INTERNATIONAL TRADE COMMISSION

[Investigation Nos. 701–TA–413 and 731–TA–913–916 and 918 (Review)]

Stainless Steel Bar From France, Germany, Italy, Korea, and The United Kingdom

AGENCY: United States International Trade Commission.

ACTION: Notice of Commission determinations to conduct full five-year reviews concerning the countervailing duty order on stainless steel bar from Italy and the antidumping duty orders on stainless steel bar from France, Germany, Italy, Korea, and the United Kingdom.

SUMMARY: The Commission hereby gives notice that it will proceed with full reviews pursuant to section 751(c)(5) of the Tariff Act of 1930 (19 U.S.C. 1675(c)(5)) to determine whether revocation of the countervailing duty order on stainless steel bar from Italy and the antidumping duty orders on stainless steel bar from France, Germany, Italy, Korea, and the United Kingdom would be likely to lead to continuation or recurrence of material injury within a reasonably foreseeable time. A schedule for the reviews will be established and announced at a later date. For further information concerning the conduct of these reviews and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

EFFECTIVE DATE: May 7, 2007.

FOR FURTHER INFORMATION CONTACT: Mary Messer (202–205–3193), Office of Investigations, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436. Hearing-impaired persons can obtain information on this matter by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*). The public record for these reviews may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov.*

SUPPLEMENTARY INFORMATION: On May 7, 2007, the Commission determined that it should proceed to full reviews in the subject five-year reviews pursuant to section 751(c)(5) of the Act. The Commission found that the domestic interested party group response to its notice of institution (72 FR 4293, January 30, 2007) was adequate and that the respondent interested party group responses with respect to France, Germany, and the United Kingdom were adequate and decided to conduct full reviews with respect to the antidumping duty orders concerning stainless steel bar from France, Germany, and the United Kingdom. The Commission found that the respondent interested party group responses with respect to Italy and Korea were inadequate.[1] However, the Commission determined to conduct full reviews concerning the countervailing duty order on stainless steel bar from Italy and the antidumping duty orders on stainless steel bar from Italy and Korea to promote administrative efficiency in light of its decision to conduct full reviews with respect to the orders concerning stainless steel bar from France, Germany, and the United Kingdom. A record of the Commissioners' votes, the Commission's statement on adequacy, and any individual Commissioner's statements will be available from the Office of the Secretary and at the Commission's Web site.

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.62 of the Commission's rules.

By order of the Commission.

Issued: May 14, 2007.

William R. Bishop,

Acting Secretary to the Commission.

[FR Doc. E7–9560 Filed 5–17–07; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 23, 2007, and published in the **Federal Register** on January 30, 2007, (72 FR 4296), American Radiolabeled Chemical, Inc., 101 Arc Drive, St. Louis, Missouri 63146, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
1-[1-(2–Thienyl) cyclohexyl] piperidine; TCP (7470).	I
Normorphine (9313)	I
Dextropropoxyphene, bulk (non-dosage form) (9273).	II

The company plans to manufacture small quantities of the listed controlled substances as radiolabeled compounds for biochemical research.

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 American Radiolabeled Chemical, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated American Radiolabeled Chemical, Inc. 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: May 14, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–9611 Filed 5–17–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated February 14, 2007 and published in the **Federal Register** on February 22, 2007, (72 FR 8017), Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Phenylacetone (8501), a basic class of controlled substance listed in schedule II.

The company plans to import Phenylacetone for use as a precursor in the manufacture of amphetamines only.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of

[1] Commissioners Lane and Williamson dissented with respect to the adequacy of the Italian respondent interested party group response, finding that the Italian respondent interested party group response was adequate.

Cambrex Charles City, Inc. to import the basic class of controlled substance 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 Cambrex Charles City, Inc. 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. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: May 14, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–9609 Filed 5–17–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 25, 2006, and published in the **Federal Register** on November 1, 2006, (71 FR 64298), Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616, made application by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Sufentanil (9740), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substances in bulk 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 Cambrex Charles City, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Cambrex Charles City, Inc. 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: May 14, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–9625 Filed 5–17–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated January 16, 2007, and published in the **Federal Register** on January 23, 2007, (72 FR 2907), Cedarburg Pharmaceuticals, Inc., 870 Badger Circle, Grafton, Wisconsin 53024, 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 schedule I and II:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Sufentanil (9740)	II
Fentanyl (9801)	II
Remifentanil (9739)	II

The company plans to manufacture the listed controlled substances in bulk 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 Cedarburg Pharmaceuticals, Inc. to manufacture the listed basic class of controlled substance is consistent with the public interest at this time. DEA has investigated Cedarburg Pharmaceuticals, Inc. 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: May 14, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–9642 Filed 5–17–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 21, 2006, and published in the **Federal Register** on December 1, 2006, (71 FR 69589–69590), Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, made application by renewal and by letter to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedule I and II:

Drug	Schedule
4–Methoxyamphetamine (7411)	I
Dihydromorphine (9145)	I
Difenoxin (9168)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Pentobarbital (2270)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

By letter dated February 15, 2007, Chattem Chemicals has withdrawn their request for N-Ethylamphetamine (1475), Secobarbital (2315), 2,5–Dimethoxyamphetamine (7396), Diphenoxylate (9170), Opium Extracts (9610), Opium Fluid Extract (9620), Opium Tincture (9630), Opium Powdered (9639), and Opium Granulated (9640).

One comment has been received; this comment has been noted and considered by DEA.