market for the *Subject Merchandise* in each *Subject Country* since the *Order Dates*, 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 each *Subject Country*, 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: October 25, 2006.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E6–18311 Filed 10–31–06; 8:45 am]

BILLING CODE 7020–02–P

JOINT BOARD FOR THE ENROLLMENT OF ACTUARIES

Advisory Committee on Actuarial Examinations; Invitation for Membership on Advisory Committee

AGENCY: Joint Board for the Enrollment of Actuaries.

ACTION: Withdrawal of notice inviting membership on advisory committee; notice inviting membership on advisory committee.

SUMMARY: The Joint Board for the Enrollment of Actuaries (Joint Board), established under the Employee Retirement Income Security Act of 1974 (ERISA), is responsible for the enrollment of individuals who wish to perform actuarial services under ERISA. The Joint Board has established an Advisory Committee on Actuarial Examinations (Advisory Committee) to assist in its examination duties mandated by ERISA. The Joint Board published a **Federal Register** notice at 71 FR 30649, May 30, 2006, inviting membership on the Advisory Committee. That notice did not reflect the Joint Board's decision to extend the appointment term of current Advisory Committee members. Therefore, this document withdraws the previous notice and gives new notice inviting membership. In accordance with the Joint Board's decision, the appointment term of current Advisory Committee members will expire on February 28, 2007. This notice describes the Advisory Committee and invites applications from those interested in serving on it.

1. General

To qualify for enrollment to perform actuarial services under ERISA, an applicant must have requisite pension actuarial experience and satisfy knowledge requirements as provided in the Joint Board's regulations. The knowledge requirements may be satisfied by successful completion of Joint Board examinations in basic actuarial mathematics and methodology and in actuarial mathematics and methodology relating to pension plans qualifying under ERISA.

The Joint Board, the Society of Actuaries, and the American Society of Pension Professionals & Actuaries jointly offer examinations acceptable to the Joint Board for enrollment purposes and acceptable to those actuarial organizations as part of their respective examination programs.

2. Programs

The Advisory Committee plays an integral role in the examination program by assisting the Joint Board in offering examinations that will enable examination candidates to demonstrate the knowledge necessary to qualify for enrollment. The purpose of the Advisory Committee, as renewed, will remain that of assisting the Joint Board in fulfilling this responsibility. The Advisory Committee will discuss the philosophy of such examinations, will review topics appropriately covered in them, and will make recommendations relative thereto. It also will recommend to the Joint Board proposed examination questions. The Joint Board will maintain liaison with the Advisory Committee in this process to ensure that its views on examination content are understood.

3. Function

The manner in which the Advisory Committee functions in preparing examination questions is intertwined with the jointly administered examination program. Under that program, the participating actuarial organizations draft questions and submit them to the Advisory Committee for its consideration. After review of the draft questions, the Advisory Committee selects appropriate questions, modifies them as it deems desirable, and then prepares one or more drafts of actuarial examinations to be recommended to the Joint Board. (In addition to revisions of the draft questions, it may be necessary for the Advisory Committee to originate questions and include them in what is recommended.)

4. Membership

The Joint Board will take steps to ensure maximum practicable representation on the Advisory Committee of points of view regarding the Joint Board's actuarial examination extant in the community at large and from nominees provided by the actuarial organizations. Since the members of the actuarial organizations comprise a large segment of the actuarial profession, this appointive process ensures expression of a broad spectrum of viewpoints. All members of the Advisory Committee will be expected to act in the public interest, that is, to produce examinations that will help ensure a level of competence among those who will be accorded enrollment to perform actuarial services under ERISA.

Membership normally will be limited to actuaries previously enrolled by the Joint Board. However, individuals having academic or other special qualifications of particular value for the Advisory Committee's work also will be considered for membership. Membership terms are at the sole discretion of the inviting authority and are not necessarily concurrent with the duration of the Advisory Committee charter. The Advisory Committee will meet about four times a year. Advisory Committee members should be prepared to devote from 125 to 175 hours, including meeting time, to the work of the Advisory Committee over the course of a year. Members will be reimbursed for travel expenses incurred, in accordance with applicable government regulations.

Actuaries interested in serving on the Advisory Committee should express their interest and fully state their qualifications in a letter addressed to: Joint Board for the Enrollment of Actuaries, c/o Internal Revenue Service, Attn: Executive Director SE: OPR, Room 7238, 1111 Constitution Avenue, NW., Washington, DC 20224.

Any questions may be directed to the Joint Board's Executive Director at 202–622–8229.

The deadline for accepting applications is December 15, 2006.

Dated: October 11, 2006.

Patrick W. McDonough,

Executive Director, Joint Board for the Enrollment of Actuaries.

[FR Doc. 06–8992 Filed 10–31–06; 8:45 am]

BILLING CODE 4830–01–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 July 25, 2006, Alcan Packaging-Bethlehem, 2400 Baglyos Circle, Bethlehem, Pennsylvania, 18020, has made application to the Drug Enforcement Administration (DEA) 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 packaging and for distribution.

Any 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 written comments or objections being sent via regular mail should 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 December 1, 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: October 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–18431 Filed 10–31–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated July 25, 2006, and published in the **Federal Register** on July 31, 2006, (71 FR 43210), Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application by letter to the Drug Enforcement Administration (DEA) to be registered as an importer of Marihuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for packaging for a clinical trial study.

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 Aptuit 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 Aptuit 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: October 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–18376 Filed 10–31–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on August 28, 2006, 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 substance in bulk for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail should 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 January 2, 2007.

Dated: October 25, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–18375 Filed 10–31–06; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By notice dated July 26, 2006, and published in the **Federal Register** on August 1, 2006, (71 FR 43526), Kenco VPI, Division of Kenco Group Inc., 350