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**Background:** In his letter the USTR requested, under the authority of section 332(g) of the Tariff Act of 1930, that the Commission provide three reports during the next 12 months relating to small and medium-sized enterprises (SMEs). In this notice the Commission is instituting the third of three investigations under section 332(g) for the purpose of preparing the third report, which is to be transmitted to the USTR by October 6, 2010. The Commission published notices of institution of the first investigation, investigation No. 332-508, in the **Federal Register** of October 28, 2009 (74 FR 55581) and the second investigation, investigation No. 332-509, in the **Federal Register** of December 1, 2009 (74 FR 62812).

As requested, in the third report the Commission will, to the extent possible:

1. Examine U.S. SMEs engaged in providing services, including the characteristics of firms that produce tradable services, the growth in these services exports, and the differences between SME and large services exporters;
2. Identify how data gaps might be overcome to further enhance our understanding of SMEs in services sector exports;
3. For both goods and services exports, identify trade barriers (nontariff barriers and tariffs) that may disproportionately affect SME export performance, as well as possible linkages between exporting and SME performance; and
4. Provide insights on the degree to which SMEs operate as multinationals, as affiliate firms, or as contributors of indirect exports to international trade through sales to larger exporting firms.

The USTR requested that the Commission deliver the second report by October 6, 2010.

**Public Hearing:** The Commission will hold a joint public hearing in connection with this investigation and investigation No. 332-509 at the U.S.

International Trade Commission Building, 500 E Street, SW., Washington, DC, beginning at 9:30 a.m. on Tuesday, February 9, 2010 (and continuing on February 10, 2010, if needed). Requests to appear at the public hearing should be filed with the Secretary no later than 5:15 p.m., January 26, 2010, in accordance with the requirements in the "Submissions" section below. Persons wishing to appear should indicate in their request to appear whether they plan to provide testimony with respect to investigation No. 332-509, investigation No. 332-510, or both investigations. All pre-hearing briefs and statements should be filed not later than 5:15 p.m., January 28, 2010; and all post-hearing briefs and statements responding to matters raised at the hearing should be filed not later than 5:15 p.m., February 23, 2010. In the event that, as of the close of business on January 26, 2010, no witnesses are scheduled to appear at the hearing, the hearing will be canceled. Any person interested in attending the hearing as an observer or non-participant may call the Office of the Secretary (202-205-2000) after January 26, 2010, for information concerning whether the hearing will be held. The Commission is also considering holding additional hearings in Portland, Oregon and St. Louis, Missouri. Notice of the time, date, and place of those hearings will be published at a later date.

**Written Submissions:** In lieu of or in addition to participating in the hearing, interested parties are invited to file written submissions concerning this investigation. All written submissions should be addressed to the Secretary, and all such submissions (other than pre- and post-hearing briefs and statements) should be received not later than 5:15 p.m., May 28, 2010. All written submissions must conform with the provisions of section 201.8 of the Commission's Rules of Practice and Procedure (19 CFR 201.8). Section 201.8 requires that a signed original (or a copy so designated) and fourteen (14) copies of each document be filed. In the event that confidential treatment of a document is requested, at least four (4) additional copies must be filed, in which the confidential information must be deleted (see the following paragraph for further information regarding confidential business information). The Commission's rules authorize filing submissions with the Secretary by facsimile or electronic means only to the extent permitted by section 201.8 of the rules (see Handbook for Electronic Filing Procedures, <http://www.usitc.gov/secretary/>

[fed\\_reg\\_notices/rules/documents/handbook\\_on\\_electronic\\_filing.pdf](http://www.usitc.gov/fed_reg_notices/rules/documents/handbook_on_electronic_filing.pdf)). Persons with questions regarding electronic filing should contact the Office of the Secretary (202-205-2000).

Any submissions that contain confidential business information must also conform with the requirements of section 201.6 of the Commission's Rules of Practice and Procedure (19 CFR 201.6). Section 201.6 of the rules requires that the cover of the document and the individual pages be clearly marked as to whether they are the "confidential" or "non-confidential" version, and that the confidential business information be clearly identified by means of brackets. All written submissions, except for confidential business information, will be made available for inspection by interested parties.

In his request letter, the USTR stated that his office intends to make the Commission's reports available to the public in their entirety, and asked that the Commission not include any confidential business information or national security classified information in the reports that the Commission transmits to his office. Any confidential business information received by the Commission in this investigation and used in preparing this report will not be published in a manner that would reveal the operations of the firm supplying the information.

By order of the Commission.

Issued: December 7, 2009.

**Marilyn R. Abbott,**  
*Secretary to the Commission.*

[FR Doc. E9-29518 Filed 12-10-09; 8:45 am]  
**BILLING CODE 7020-02-P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a regulation under 21 U.S.C. 952(a)(2), authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on October 20, 2009, Tocris Cookson, Inc., 16144 Westwoods Business Park, Ellisville, Missouri 63021-4500, 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 schedules I and II:

Drug	Schedule
Marihuana (7360) .....	I
Tetrahydrocannabinols (7370) .....	I
4-Bromo-2, dimethoxyamphetamine. (7391) .....	5- I
3, Methyleneedioxymethamphetamine. (7405) .....	4- I
Amphetamine (1100) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Diprenorphine (9058) .....	II
Fentanyl (9801) .....	II

The company plans to import small quantities of the above listed controlled substances for non-clinical, laboratory-based research only.

In reference to drug code 7360 (Marihuana), the company plans to import synthetic cannabinoid agonists. In reference to drug code 7370 (Tetrahydrocannabinols), the company will import a synthetic Delta-9-THC. No other activity for these drug codes are authorized for this registration.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than January 11, 2010.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the **Federal Register** on September 23, 1975, (40 FR 43745-46), all applicants for registration to import a basic class of any controlled substances in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21

CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: December 1, 2009.

**Joseph T. Rannazzisi,**

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

[FR Doc. E9-29542 Filed 12-10-09; 8:45 am]  
**BILLING CODE 4410-09-P**

## DEPARTMENT OF JUSTICE

### Office of Justice Programs

[OJP (NIJ) Docket No. 1510]

### Vehicular Digital Multimedia Evidence Recording System Standard Special Technical Committee

**AGENCY:** National Institute of Justice.

**ACTION:** Notice of request for proposals for certification and testing expertise.

**SUMMARY:** The National Institute of Justice (NIJ) is in the process of developing a new Vehicular Digital Multimedia Evidence Recording System Standard and corresponding certification program requirements. This work is being performed by a Special Technical Committee (STC), comprised of practitioners from the field, researchers, testing experts, certification experts, and representatives from stakeholder organizations. It is anticipated that the STC members will participate in six 2-day meetings over a 9-month time period with the goal of completing development of the standard and certification program requirements. It is anticipated that STC meetings will begin in mid-January 2010. Travel expenses and per diem will be reimbursed for all STC meetings; however, participation time will not be funded.

NIJ is seeking representatives from (1) certification bodies and (2) test laboratories with experience in programs for similar types of electronic equipment. Additional preferred knowledge includes experience with in-car video systems or experience with law enforcement operations. There are up to four positions to be filled on the STC, and NIJ will accept the first 20 submissions for review.

Interested parties are requested to nominate individuals from their organizations and submit no more than two pages describing the nominee's applicable experience, preferred knowledge, and affiliations with standards development organizations. This information shall be submitted to Frances Scott at [frances.scott@usdoj.gov](mailto:frances.scott@usdoj.gov) by December 22, 2009. The submissions will be reviewed, and participants will be notified regarding their acceptance by January 8, 2009.

### FOR FURTHER INFORMATION CONTACT:

Cassandra Robinson by telephone at 202-305-2296 [Note: this is not a toll-free

Drug	Schedule
Tetrahydrocannabinols (7370) .....	I
Dihydromorphine (9145) .....	I
Dihydrocodeine (9120) .....	II
Oxycodone (9143) .....	II
Hydromorphone (9150) .....	II
Hydrocodone (9193) .....	II
Remifentanil (9739) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The company plans to manufacture the listed controlled substances in bulk for distribution to its customers.

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.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substances, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than February 9, 2010.

Dated: December 1, 2009.

**Joseph T. Rannazzisi,**

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

[FR Doc. E9-29527 Filed 12-10-09; 8:45 am]  
**BILLING CODE 4410-09-P**