

Wang (a/k/a Alice B. Wang) of Arcadia, CA; Trendy Creations, Inc. of Chatsworth, CA; The Inspired Bagger of Dallas, TX; and House of Bags of Los Angeles, CA.

The complainant, proposed respondents, other interested parties, and members of the public are invited to file comments, not to exceed five pages in length, on any public interest issues raised by the complaint. Comments should address whether issuance of an exclusion order and/or a cease and desist order in this investigation would negatively affect the public health and welfare in the United States, competitive conditions in the United States economy, the production of like or directly competitive articles in the United States, or United States consumers.

In particular, the Commission is interested in comments that:

- (i) Explain how the articles potentially subject to the orders are used in the United States;
- (ii) Identify any public health, safety, or welfare concerns in the United States relating to the potential orders;
- (iii) Indicate the extent to which like or directly competitive articles are produced in the United States or are otherwise available in the United States, with respect to the articles potentially subject to the orders; and
- (iv) Indicate whether Complainant, Complainant's licensees, and/or third party suppliers have the capacity to replace the volume of articles potentially subject to an exclusion order and a cease and desist order within a commercially reasonable time.

Written submissions must be filed no later than by close of business, five business days after the date of publication of this notice in the **Federal Register**. There will be further opportunities for comment on the public interest after the issuance of any final initial determination in this investigation.

Persons filing written submissions must file the original document and 12 true copies thereof on or before the deadlines stated above with the Office of the Secretary. Submissions should refer to the docket number ("Docket No. 2772") in a prominent place on the cover page and/or the first page. The Commission's rules authorize filing submissions with the Secretary by facsimile or electronic means only to the extent permitted by section 201.8 of the rules (see Handbook for Electronic Filing Procedures, [http://www.usitc.gov/secretary/fed\\_reg\\_notices/rules/documents/handbook\\_on\\_electronic\\_filing.pdf](http://www.usitc.gov/secretary/fed_reg_notices/rules/documents/handbook_on_electronic_filing.pdf)). Persons with questions regarding

electronic filing should contact the Secretary (202-205-2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of sections 201.10 and 210.50(a)(4) of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.50(a)(4)).

By order of the Commission.

Issued: December 6, 2010.

**William R. Bishop,**

*Acting Secretary to the Commission.*

[FR Doc. 2010-30951 Filed 12-8-10; 8:45 am]

**BILLING CODE P**

## INTERNATIONAL TRADE COMMISSION

[Investigation No. 731-TA-841 (Second Review)]

### Non-Frozen Apple Juice Concentrate From China

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

**ACTION:** Termination of five-year review.

**SUMMARY:** The subject five-year review was initiated in October 2010 to determine whether revocation of the antidumping duty order on non-frozen apple juice concentrate from China would be likely to lead to continuation or recurrence of material injury. On November 15, 2010, the Department of Commerce published notice that it was revoking the order effective November 2, 2010, "because the domestic interested parties did not participate in this sunset review \* \* \*" (75 FR 69628). Accordingly, pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)), the subject review is terminated.

**DATES:** *Effective Date:* November 2, 2010.

**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 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 review is being terminated under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.69 of the Commission's rules (19 CFR 207.69).

By order of the Commission.

Issued: December 3, 2010.

**Marilyn R. Abbott,**

*Secretary to the Commission.*

[FR Doc. 2010-30906 Filed 12-8-10; 8:45 am]

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## DEPARTMENT OF JUSTICE

### Notice of Lodging of Consent Decree Under the Residential Lead-Based Paint Hazard Reduction Act

Notice is hereby given that on December 3, 2010 a proposed Consent Decree in *United States v. Combined Development Co. I, LLC, et al.*, Civil Action No. 1:10-cv-853 was lodged with the United States District Court for the Southern District of Ohio.

The consent decree settles claims against the owners and managers of 166 housing units in nine separate properties located in or near Cincinnati, Ohio. The claims were brought on behalf of the Environmental Protection Agency ("U.S. EPA") and the Department of Housing and Urban Development ("HUD") under the Residential Lead-Based Paint Hazard Reduction Act, 42 U.S.C. 4851 *et seq.* ("Lead Hazard Reduction Act"). The United States alleged in the complaint that the Defendants failed to make one or more of the disclosures or to complete one or more of the disclosure activities required by the Lead Hazard Reduction Act.

Under the Consent Decree, the Defendants will certify that they are complying with residential lead paint notification requirements. The Defendants will submit a plan for window replacement work and will replace all windows known to or believed to contain lead-based paint in all residential properties owned or managed by Defendants that are not certified lead-based paint free. In

addition, Defendants will abate lead-based paint hazards on friction and impact surfaces, stabilize other lead-based paint hazards, and pay an administrative penalty of \$7,500.

The Department of Justice will receive for a period of thirty (30) days from the date of this publication comments relating to the Proposed Consent Decree. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and either e-mailed to [pubcomment-ees.enrd@usdoj.gov](mailto:pubcomment-ees.enrd@usdoj.gov) or mailed to U.S. Department of Justice, Washington, DC 20044-7611 P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044-7611, and should refer to *United States v. Combined Development Co. I, LLC, et al.*, D.J. Ref. # 90-5-1-1-09435.

The Proposed Consent Decree may be examined at the Department of Housing and Urban Development, Office of General Counsel, 451 7th St. NW., Room 9262, Washington, DC 20410; at the office of the United States Attorney for the Southern District of Ohio, 303 Marconi Blvd., Suite 200, Columbus, Ohio 43215 (Attn. Assistant United States Attorney Andrew M. Malek); and at U.S. EPA Region 5, 77 W. Jackson Blvd., Chicago, IL 60604. During the public comment period, the Consent Decree may also be examined on the following Department of Justice Web site, to [http://www.usdoj.gov/enrd/Consent\\_Decrees.html](http://www.usdoj.gov/enrd/Consent_Decrees.html). A copy of the Consent Decree may also be obtained by mail from the Consent Decree Library, P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044-7611 or by faxing or e-mailing a request to Tonia Fleetwood ([tonia.fleetwood@usdoj.gov](mailto:tonia.fleetwood@usdoj.gov)), fax no. (202) 514-0097, phone confirmation number (202) 514-1547. In requesting a copy from the Consent Decree Library, please enclose a check in the amount of \$9.25 (25 cents per page reproduction cost) payable to the U.S. Treasury or, if by e-mail or fax, forward a check in that amount to the Consent Decree Library at the stated address.

**Maureen Katz,**  
Assistant Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2010-30900 Filed 12-8-10; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Application**

Pursuant to 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 21 U.S.C. 952(a)(2) 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 21 CFR 1301.34(a), this is notice that on October 19, 2010, Mylan Pharmaceuticals, Inc., 781 Chestnut Ridge Road, Morgantown, West Virginia 26505, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule II:

Drug	Schedule
Methylphenidate (1724) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Fentanyl (9801) .....	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than January 10, 2011.

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 published in

the **Federal Register** on September 23, 1975, (40 FR 43745-46), 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(b), (c), (d), (e), and (f) are satisfied.

Dated: November 29, 2010.

**Joseph T. Rannazzisi,**  
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2010-30901 Filed 12-8-10; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Importer of Controlled Substances; Notice of Registration**

By Notice dated August 3, 2010, and published in the **Federal Register** on September 1, 2010, (75 FR 53719), Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Heroin (9200) .....	I
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II

The company plans to import these controlled substances for the manufacture of reference standards.

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 Alltech Associates, Inc. to import the basic classes of 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. DEA has investigated Alltech Associates, 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