the patentee and the applicable standard-setting organization, *i.e.,* the International Telecommunication Union (ITU)?

2. Please summarize the history to date of negotiations between LSI and Funai and between LSI and Realtek concerning any potential license to the '663, the '958, and the '867 patents, either alone, in conjunction with each other and/or the '087 patent, and/or in conjunction with non-asserted patents. Please provide copies of, or cite to their location in the record evidence, all offers and communications related to the negotiations including any offer or counteroffer made by Funai and Realtek.

3. Please summarize all licenses to the '663, the '958, and the '867 patents granted by LSI to any entity including evidence of the value of each patent if such patent was licensed as part of a patent portfolio. Please provide copies of, or cite to their location in the record evidence, all agreements wherein LSI grants any entity a license to these patents. Please also provide a comparison of the offers made to Funai and/or Realtek with offers made to these other entities.

4. If applicable, please discuss the industry practice for licensing patents involving technology similar to the technology in the '663, the '958, and the '867 patents individually or as part of a patent portfolio.

5. Please identify the forums in which you have sought and/or obtained a determination of a RAND rate for the '663, the '958, and the '867 patents. LSI, Funai and Realtek are each requested to submit specific licensing terms for the '663, the '958, and the '867 patents that each believes are reasonable and non-discriminatory.

6. Please discuss and cite any record evidence of any party attempting to gain undue leverage, or constructively refusing to negotiate a license, with respect to the '663, the '958, and the '867 patents. Please specify how that evidence is relevant to whether section 337 remedies with respect to such patents would be detrimental to competitive conditions in the U.S. economy and any other statutory public interest factor.

If the Commission orders some form of remedy, the U.S. Trade Representative, as delegated by the President, has 60 days to approve or disapprove the Commission's action. See Presidential Memorandum of July 21, 2005, 70 FR 43251 (July 26, 2005). During this period, the subject articles would be entitled to enter the United States under bond, in an amount determined by the Commission. The Commission is therefore interested in receiving submissions concerning the amount of the bond that should be imposed if a remedy is ordered.

Written Submissions: The parties to the investigation are requested to file written submissions on the issues identified in this notice. Parties to the investigation, interested government agencies, and any other interested parties are encouraged to file written submissions on the issues of remedy, the public interest, and bonding. Such submissions should address the recommended determination by the ALJ on remedy and bonding with respect to the asserted patents. Complainant is also requested to submit proposed remedial orders for the Commission's consideration. Complainant is further requested to state the date that the patents expire and the HTSUS numbers under which the accused products are imported. The written submissions and proposed remedial orders must be filed no later than close of business on Friday, November 1, 2013. Initial submissions by the parties are limited to 100 pages, not including submissions related to remedy, bonding, and the public interest. Reply submissions must be filed no later than the close of business on Monday, November 11, 2013. All reply submissions are limited to 60 pages, not including submissions related to remedy, bonding, and the public interest. No further submissions on these issues will be permitted unless otherwise ordered by the Commission.

Persons filing written submissions must file the original document electronically on or before the deadlines stated above and submit 8 true paper copies to the Office of the Secretary by noon the next day pursuant to section 210.4(f) of the Commission's Rules of Practice and Procedure (19 CFR 210.4(f)). Submissions should refer to the investigation number (''Inv. No. 337–TA–837'') in a prominent place on the cover page and/or the first page. (See Handbook for Electronic Filing Procedures, *http://www.usitc.gov/secretary/fed_reg_notices/rules/handbook_on_electronic_filing.pdf).* Persons with questions regarding filing should contact the Secretary (202–205–2000).

Any person desiring to submit a document to the Commission in confidence must request confidential treatment. All such requests should be directed to the Secretary to the Commission and must include a full statement of the reasons why the Commission should grant such treatment. See 19 CFR 201.6. Documents for which confidential treatment by the Commission is properly sought will be treated accordingly. A redacted non-confidential version of the document must also be filed simultaneously with the any confidential filing. All non-confidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.

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 sections 210.42–46 and 210.50 of the Commission's Rules of Practice and Procedure (19 CFR 210.42–46 and 210.50).

Issued: October 17, 2013.

By order of the Commission.

Lisa R. Barton,

Secretary to the Commission.

[FR Doc. 2013–24752 Filed 10–22–13; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Office on Violence Against Women

[OMB Number 1122–0001]

Certification of Compliance With the Statutory Eligibility Requirements of the Violence Against Women Act as Amended for Applicants to the STOP (Services* Training* Officers* Prosecutors) Violence Against Women Formula Grant Program; Agency Information Collection Activities: Revision of a Currently Approved Collection

ACTION: 30-Day Notice.

The Department of Justice, Office on Violence Against Women (OVW) will be submitting 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. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** Volume 78, page 39325 on July 1, 2013, allowing for a 60 day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until November 22, 2013. This process is conducted in accordance with 5 CFR 1320.10.

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 The Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk

Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395–7285.

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:

(1) 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;

(2) 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;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) 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:* Revision of a currently approved collection

(2) *Title of the Form/Collection:* ''Certification of Compliance with the Statutory Eligibility Requirements of the Violence Against Women Act as Amended'' for Applicants to the STOP Formula Grant Program

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: 1122–0001. U.S. Department of Justice, Office on Violence Against Women

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: The affected public includes STOP formula grantees (50 states, the District of Columbia and five territories (Guam, Puerto Rico, American Samoa, Virgin Islands, Northern Mariana Islands). The STOP Violence Against Women Formula Grant Program was authorized through the Violence Against Women Act of 1994 and reauthorized and amended by the Violence Against Women Act of 2000, the Violence Against Women Act of 2005 and the Violence Against Women Act of 2013. The purpose of the STOP Formula Grant Program is to promote a coordinated, multi-disciplinary approach to improving the criminal justice system's response to violence against women. It envisions a partnership among law enforcement, prosecution, courts, and victim advocacy organizations to enhance victim safety and hold offenders accountable for their crimes of violence against women. The Department of Justice's Office on Violence Against Women (OVW) administers the STOP Formula Grant Program funds which must be distributed by STOP state administrators according to statutory formula (as amended by VAWA 2000, VAWA 2005 and VAWA 2013).

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond/reply:* It is estimated that it will take the approximately 56 respondents (state administrators from the STOP Formula Grant Program) less than one hour to complete a Certification of Compliance with the Statutory Eligibility Requirements of the Violence Against Women Act, as Amended.

(6) *An estimate of the total public burden (in hours) associated with the collection:* The total annual hour burden to complete the Certification is less than 56 hours.

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, 145 N Street NE., Washington, DC 20530.

Dated: October 17, 2013.

Jerri Murray,

Department Clearance Officer for PRA, United States Department of Justice.

[FR Doc. 2013–24733 Filed 10–22–13; 8:45 am]

BILLING CODE 4410–FX–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0013]

Agency Information Collection Activities; Proposed Collection; Comments Requested: Application for Permit To Import Controlled Substances for Domestic and/or Scientific Purposes Pursuant to 21 U.S.C. 952 (DEA Form 357)

ACTION: 30-Day Notice.

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. The proposed information collection is published to obtain comments from the public and affected agencies. This proposed information collection was previously published in the **Federal Register** Volume 78, Number 154, page 48718 on August 9, 2013, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until November 22, 2013. This process is conducted in accordance with 5 CFR 1320.10.

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 the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. The best way to ensure your comments are received is to email them to *oira_submission@omb.eop.gov* or fax them to (202) 395–7285. 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 agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- Enhance the quality, utility, and clarity of the information to be collected; 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 Information Collection 1117–0013

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

(2) *Title of the Form/Collection:* Application for Permit to Import Controlled Substances for Domestic and/or Scientific Purposes pursuant to 21 U.S.C. 952 (DEA Form 357).

(3) *Agency form number, if any, and the applicable component of the Department sponsoring the collection:*