solutions for the problems affecting the Bay-Delta system has been chartered under the Federal Advisory Committee Act (FACA) as the Bay-Delta Advisory Council (BDAC) to advise CALFED on the program mission, problems to be addressed, and objectives for the CALFED Bay-Delta Program. BDAC provides a forum to help ensure public participation, and will review reports and other materials prepared by CALFED staff. BDAC has established a subcommittee called the Ecosystem Roundtable to provide input on annual workplans to implement ecosystem restoration projects and programs.

Minutes of the meeting will be maintained by the CALFED Bay-Delta Program, Suite 1155, 1416 Ninth Street, Sacramento, CA 95814, and will be available for public inspection during regular business hours, Monday through Friday within 30 days following the meeting.

Dated: November 19, 1996.

Roger Patterson,

Regional Director, Mid-Pacific Region.

[FR Doc. 96–30203 Filed 11–26–96; 8:45 am]

BILLING CODE 4310–94P–M

INTERNATIONAL TRADE COMMISSION

[Inv. No. 337–TA–387]

Certain Self-Powered Fiber Optic Modems; Notice of Commission Decision Not To Review an Initial Determination Terminating the Investigation on the Basis of a Settlement Agreement

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the Commission has determined not to review the presiding administrative law judge's (ALJ's) initial determination (ID) in the above-captioned investigation terminating the investigation on the basis of a settlement agreement.

FOR FURTHER INFORMATION CONTACT: Cynthia P. Johnson, Esq., Office of the General Counsel, U.S. International Trade Commission, telephone 202–205–3098.

SUPPLEMENTARY INFORMATION: This patent-based section 337 investigation was instituted by the Commission on April 25, 1996, on behalf of Patton Electronics Co. (Patton) of Gaithersburg, Maryland. The complaint alleges violations of section 337 based on the importation into the United States, the sale for importation, and the sale within the United States after importation of certain self-powered fiber optic modems that allegedly infringe claims 1, 2, 3, 7, and 8 of U.S. Letters Patent 4,161,650, (the '650 patent) and that there exists an industry in the United States as required by subsection (a)(2) of section 337. The notice of investigation named RAD Data Communications, Ltd., of Tel Aviv, Israel and RAD Data Communications, Inc. (collectively ''RAD'') of Mahwah, New Jersey as respondents.

On October 11, 1996, Patton and RAD filed a joint motion to terminate the investigation based on a settlement agreement. On October 23, 1996, the Commission investigative attorney (IA) filed a response in support of the joint motion to terminate the investigation. On October 24, 1996, the ALJ issued an ID (Order No. 16) granting the joint motion to terminate the investigation on the basis of a settlement agreement. No petitions for review of the ID were filed.

This action is taken under the authority of section 337 of the Tariff Act of 1930, 19 U.S.C. 1337, and Commission rule 210.42, 19 C.F.R. 210.42.

Copies of the ALJ's ID 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 on 202–205–1810.

Issued: November 21, 1996.

By order of the Commission.

Donna R. Koehnke,

Secretary.

[FR Doc. 96–30321 Filed 11–26–96; 8:45 am]

BILLING CODE 7020–02–P

[Investigation No. 337–TA–391]

Certain Toothbrushes and the Packaging Thereof; Notice of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Institution of investigation pursuant to 19 U.S.C. 1337.

SUMMARY: Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on October 25, 1996, under section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337, on behalf of Procter & Gamble Company, One Procter & Gamble Plaza, Cincinnati, OH 45202. An amended complaint was filed on November 14, 1996, and supplementary letters were filed on November 18 and 19, 1996. The complaint, as amended and supplemented, alleges violations of section 337 in the importation into the United States, the sale for importation, and the sale within the United States after importation of certain toothbrushes and packaging thereof by reason of infringement of U.S. Patent Des. 328,392 and U.S. Copyright Registration No. TX 4–103–537. The complaint further alleges that there exists an industry in the United States as required by subsection (a)(2) of section 337.

The complainant requests that the Commission institute an investigation and, after a hearing, issue permanent exclusion orders and permanent cease and desist orders.

ADDRESSES: The complaint, except for any confidential information contained therein, is 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, SW, Room 112, Washington, D.C. 20436, telephone 202–205–2000. Hearing-impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810.

FOR FURTHER INFORMATION CONTACT: Thomas S. Fusco, Esq., Office of Unfair Import Investigations, U.S. International Trade Commission, telephone 202–205–2571.

Authority

The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, and in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (1996).

Scope of Investigation

Having considered the complaint, the U.S. International Trade Commission, on November 21, 1996, *Ordered that*—

(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine—

(a) Whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the sale for importation, or the sale within the United States after importation of certain toothbrushes by reason of infringement of U.S. Patent Des. 328,392;

(b) Whether there is a violation of subsection (a)(1)(B) of section 337 in the importation into the United States, the

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.