INTERNATIONAL TRADE COMMISSION

[Investigation No. 337–TA–952]

Certain Electronic Devices, Including Wireless Communication Devices, Computers, Tablet Computers, Digital Media Players, and Cameras; Commission Determination to Affirm an Initial Determination Granting a Joint Motion to Terminate the Investigation on the Basis of Settlement; Termination of Investigation

AGENCY: U.S. International Trade Commission.

ACTION: Notice.

SUMMARY: Notice is hereby given that the U.S. International Trade Commission has determined to affirm the administrative law judge's (ALJ) initial determination (ID) (Order No. 52) granting a joint motion to terminate the above-referenced investigation on the basis of a settlement agreement.

FOR FURTHER INFORMATION CONTACT: Ron Traud, Office of the General Counsel, U.S. International Trade Commission, 500 E Street SW., Washington, DC 20436, (202) 205–3427. 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, (202) 205–2000. General information concerning the Commission may also be obtained by accessing its Internet server at *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: On April 3, 2015, the Commission instituted this investigation (the 952 investigation) based on a complaint filed by Ericsson Inc. of Plano, Texas and Telefonaktiebolaget LM Ericsson of Sweden (collectively, ''Ericsson''). 80 FR 18254 (Apr. 3, 2015). The complaint alleged violations of 19 U.S.C. 1337 (Section 337) based upon the importation into the United States, the sale for importation, and the sale within the United States after importation of certain electronic devices, including wireless communication devices, computers, tablet computers, digital media players, and cameras by reason of infringement of certain claims of U.S. Patent Nos. 6,633,550; 6,157,620; 6,029,052; 8,812,059; 6,291,966; and 6,122,263. *Id.* at 18255. The Commission's Notice of Investigation named Apple Inc. of Cupertino, California (Apple) as respondent and also named the Office of Unfair Import Investigations (OUII) as a party. *Id.*

On December 29, 2015, Ericsson and Apple (collectively, the private parties) filed a joint motion to terminate the investigation pursuant to Commission Rule 210.21(b) on the basis of a settlement. *See* Order No. 51 at 1 (Jan. 12, 2016). On January 12, the ALJ (Judge Shaw) denied the motion because the private parties failed to provide a copy of the Agreement. *See id.* On February 1, 2016, the private parties filed a second amended joint motion (the Joint Motion) to terminate the investigation in view of a settlement agreement. *See* Order 52 at 1 (Mar. 9, 2016) [hereinafter, the Subject ID]. The motion included both a confidential, un-redacted and a public, redacted copy of the settlement agreement (the Agreement). *Id.* at 2. The Agreement and a corresponding motion to terminate were also filed in Investigation No. 337–TA–953 (the 953 investigation). *Id.*

On February 3, 2016, the ALJ presiding in the 953 investigation (Judge Lord) denied the motion to terminate that investigation, reasoning that the public version of the Agreement was over-redacted. *See id.* Pursuant to Commission Rules 210.24(b)(2)–(3) and 210.5(e), Ericsson filed a petition for interlocutory Commission review of only five of Judge Lord's confidentiality determinations. *See* Complainant Ericsson's Application for Commission Review of Certain Confidentiality Determinations in Order No. 45 (Feb. 11, 2016). Ericsson submitted with its appeal a revised, less-redacted public version of the Agreement (the Final Public Version). *Id.*

On March 9, 2016, Judge Shaw issued the Subject ID, which grants the Joint Motion. Subject ID, at 3. The Subject ID concludes that termination of the 952 investigation based on the private parties' settlement is in the public interest. *Id.* at 2. The Subject ID then declares that the private parties should file another public version of the Agreement in accordance with Judge Lord's ruling in the 953 investigation, as affirmed or modified by the Commission. *See id.* at 2–3. No petitions for review of the Subject ID were filed. On April 8, 2016, the Commission determined to review the Subject ID. Notice of Commission Determination to Review an Initial Determination Granting a Joint Motion to Terminate the Investigation on the Basis of Settlement, at 2 (Apr. 8, 2016).

On May 4, 2016, the Commission granted Ericsson's interlocutory appeal in the 953 investigation, reversed the ALJ on all five of the appealed confidentiality determinations, and remanded to the ALJ. Order Granting Appeal for Interlocutory Review of Order No. 45, Upon Review, Reversing, and Remanding to the Administrative Law Judge, at 3 (May 4, 2016).

On May 9, 2016, Ericsson filed with the Commission for purposes of the 952 investigation the Final Public Version. Letter to Secretary Lisa R. Barton enclosing Proposed Public Version of Parties' Global Patent License Agreement for Consideration in the Pending Initial Determination Terminating the Investigation Based on a Settlement Agreement (May 9, 2016).

The Commission hereby affirms the Subject ID, which grants the private parties' motion to terminate the investigation.

The authority for the Commission's determination is contained in Section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in Part 210 of the Commission's Rules of Practice and Procedure (19 CFR part 210).

By order of the Commission.
Issued: May 25, 2016.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2016–12711 Filed 5–27–16; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Rhodes Technologies

AGENCY: Drug Enforcement Administration, DOJ.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before August 1, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on March 26, 2016, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816 applied to be registered as a bulk manufacturer the following basic classes of controlled substances:

Controlled substance	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Methylphenidate (1724)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Morphine (9300)	II
Oripavine (9330)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to dosage form manufacturers.

In reference to drug code 7370 the company plans to bulk manufacture a synthetic tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

Dated: May 19, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–12752 Filed 5–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Wildlife Laboratories, Inc.

AGENCY: Drug Enforcement Administration, DOJ.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before June 30, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before June 30, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA **Federal Register** Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on February 19, 2016, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550–8055 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
Etorphine (except HCl) (9056)	I
Etorphine HCl (9059)	II

The company plans to import the listed controlled substances for sale to its customer.

Dated: May 19, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016–12751 Filed 5–27–16; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Office of the Secretary

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Trade Adjustment Assistance Program Reserve Funding Request

ACTION: Notice.

SUMMARY: The Department of Labor (DOL) is submitting the Employment and Training Administration (ETA) sponsored information collection request (ICR) titled, ''Trade Adjustment Assistance Program Reserve Funding Request,'' to the Office of Management and Budget (OMB) for review and approval for continued use, without change, in accordance with the Paperwork Reduction Act of 1995 (PRA), 44 U.S.C. 3501 *et seq.* Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that agency receives on or before June 30, 2016.

ADDRESSES: A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response, and estimated total burden may be obtained free of charge from the RegInfo.gov Web site at *http://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=201605-1205-009* (this link will only become active on the day following publication of this notice) or by contacting Michel Smyth by telephone at 202–693–4129, TTY 202–693–8064, (these are not toll-free numbers) or by email at *DOL_PRA_PUBLIC@dol.gov.*

Submit comments about this request by mail or courier to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for DOL–ETA, Office of Management and Budget, Room 10235, 725 17th Street NW.,