determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in section 210.50 of the Commission's Rules of Practice and Procedure (19 CFR 210.50).

The Commission instituted this investigation on November 22, 1996, based on a complaint filed by The Procter & Gamble Company (P&G) concerning allegations of unfair acts in violation of section 337 in the importation and sale of certain toothbrushes covered by U.S. Letters Patent Des. 328,392. The complaint, as amended, also alleged copyright infringement by certain respondents, but those allegations were subsequently withdrawn from the investigation.

The Commission found Shummi Enterprise Co., Ltd (Shummi) and Shumei Industrial Co., Ltd. (Shumei) in violation of section 337 and found Giftline International Corporation (Giftline) in default for failure to respond to the complaint and notice of investigation.

On July 2, 1997, the presiding administrative law judge (ALJ) issued a recommended determination (RD) on the issues of remedy and bonding for respondents Shummi and Shumei. The ALJ recommended a limited exclusion order and a bond in the amount of 100 percent of entered value during the 60-day Presidential review period.

On August 20, 1997, the Commission published a notice requesting written submissions on the issues of remedy, the public interest, and bonding. The Commission investigative attorney and complainant P&G filed submissions on these issues, essentially concurring with the ALJ's recommendations as to Shumei and Shummi and arguing for the same remedy and bond to apply to Giftline. No other submissions were filed.

Having reviewed the record in this investigation, including the parties' written submissions, the Commission determined that the appropriate form of relief is a limited exclusion order prohibiting the unlicensed entry of infringing toothbrushes that are manufactured abroad by or on behalf of Shummi, Shumei, or Giftline. The Commission further determined that the public interest factors enumerated in subsection 337(d) do not preclude issuance of the limited exclusion order, and that the bond during the Presidential review period shall be in the amount of one hundred (100) percent of the entered value of the articles in question.

Copies of the Commission's order, the public version of the Commission's opinion in support thereof, and all other nonconfidential documents filed in connection with this investigation are or will be available for inspection during official business hours (8:45 a.m. to 5:15 p.m.) in the Office of the Secretary, U.S. International Trade Commission, 500 E Street, S.W., Washington, D.C. 20436, telephone (202) 205–2000. Hearing impaired persons are advised that information on the matter can be obtained by contacting the Commission's TDD terminal at 202–205–1810.

Issued: October 15, 1997.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 97–27938 Filed 10–21–97; 8:45 am]

BILLING CODE 7020–02–P

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 June 6, 1997, Arenol Corporation, 189 Meister Avenue, Somerville, New Jersey 08876, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

Drug	Schedule
Methamphetamine (1105)	II
Phenylacetone (8501)	II

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

Any manufacturer holding, or applying for, registration as a bulk manufacturer of these basic classes of controlled substances 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than November 21, 1997.

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 basic classes of any controlled substances 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: October 7, 1997.

John H. King,

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

[FR Doc. 97–27894 Filed 10–21–97; 8:45 am]

BILLING CODE 4410–09–M

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 September 10, 1997, Norac Company, Inc., 405 S. Motor Avenue, Azusa, California 91702, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of tetrahydrocannabinols (7370) a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture medication for the treatment of AIDS wasting syndrome and as an antiemetic.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 22, 1997.

Dated: October 3, 1997.

John H. King,

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

[FR Doc. 97–27895 Filed 10–21–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacture 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 August 4, 1997, Novartis Pharmaceuticals Corp., Attn: Compliance, East Hanover, 556 Morris Avenue, Summit, New Jersey 07901, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the Schedule II controlled substance methylphenidate (1724).

The firm plans to manufacture the finished product 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 22, 1997.

Dated: October 6, 1997.

John H. King,

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

[FR Doc. 97–27896 Filed 10–21–97; 8:45 am]

BILLING CODE 4410–09–M

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 September 10, 1997, Nycomed, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, made application to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of meperidine (9230) a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture bulk product for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance 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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 22, 1997.

Dated: October 3, 1997.

John H. King,

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

[FR Doc. 97–27897 Filed 10–21–97; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Office of the Secretary

Advisory Council on Employee Welfare and Pension Benefit Plans; Reopening and Extending the Time for Receipt of Nominations for Vacancies Until November 7, 1997

Section 512 of the Employee Retirement Income Security Act of 1974 (ERISA), 88 Stat. 895, 29 U.S.C. 1142, provides for an ''Advisory Council on Employee Welfare and Pension Benefit Plans'' (the Council), to consist of 15 members to be appointed by the Secretary of Labor (the Secretary) as follows: Three representatives of employee organizations (at least one of whom shall be representative of an organization whose members are participants in a multiemployer plan); three representatives of employers (at least one of whom shall be representative of employers maintaining or contributing to multiemployer plans); one representative each from the fields of insurance, corporate trust, actuarial counseling, investment counseling, investment management and accounting; and three representatives from the general public (one of whom shall be a person representing those receiving benefits from a pension plan). No more than eight members of the Council shall be members of the same political party.

Members shall be qualified to appraise the programs instituted under ERISA. Appointments are for terms of three years. The prescribed duties of the Council are to advise the Secretary with respect to the carrying out of his or her functions under ERISA, and to submit to the Secretary, or his or her designee, recommendations with respect thereto. The Council will meet at least four times each year, and recommendations of the Council to the Secretary will be included in the Secretary's annual report to the Congress on ERISA.

The terms of five members of the Council expire Friday, November 14, 1997. The groups or fields represented are as follows: employee organizations (multiemployer plans), investment counseling, actuarial counseling, employers and the general public (pensioners). In addition, this year nominations also are being sought for individuals interested in an appointment to fill one year of an unexpired three-year term of a council member who died while serving on the Council. That unexpired term calls for naming an employee organization (multiemployer) representative.

Accordingly, notice is hereby given that any person or organization desiring to recommend one or more individuals for appointment to the Advisory Council on Employee Welfare and Pension Benefit Plans to represent any of the groups or fields specified in the preceding paragraph, may submit recommendations to Sharon Morrissey, Executive Secretary, ERISA Advisory Council, Frances Perkins Building, U.S. Department of Labor, 200 Constitution Avenue, NW, Suite N–5677, Washington, D.C. 20210. This notice is being issued to reopen and further extend the period in which recommendations can be delivered or mailed. The new date for receipt of recommendations is on or before November 7, 1997. Nominations for a particular category of membership should come from organizations or individuals within that category. A summary of the candidate's qualifications should be included with the nomination.

Signed at Washington, D.C., this 16 day of October, 1997.

Olena Berg,

Assistant Secretary of Labor, Pension and Welfare Benefits Administration.

[FR Doc. 97–27958 Filed 10–21–97; 8:45 am]

BILLING CODE 4510–29–M