(1) Pursuant to subsection (b) of section 337 of the Tariff Act of 1930, as amended, an investigation be instituted to determine 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 lighting products, components thereof, and products containing the same by reason of infringement of one or more of claims 23, 26, and 27 of U.S. Patent No. 6,082,878, and claims 1 and 7 of U.S. Patent No. 5,662,413, and whether an industry in the United States exists 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—Cooper Lighting, Inc., 1121 Highway 74 South, Peachtree City, GA 30269.

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

Cordelia Lighting, Inc., 20101 South Santa Fe Avenue, Rancho Dominguez, CA 90221.

Jimway, Inc., 20101 South Santa Fe Avenue, Rancho Dominguez, CA 90221.

(c) The Commission investigative attorney, party to this investigation, is Bryan F. Moore, Esq., Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street, SW., Room 401–I, Washington, DC 20436; and

(3) For the investigation so instituted, the Honorable Charles E. Bullock 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 section 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 not later than 20 days after the date of service by the Commission of the complaint and the notice of investigation. Extensions of time for submitting responses to the complaint and the notice of investigation 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 respondent, to find the facts to be as alleged in the complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of a limited exclusion order or cease and desist order or both directed against the respondent.

Issued: February 21, 2007.

By order of the Commission.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E7–3364 Filed 2–26–07; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–550]

In the Matter of Certain Modified Vaccinia Ankara (''MVA'') Viruses and Vaccines and Pharmaceutical Compositions Based Thereon; Notice of Commission Decision to Remand the Final Initial Determination Finding No Violation of Section 337 and To Extend the Target Date for Completion of the Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined to remand the final initial determination (''final ID'') of the presiding administrative law judge (''ALJ'') finding no violation of section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 (''section 337''), in the above-captioned investigation and to extend the target date for completion of the investigation to October 19, 2007.

FOR FURTHER INFORMATION CONTACT: James A. Worth, Office of the General Counsel, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436, telephone (202) 205–3065. Copies of non-confidential 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, SW., Washington, DC 20436, telephone (202) 205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*). The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at *http://edis.usitc.gov*. Hearing-impaired persons are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on (202) 205–1810.

SUPPLEMENTARY INFORMATION: This investigation was instituted on September 23, 2005, based on a complaint filed by Bavarian Nordic A/S of Denmark (''Bavarian Nordic''). The complaint alleged 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 Modified Vaccinia Ankara (''MVA'') viruses and vaccines and pharmaceutical compositions based thereon by reason of infringement of various claims of United States Patent Nos. 6,761,893 and 6,913,752. The complaint also alleged violations of section 337 in the importation of certain MVA viruses and vaccines and pharmaceutical compositions based thereon or in the sale of such articles by reason of misappropriation of trade secrets, the threat or effect of which is to destroy or substantially injure an industry in the United States. The complaint named a single respondent, Acambis PLC (''Acambis'') of the United Kingdom. Only the patent allegations remain in this investigation.

After a hearing and post-hearing briefing, the ALJ issued a final initial determination (''final ID'') on September 6, 2006, finding no violation of section 337. The ALJ held that the patents were infringed but invalid.

Bavarian Nordic, Acambis, and the Commission investigative attorney filed petitions for review of the final ID. By notice of November 22, 2006, the Commission determined to review the final ID in its entirety, as well as Order No. 10, and to ask the parties for briefing on the issues on review and on remedy, public interest and bonding. The parties submitted their initial and reply briefs on December 12 and December 22, 2006, respectively.

By notice of January 19, 2007, the Commission requested briefing on whether this investigation has become or will shortly become moot, and if so, whether the investigation should be terminated. The parties submitted briefing on January 26, 2007.

The Commission has determined to remand the final ID to the ALJ and to extend the target date for completion of the investigation by eight months to October 19, 2007.

This action is taken under the authority of section 337 of the Tariff Act

of 1930, as amended (19 U.S.C. 1337), and in sections 210.45(c), 210.51(a) of the Commission's Rules of Practice and Procedure (19 CFR 210.45(c), 210.51(a)).

By order of the Commission.

Issued: February 21, 2007.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E7–3390 Filed 2–26–07; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated October 6, 2006, and published in the **Federal Register** on October 18, 2006, (71 FR 61510), Boehringer Ingelheim Chemicals Inc., 2820 N. Normandy Drive, Petersburg, Virginia 23805, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in schedules I and II:

Drug	Schedule
Tetrahydrocannabinols (7370)	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Methadone (9250)	II
Methadone Intermediate (9254)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Fentanyl(9801)	II

The company plans to manufacture the listed controlled substances in bulk for sale to its customers for formulation into finished pharmaceuticals.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Boehringer Ingelheim Chemicals Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Boehringer Ingelheim Chemicals Inc. 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: February 16, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–3296 Filed 2–26–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated November 21, 2006 and published in the **Federal Register** on December 1, 2006, (71 FR 69591–69592), Johnson Matthey, Inc., Pharmaceutical Materials, 2003 Nolte Drive, West Deptford, New Jersey 08066–1742, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule II:

Drug	Schedule
Phenylacetone (8501)	II
Raw Opium (9600)	II
Concentrate of Poppy Straw (9670)	II

The company plans to import the listed controlled substances as raw materials for use in the manufacture of bulk controlled substances for distribution to its customers.

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 Johnson Matthey Inc to import the basic classes of controlled substances 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 Johnson Matthey Inc 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 classes of controlled substances listed.

Dated: February 16, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–3299 Filed 2–26–07; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated November 8, 2006 and published in the **Federal Register** on November 17, 2006, (71 FR 66974), Kenco VPI, Division of Kenco Group Inc., 350 Corporate Place, Chattanooga, Tennessee 37419, made application by renewal 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 distribution to its customers.

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 Kenco VPI 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 Kenco VPI 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: February 16, 2007.

Joseph T. Rannazzisi,

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

[FR Doc. E7–3298 Filed 2–26–07; 8:45 am]

BILLING CODE 4410–09–P