sale for importation, or the sale within the United States after importation of certain toothbrushes and/or the packaging thereof, by reason of infringement of U.S. Copyright Registration No. TX 4–103–537; and

(c) Whether there exists an industry in the United States as required by subsection (a)(2) of section 337.

(2) For the purpose of the investigation so instituted, the following are hereby named as parties upon which this notice of investigation shall be served:

(a) The complainant is—The Procter & Gamble Company, One Procter & Gamble Plaza, Cincinnati, Ohio 45202.

(b) The respondents are the following companies alleged to be in violation of section 337, and are the parties upon which the complaint is to be served:

Shummi Enterprise Co., Ltd., No. 15, Alley 8, Lane 53, Nanking East Road, Section 4, Taipei, Taiwan.

Shumei Industrial Co., Ltd., Ping-Di, Central, Lung-Kang District, Shenzhen, China.

Giftline International Corporation, 1/F, No. 33, Alley 6, Lane 133, Nanking East Road, Section 4, Taipei, Taiwan.

Lollipop Imports & Exports of Brooklyn, Inc., 774 Broadway, Brooklyn, New York 11206.

MAS Marketing, Inc., 23800 Commerce Park D, Cleveland, Ohio 44122.

(c) Thomas S. Fusco, Esq., Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street, SW, Room 401–O, Washington, D.C. 20436, who shall be the Commission investigative attorney, party to this investigation; and

(3) For the investigation so instituted, the Honorable Sidney Harris is designated as the presiding Administrative Law Judge.

Responses to the complaint and the notice of investigation must be submitted by the named respondents in accordance with § 210.13 of the Commission's Rules of Practice and Procedure, 19 CFR 210.13. Pursuant to 19 CFR 201.16(d) and 210.13(a) such responses will be considered by the Commission if received no later than 20 days after the date of service of the complaint. Extensions of time for submitting responses to the complaint will not be granted unless good cause therefor is shown.

Failure of a respondent to file a timely response to each allegation in the complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the complaint and this notice, and to authorize the administrative law judge and the Commission, without further notice to the respondents, to find the facts to be as alleged in the complaint and this notice and to enter both an initial determination and a final determination containing such findings, and may result in the issuance of a limited exclusion order or a cease and desist order or both directed against such respondent.

Issued: November 22, 1996.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 96–30320 Filed 11–26–96; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.43(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on September 10, 1996, Hoffman-LaRoche, Inc., 340 Kingsland Street, Nutley, New Jersey 07110, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of levorphanol (9220) a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture finished dosage forms 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 above application.

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 January 27, 1997.

Dated: October 28, 1996.

Gene R. Haislip,

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

[FR Doc. 96–30353 Filed 11–26–96; 8:45 am]

BILLING CODE 4410–09–M

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 1311.42 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on September 18, 1996, North Pacific Trading Company, 1505 SE Gideon Street, Portland, Oregon 97202, made application by renewal to the Drug Enforcement Administration to be registered as an importer of marihauna (7360) a basic class of controlled substance listed in Schedule I.

This application is exclusively for the importation of marihuana seed which will be rendered non-viable and used as bird food.

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 1301.54 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections, or requests for a hearing may be addressed 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 (30 days from publication).

This procedure is to be conducted simultaneously with and independent of the procedures described in 21 CFR 1311.42 (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 1311.42 (a), (b), (c), (d), (e), and (f) are satisfied.