# INTERNATIONAL TRADE COMMISSION

[Investigation No. 337-TA-1071]

# Certain Wireless Audio Systems and Components Thereof Institution of Investigation

AGENCY: U.S. International Trade

Commission **ACTION:** Notice.

**SUMMARY:** Notice is hereby given that a complaint was filed with the U.S. International Trade Commission on August 10, 2017, under section 337 of the Tariff Act of 1930, as amended, on behalf of Broadcom Limited of San Jose, California and Avago Technologies General IP (Singapore) Pte. Ltd. of Singapore. An amended complaint was filed on August 16, 2017, and supplements to the amended complaint were filed on August 30, 2017. The amended complaint, as supplemented, alleges violations of 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 wireless audio systems and components thereof by reason of infringement of U.S. Patent No. 6,684,060 ("the '060 patent"). The amended complaint further alleges that an industry in the United States exists as required by the applicable Federal Statute.

The complainants request that the Commission institute an investigation and, after the investigation, issue a limited exclusion order and cease and desist orders.

ADDRESSES: The amended complaint, as supplemented, 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, DC 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. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at (202) 205-2000. General information concerning the Commission may also be obtained by accessing its internet server at https://www.usitc.gov. The public record for this investigation may be viewed on the Commission's electronic docket (EDIS) at https://edis.usitc.gov.

**FOR FURTHER INFORMATION CONTACT:** Pathenia M. Proctor, The Office of

Unfair Import Investigations, U.S. International Trade Commission, telephone (202) 205–2560.

# SUPPLEMENTARY INFORMATION:

Authority: The authority for institution of this investigation is contained in section 337 of the Tariff Act of 1930, as amended, 19 U.S.C. 1337 and in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (2017).

Scope of Investigation: Having considered the amended complaint, the U.S. International Trade Commission, on September 8, 2017, ordered that—

- (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 wireless audio systems and components thereof by reason of infringement of claim 20 of the '060 patent; 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 complainants are: Broadcom Limited, 1320 Ridder Park Drive, San Jose, CA 95131 Avago Technologies General IP (Singapore) Pte. Ltd., 1 Yinshun Avenue 7, Singapore, 768923
- (b) The respondents are the following entities alleged to be in violation of section 337, and are the parties upon which the amended complaint is to be served:

DTS, Inc., 5220 Las Virgenes Road, Calabasas, CA 91302

Phorus, Inc., 5220 Las Virgenes Road, Calabasas, CA 91302

MartinLogan, Ltd., 2101 Delaware Street, Lawrence, KS 66046–3149 Paradigm Electronics Inc., 205 Annagem Boulevard, Mississauga, ON L5T 2V1, Canada

Anthem Electronics, Inc., 205 Annagem Boulevard, Mississauga, ON L5T 2V1, Canada

Wren Sound Systems, LLC, 169 Gateshead Way, Phoenixville, PA 19460

McIntosh Laboratory, Inc., 2 Chambers Street, Binghamton, NY 13903–2699 Definitive Technology, 11433 Cronridge Drive, Suite K, Owings Mills, MD 21117

Polk Audio Inc., 1 Viper Way, Vista, CA 92081

(c) The Office of Unfair Import Investigations, U.S. International Trade Commission, 500 E Street SW., Suite 401, Washington, DC 20436; and

(3) For the investigation so instituted, the Chief Administrative Law Judge, U.S. International Trade Commission, shall designate the presiding Administrative Law Judge.

Responses to the amended 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(e) 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 amended complaint and the notice of investigation. Extensions of time for submitting responses to the amended 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 amended complaint and in this notice may be deemed to constitute a waiver of the right to appear and contest the allegations of the amended 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 amended complaint and this notice and to enter an initial determination and a final determination containing such findings, and may result in the issuance of an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission. Issued: September 11, 2017.

#### Lisa R. Barton,

 $Secretary\ to\ the\ Commission.$  [FR Doc. 2017–19597 Filed 9–14–17; 8:45 am]

BILLING CODE 7020-02-P

### **DEPARTMENT OF JUSTICE**

# **Drug Enforcement Administration**

[Docket No. DEA-392]

Importer of Controlled Substances Application: Cerilliant Corporation

**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 on or before October 16, 2017. Such persons may also file a written request

for a hearing on the application on or before October 16, 2017.

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

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his 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 Assistant Administrator of the DEA Diversion Control Division ("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 July 13, 2017, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
3-Fluoro-N-methylcathinone (3–FMC)	1233	I
Cathinone	1235	1
Methcathinone	1237	1
4-Fluoro-N-methylcathinone (4–FMC)	1238	1
Pentedrone (α-methylaminovalerophenone)	1246	1
Mephedrone (4-Methyl-N-methylcathinone)	1248	1
4-Methyl-N-ethylcathinone (4–MEC)	1249	1
Naphyrone	1258	!
N-Ethylamphetamine	1475	!
N,N-limethylamphetamine	1480	!
Fenethylline	1503	!
Methaqualone	2565	!
JWH–250 (1-Pentyl-3-(2-methoxyphenylacetyl) indole)	6250	!
SR–18 (Also known as RCS–8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl) indole)	7008	!
5-Flouro-UR-144 and XLR11 [1-(5-Fluoro-pentyl)1H-indol-3-yl](2,2,3,3-tetramethylcyclopropyl)methanone	7011	!
	7012	
JWH-019 (1-Hexyl-3-(1-naphthoyl)indole)	7019	
AB-PINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-pentyl-1H-indazole-3-carboxamide)	7023 7024	
AB-CHMINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl)-1-(cyclohexylmethyl)-1H-indazole-3-carboxamide	7024 7031	1
ADB-PINACA (N-(1-amino-3,1-idinyi-1-oxobutan-2-yl)-1-pentyl-1H-indazole-3-carboxamide)	7035	1
APINACA (N-(1-amino-3,3-dimetryi-1-oxobutar-z-yi)-1-perityi-1H-indazole-3-carboxamide)	7035	1
JWH–081 (1-Pentyl-3-(1-(4-methoxynaphthoyl) indole)	7048	1
SR=19 (Also known as RCS=4) (1-Pentyl-3-[(4-methoxy)-benzoyl] indole	7104	
JWH–018 (also known as AM678) (1-Pentyl-3-(1-naphthoyl)indole)	7118	li
JWH-122 (1-Pentyl-3-(4-methyl-1-naphthoyl) indole)	7110	li
UR-144 (1-Pentyl-1H-indol-3-yl)(2,2,3,3-tetramethylcyclopropyl)methanone	7144	li
JWH-073 (1-Butyl-3-(1-naphthoyl)indole)	7173	l i
JWH–200 (1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole)	7200	li
AM2201 (1-(5-Fluoropentyl)-3-(1-naphthoyl) indole)	7201	li
JWH-203 (1-Pentyl-3-(2-chlorophenylacetyl) indole)	7203	l i
PB-22 (Quinolin-8-yl 1-pentyl-1H-indole-3-carboxylate)	7222	l i
5F-PB-22 (Quinolin-8-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate)	7225	l i
Alpha-ethyltryptamine	7249	li
bogaine	7260	1
CP-47,497 (5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl-phenol)	7297	1
CP-47,497 C8 Homologue (5-(1,1-Dimethyloctyl)-2-[(1F,3S)3-hydroxycyclohexyl-phenol)	7298	1
	7315	1
2,5-Dimethoxy-4-(n)-propylthiophenethylamine (2C-T-7)	7348	1
Marihuana	7360	1
Parahexyl	7374	1
Mescaline	7381	1
2-(4-Ethylthio-2,5-dimethoxyphenyl) ethanamine (2C-T-2)	7385	1
3,4,5-Trimethoxyamphetamine	7390	1
4-Bromo-2,5-dimethoxyamphetamine	7391	1
4-Bromo-2,5-dimethoxyphenethylamine	7392	1
4-Methyl-2,5-dimethoxyamphetamine	7395	I
2,5-Dimethoxyamphetamine	7396	1
JWH-398 (1-Pentyl-3-(4-chloro-1-naphthoyl) indole)	7398	I
3,4-Methylenedioxyamphetamine	7400	1
5-Methoxy-3,4-methylenedioxyamphetamine	7401	I
N-Hydroxy-3,4-methylenedioxyamphetamine	7402	1
3,4-Methylenedioxy-N-ethylamphetamine	7404	ļ ļ
3,4-Methylenedioxymethamphetamine	7405	
4-Methoxyamphetamine	7411	H

Controlled substance	Drug code	Schedule
5-Methoxy-N-N-dimethyltryptamine	7431	1
Alpha-methyltryptamine	7432	1
Bufotenine	7433 7434	1
Dimethyltryptamine	7435	i
Psilocybin		i
Psilocýn	7438	1
5-Methoxy-N,N-diisopropyltryptamine		1
N-Ethyl-1-phenylcyclohexylamine	7455	ļ
1-(1-Phenylcyclohexyl)pyrrolidine		1
N-Benzylpiperazine	7470 7493	i
4-Methyl-alphapyrrolidinopropiophenone (4-MePPP)	7498	i
2-(2,5-Dimethoxy-4-methylphenyl) ethanamine (2C-D)		i
2-(2,5-Dimethoxy-4-ethylphenyl) ethanamine (2C-E)	7509	1
2-(2,5-Dimethoxyphenyl) ethanamine (2C-H)		1
2-(4-lodo-2,5-dimethoxyphenyl) ethanamine (2C–l)	7518	!
2-(4-Chloro-2,5-dimethoxyphenyl) ethanamine (2C–C)	7519 7521	1
2-(2,5-Dimethoxy-4-Intro-priefryf) ethanamine (2C-P)		i
2-(4-Isopropylthio)-2,5-dimethoxyphenyl) ethanamine (2C–T–4)	7532	i
MDPV (3,4-Methylenedioxypyrovalerone)	7535	İ
2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine (25B–NBOMe)	7536	1
2-(4-chloro-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine (25C-NBOMe)	7537	İ
2-(4-iodo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) éthanamine (25I–NBOMe)	7538	Į.
Methylone (3,4-Methylenedioxy-N-methylcathinone)	7540	1
Butylone	7541	!
Pentylonealpha-pyrrolidinopentiophenone (α-PVP)	7542 7545	i
alpha-pyrrolidinobutiophenone (α-PBP)	7546	i
AM–694 (1-(5-Fluoropentyl)-3-(2-iodobenzoyl) indole)	7694	i
Desomorphine	9055	I
Etorphine (except HCI)	9056	1
Codeine methylbromide	9070	1
Heroin	9200	!
Morphine-N-oxide	9307 9313	ļ
NormorphinePholcodine	9314	i
U–47700 (3,4-dichloro-N-[2-dimethylamino)cyclohexy]-N-methylbenzamide)		i
AH–7921 (3,4-dichloro-N-[(1-dimethylamino)cyclohexylmethyl]benzamide))	9551	i
Acetylmethadol	9601	1
Allylprodine	9602	I
Alphacetylmethadol except levo-alphacetylmethadol	9603	!
Alphameprodine	9604	1
Alphamethadol	9605 9607	1
Betameprodine	9608	i
Betamethadol	9609	i
Betaprodine	9611	i
Dextromoramide	9613	1
Dipipanone	9622	1
Hydroxypethidine	9627	1
Noracymethadol	9633	1
Norlevorphanol	9634 9635	I I
Racemoramide	9645	í
Trimeperidine	9646	i
1-Methyl-4-phenyl-4-propionoxypiperidine	9661	I
Tilidine	9750	1
Para-Fluorofentanyl	9812	!
3-Methylfentanyl	9813	I
Alpha-methylfentanyl	9814	I
Acetyl-alpha-methylfentanylBeta-hydroxyfentanyl	9815 9830	i
Beta-hydroxy-3-methylfentanyl	9831	i
Alpha-methylthiofentanyl	9832	i
3-Methylthiofentanyl	9833	i
Thiofentanyl	9835	1
Methamphetamine	1105	II
Methylphenidate	1724	II
	2125	II
Amobarbital		
	2270 2315	II II

Controlled substance	Drug code	Schedule
Nabilone	7379	П
1-Phenylcyclohexylamine	7460	П
Phencyclidine	7471	П
Phenylacetone	8501	П
1-Piperidinocyclohexanecarbonitrile	8603	П
Alphaprodine	9010	П
Dihydrocodeine	9120	П
Ecgonine	9180	П
Ethylmorphine	9190	П
Levomethorphan	9210	П
Levorphanol	9220	ii
Meperidine	9230	ii
Dextropropoxyphene, bulk (non-dosage forms)	9273	ii
Levo-alphacetylmethadol	9648	ii
Noroxymorphone	9668	ii
Racemethorphan	9732	ii
Alfentanil	9737	ii
Remifentanil	9739	lii
Sufentanil	9740	ii
Carfentanil	9743	lii
Tapentadol	9780	l ii

The company plans to import small quantities of the listed controlled substances for the manufacture of analytical reference standards and distribution to their research and forensic customers.

In reference to drug code 7360 the company plans to import a synthetic cannabidiol. No other activity for this drug code is authorized for this registration.

Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of FDA approval or non-approved finished dosage forms for commercial sale.

Dated: August 31, 2017.

# Demetra Ashley,

Acting Assistant Administrator. [FR Doc. 2017–19067 Filed 9–14–17; 8:45 am]

BILLING CODE 4410-09-P

#### **DEPARTMENT OF JUSTICE**

### **National Institute of Justice**

[OMB Number 1121-NEW]

Agency Information Collection Activities: Proposed New Information Collection Activity; Comment Request, Proposed Study Entitled "Evaluation of the Bureau of Justice Assistance Sexual Assault Kit Initiative"

**AGENCY:** National Institute of Justice, U.S. Department of Justice.

**ACTION:** 30-Day notice.

**SUMMARY:** The Department of Justice (DOJ), Office of Justice Programs, National Institute of Justice, will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

**DATES:** Comments are encouraged and will be accepted for 60 days until November 14, 2017.

# FOR FURTHER INFORMATION CONTACT:

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to Christine Crossland, National Institute of Justice, Office of Research & Evaluation, 810 Seventh Street NW., Washington, DC 20531 (overnight 20001) or via email at christine.crossland@ojp.usdoj.gov.

Written comments and/or suggestions can also be sent to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503 or sent to OIRA\_submissions@omb.eop.gov.

SUPPLEMENTARY INFORMATION: This process is conducted in accordance with 5 CFR 1320.10. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

—Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the National Institute of Justice, including whether the information will have practical utility;
—Evaluate the accuracy of the agency's

estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
—Evaluate whether, and if so how, the

 Evaluate whether, and if so how, the quality, utility, and clarity of the information to be collected can be enhanced; and

—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

# Overview of This Information Collection

1. *Type of Information Collection:* Site visits, which will include individual and group interviews.

2. The Title of the Form/Collection: Evaluation of the Bureau of Justice Assistance Sexual Assault Kit Initiative.

3. The agency form number, if any, and the applicable component of the Department sponsoring the collection: The applicable component within the U.S. Department of Justice is the National Institute of Justice in the Office of Justice Programs.

4. Affected public who will be asked or required to respond, as well as a brief abstract: Sexual assault kits (SAKs) are invaluable tools used in investigations to collect evidence such as DNA and to document injuries from alleged victims; this evidence in turn is used to identify and prosecute offenders and to exonerate innocent suspects. Despite the importance of SAKs, backlogs of