, 2003.

Laura M. Nagel,

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

[FR Doc. 03–18477 Filed 7–18–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 14, 2003, and published in the **Federal Register** on April 2, 2003, (68 FR 16088), Cedarburg Pharmaceuticals, LLC, 870 Badget Circle, Grafton, Wisconsin 53204, made application by letter to the Drug Enforcement Administration to be registered as a bulk manufacturer of dihydromorphine (9145), a basic class of controlled substance listed in Schedule I.

The firm plans to use this substance in the conversion process to produce a Schedule II controlled substance, hydromorphone.

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 Cedarburg Pharmaceuticals, LLC, to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated

Cedarburg Pharmaceuticals, LLC. 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: July 3, 2003.

Laura M. Nagel,

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

[FR Doc. 03–18481 Filed 7–18–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 21, 2003, and published in the **Federal Register** on April 9, 2003, (68 FR 17405), Dade Behring Inc., Route 896 Corporate Boulevard, Building 100, Attn: RA/QA, P.O. Box 6101, Newark, Delaware 19714, made application by letter, to the Drug Enforcement Administration to be registered as a bulk manufacturer of the basic classes of controlled substances listed below:

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

The firm plans to produce bulk products used for the manufacture or reagents and drug calibrator/controls, DEA exempt 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 Dade Behring Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Dade Behring Inc. 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 above is granted.

Dated: July 2, 2003.

Laura M. Nagel,

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

[FR Doc. 03–18479 Filed 7–18–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(1)), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in Schedule I or II and prior to issuing a regulation under Section 1002(a) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Section 1301.34 of Title 21, Code of Federal Regulation (CFR), notice is hereby given that on October 24, 2002, Noramco Inc., 500 Old Swedes Landing Road, Wilmington, Delaware 19801, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Opium, (raw) (9600)	II
Poppy Straw Concentrate	II

The firm plans to import the listed controlled substances for the bulk manufacture of other controlled substances.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substances may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing 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: Federal Register Representative, Office of Chief Counsel (CCD) and must be filed no later than September 19, 2003.

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice at 40 FR 43745–46 (September 23, 1975), all applicants for registration to import basic class of any controlled substance in Schedule I or II are and will continue to be required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration that the requirements for such registration pursuant to 21 U.S.C. 958(a), 21 U.S.C. 823(a), and 21 CFR 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: July 2, 2003.

Laura M. Nagel,

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

[FR Doc. 03–18480 Filed 7–18–03; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 11, 2003, and published in the **Federal Register** on April 2, 2003, (68 FR 16091), Organichem Corporation, 33 Riverside Avenue, Rensselar, New York 12144, 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
Amphetamine (1100)	II
Pentobarbital (2270)	II
Methylphenidate (1724)	II
Meperidine (9230)	II

The firm plans to manufacture bulk products for use internally and for distribution to its customers.

No comments or objections have been received. DEA has considered the factors in Title 21, United States Code,