

International Trade Commission, 500 E Street SW, Washington, DC 20436, telephone 202-205-2000.

Issued: April 19, 1999.  
By order of the Commission.

**Donna R. Koehnke,**  
Secretary.

[FR Doc. 99-10535 Filed 4-26-99; 8:45 am]

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**INTERNATIONAL TRADE COMMISSION**

[Investigation No. 701-TA-378 (Final)]

**Certain Stainless Steel Plate From Korea**

**AGENCY:** United States International Trade Commission.

**ACTION:** Termination of investigation.

**SUMMARY:** On March 31, 1999, the Department of Commerce published notice in the **Federal Register** of a negative final determination of countervailable subsidies in connection with the subject investigation (64 FR 15530). Accordingly, pursuant to § 207.40(a) of the Commission's Rules of Practice and Procedure (19 CFR 207.40(a)), the countervailing duty investigation concerning certain stainless steel plate from Korea (investigation No. 701-TA-378 (Final)) is terminated.

**EFFECTIVE DATE:** March 31, 1999.

**FOR FURTHER INFORMATION CONTACT:** Woodley Timberlake (202-205-3188), Office of Investigations, U.S. International Trade Commission, 500 E Street SW, Washington, DC 20436. Hearing-impaired individuals are advised that information on this matter can be obtained 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>).

**Authority:** This investigation is being terminated under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to § 201.10 of the Commission's rules (19 CFR 201.10).

Issued: April 22, 1999.  
By order of the Commission.

**Donna R. Koehnke,**  
Secretary.

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**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(i)), 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 Regulations (CFR), notice is hereby given that on January 21, 1999, Mallinckrodt Chemical, Inc., Mallinckrodt & Second Streets, St. Louis, Missouri 63147, made application by renewal of the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Phenylacetone (8501) .....	II
Coca Leaves (9040) .....	II
Opium, raw (9600) .....	II
Opium poppy (9650) .....	II
Poppy Straw Concentrate (9670) .....	II

The firm plans to import the listed controlled substances to manufacture bulk finished products.

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance 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: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

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 a 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 1301.34(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: April 12, 1999.

**John H. King,**

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

[FR Doc. 99-10511 Filed 4-26-99; 8:45 am]

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**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 January 25, 1999, Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, 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
Cocaine (9041) .....	II
Benzoylcgonine (9180) .....	II

The firm plans to manufacture bulk controlled substances for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances 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, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (60 days from publication).