in section 210.10 of the Commission's Rules of Practice and Procedure, 19 CFR 210.10 (2014).

Scope of Investigation: Having considered the complaint, the U.S. International Trade Commission, on March 30, 2015, *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 standard compliant electronic devices, including communication devices and tablet computers, by reason of infringement of one or more of claims 1, 2, 4, 6, 7, 9, 11–13, 15–19, 21, and 22 of the '996 patent; claims 7–12 and 18–23 of the '270 patent; claims 28–54 of the '359 patent; claims 1, 8–10, 12, 23, 24, 26, 27, 29–31, 38–40, 42, 49, 50, 52, 53, 57, 58, 64–66, and 68 of the '556 patent; claims 19, 20, 22–27, and 29–32 of the '805 patent; claims 1, 3, 5–8, 10, and 12–15 of the '130 patent; claims 1, 2, 4, 5, 7, 11, and 13–15 of the '381 patent; and claims 1, 3, 4, 6, 8, 9, 11, 12, 14, 16, 25, and 26 of the '476 patent; and whether an industry in the United States exists as required by subsection (a)(2) of section 337;

(2) Pursuant to Commission Rule 210.50(b)(1), 19 CFR 210.50(b)(1), the presiding administrative law judge shall take evidence or other information and hear arguments from the parties and other interested persons with respect to the public interest in this investigation, as appropriate, and provide the Commission with findings of fact and a recommended determination on this issue, which shall be limited to the statutory public interest factors set forth in 19 U.S.C. 1337(d)(1), (f)(1), (g)(1).

(3) 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: Ericsson Inc., 6300 Legacy Drive, Plano, TX 75024.

Telefonaktiebolaget LM Ericsson, Torshamnsgatan 21, Kista, Stockholm, Sweden.

(b) The respondent is the following entity alleged to be in violation of section 337, and is the party upon which the complaint is to be served: Apple Inc., a/k/a Apple Computer, Inc., 1 Infinite Loop, Cupertino, CA 95014.

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

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

A response to the complaint and the notice of investigation must be submitted by the named respondent 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 a response 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 a response to the complaint and the notice of investigation will not be granted unless good cause therefor is shown.

Failure of the 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 an exclusion order or a cease and desist order or both directed against the respondent.

By order of the Commission.

Issued: March 30, 2015.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2015–07647 Filed 4–2–15; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0024]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection; Reports of Regulated Transactions Involving Extraordinary Quantities, Uncommon Methods of Payment, and Unusual/Excessive Loss or Disappearance, and Regulated Transactions in Tableting/Encapsulating Machines

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 30-day notice.

SUMMARY: The Department of Justice (DOJ), Drug Enforcement Administration (DEA), 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. This proposed information collection was previously published in the **Federal Register** at 80 FR 6766, February 06, 2015, allowing for a 60 day comment period.

DATES: Comments are encouraged and will be accepted for an additional 30 days until May 4, 2015.

SUPPLEMENTARY INFORMATION: 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 agency, 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 quality, utility, and clarity of the information proposed 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 forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.

2. *Title of the Form/Collection:* Reports of Regulated Transactions Involving Extraordinary Quantities, Uncommon Methods of Payment, and Unusual/Excessive Loss or Disappearance, and Regulated Transactions in Tableting/Encapsulating Machines.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Notification of extraordinary quantities, uncommon methods of payment, and unusual/excessive loss or disappearance of listed chemicals and regulated transactions in tableting/encapsulating

machines is provided in writing on an as needed basis and does not require use of a form. The applicable component within the Department of Justice is the Drug Enforcement Administration, Office of Diversion Control.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Affected public (Primary): Business or other for-profit.

Affected public (Other): Not-for-profit institutions; Federal, State, local, and tribal governments.

Abstract: Each regulated person is required to report any regulated transaction involving an extraordinary quantity of a listed chemical, an uncommon method of payment or delivery, any unusual or excessive loss or disappearance of a listed chemical, and any regulated transaction in a tableting or encapsulating machine, to include any domestic regulated transaction in a tableting or encapsulating machine and any import or export of a tableting or encapsulating machine. 21 U.S.C. 830 (b)(1)(A), (C) and (D); 21 CFR 1310.05(a)(1), (3) and (4); 21 CFR 1310.05(c).

Regulated persons include manufacturers, distributors, importers, and exporters of listed chemicals, tableting machines, or encapsulating machines, or persons who serve as brokers or traders for international transactions involving a listed chemical, tableting machine, or encapsulating machine. 21 CFR 1300.02(b).

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* The DEA estimates that 63 persons respond as needed to this collection. Responses take 20 minutes.

6. *An estimate of the total public burden (in hours) associated with the proposed collection:* The DEA estimates that this collection takes 21 annual burden hours.

If additional information is required please contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., Suite 3E.405B, Washington, DC 20530.

Jerri Murray,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2015–07666 Filed 4–2–15; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

[OMB Number 1117–0012]

Agency Information Collection Activities; Proposed eCollection, eComments Requested; Extension Without Change of a Previously Approved Collection Application for Registration, Application for Registration Renewal, Affidavit for Chain Renewal (DEA Forms 225, 225a and 225b)

AGENCY: Drug Enforcement Administration, Department of Justice.

ACTION: 30-day notice.

SUMMARY: The Department of Justice (DOJ), Drug Enforcement Administration (DEA), 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. This proposed information collection was previously published in the **Federal Register** at 80 FR, page 5137, on January 30, 2015, allowing for a 60 day comment period.

DATES: Comments are encouraged and will be accepted for an additional 30 days until May 4, 2015.

SUPPLEMENTARY INFORMATION: 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 agency, 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 quality, utility, and clarity of the information proposed 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 forms of information technology, *e.g.,* permitting electronic submission of responses.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a currently approved collection.

2. *Title of the Form/Collection:* Application for Registration, Application for Registration Renewal, Affidavit for Chain Renewal.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* The form numbers are DEA Forms 225, 225a, and 225b. The applicable component within the Department of Justice is the Drug Enforcement Administration, Office of Diversion Control.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:*

Affected public (Primary): Business or other for-profit.

A*ffected public (Other):* Not-for-profit institutions, Federal, State, local, and tribal governments.

Abstract: The Controlled Substances Act requires all businesses and individuals who manufacture, distribute, import, export, and conduct research and laboratory analysis with controlled substances to register with the DEA. 21 U.S.C. 822, 21 CFR 1301.11 and 1301.13. Registration is a necessary control measure that prevents diversion by ensuring the closed system of distribution of controlled substances can be monitored by the DEA and that the businesses and individuals handling controlled substances are qualified to do so and are accountable.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* DEA Form 225 is only for registration of controlled substance manufacturers, distributors, reverse distributors, importers, exporters, researchers, canine handlers, and analytical laboratories, as well as list 1 chemical manufacturers and importers. DEA Form 225 is submitted on an as-needed basis by persons seeking to become registered. DEA Form 225a is submitted annually thereafter to renew existing registrations. DEA Form 225b is submitted annually for renewals of chain registrants. Chain registrants are those corporations and laboratories that maintain separate registrations at multiple locations (*e.g.,* distributors) and may renew all their registrations using a single DEA Form 225b.