Date, and significant changes, if any, that are likely to occur within a reasonably foreseeable time. Supply conditions to consider include technology; production methods; development efforts; ability to increase production (including the shift of production facilities used for other products and the use, cost, or availability of major inputs into production); and factors related to the ability to shift supply among different national markets (including barriers to importation in foreign markets or changes in market demand abroad). Demand conditions to consider include end uses and applications; the existence and availability of substitute products; and the level of competition among the *Domestic Like Product* produced in the United States, *Subject Merchandise* produced in the *Subject Countries*, and such merchandise from other countries.

(11) (*Optional*) A statement of whether you agree with the above definitions of the *Domestic Like Product* and *Domestic Industry;* if you disagree with either or both of these definitions, please explain why and provide alternative definitions.

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.61 of the Commission's rules.

By order of the Commission.

Issued: July 26, 2006.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E6–12275 Filed 7–31–06; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[USITC SE–06–048]

Government in the Sunshine Act Meeting Notice

AGENCY HOLDING THE MEETING: United States International Trade Commission.

TIME AND DATE: August 7, 2006 at 11 a.m.

PLACE: Room 101, 500 E Street SW., Washington, DC 20436, Telephone: (202) 205–2000.

STATUS: Open to the public.

MATTERS TO BE CONSIDERED: 1. Agenda for future meetings: none.

2. Minutes.

3. Ratification List.

4. Inv. No. 731–TA–1104 (Preliminary) (Certain Polyester Staple Fiber from China)—briefing and vote. (The Commission is currently scheduled to transmit its determination to the Secretary of Commerce on August 7, 2006; Commissioners' opinions are currently scheduled to be transmitted to the Secretary of Commerce on or before August 14, 2006.).

5. Outstanding action jackets: none.

In accordance with Commission policy, subject matter listed above, not disposed of at the scheduled meeting, may be carried over to the agenda of the following meeting.

By order of the Commission.

Issued: July 27, 2006.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. 06–6644 Filed 7–28–06; 1:12 pm]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Prior to issuing a registration under 21 U.S.C. 952(a)(2)(B), and in accordance with 21 CFR 1301.34(a), this is notice that on April 13, 2005, Kenco VPI, Division of Kenco Group Inc., 350 Corporate Place, Chattanooga, TN 37419, has made application to the Drug Enforcement Administration (DEA) by letter to be registered as an importer of Nabilone (7379), a basic class of controlled substance listed in Schedule II.

The company plans to import the listed controlled substance for distribution to its customers.

Kenco VPI has been an importer of Schedule III–V controlled substances since June 14, 2004. On April 14, 2005, the DEA added Schedule II to the firm's importer registration. The DEA also added the drug code for Nabilone, a Schedule II controlled substance, to the firm's registration on April 28, 2005. Both amendments to the registration were made without benefit of the required legal process for modifying the DEA registration. Kenco VPI is currently complying with the legal requirements to register as a Schedule III importer. In addition the firm was given authorization to import the Nabilone product into the United States on May 12, 2005. The Nabilone product was approved by the Food & Drug Administration on May 15, 2006. DEA has agreed to allow Kenco VPI to continue to import the Nabilone product into the United States, while the firm is completing the required legal process.

Any manufacturer who on April 13, 2005, was registered, or 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. For purposes of this Notice, DEA has chosen recognized applicable manufacturers registered on April 13, 2005, the date on which Kenco submitted its initial request to have Nabilone added to its DEA importer registration. By employing this date, DEA seeks to equitably address its initial failure to publish Kenco's request to import Nabilone, while at the same time allowing those entities that would have been in a position to request a hearing on April 13, 2005, had DEA filed a timely notice, the right to request a hearing.

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA **Federal Register** Representative/ODL; or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA **Federal Register** Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than August 31, 2006.

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 listed 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: July 26, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–12256 Filed 7–31–06; 8:45 am]

BILLING CODE 4410–09–P