Persons filing written submissions must file the original document and 12 true copies thereof on or before the deadlines stated above with the Office of the Secretary. Submissions should refer to the docket number (“Docket No. 2818”) in a prominent place on the cover page and/or the first page. 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電子 filing.pdf).

Persons with questions regarding electronic filing should contact the Secretary (202–205–1810). 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. All nonconfidential written submissions will be available for public inspection at the Office of the Secretary. This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of sections 201.10 and 210.50(a)(4) of the Commission’s Rules of Practice and Procedure (19 CFR 201.10, 210.50(a)(4)).

By order of the Commission.
Issued: June 15, 2011.

James R. Holbein,
Secretary to the Commission.

[FR Doc. 2011–15242 Filed 6–17–11; 8:45 am]
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). Persons with questions regarding electronic filing should contact 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 its request letter, the Committee stated that it intends to make the Commission’s report available to the public in its entirety, and asked that the Commission not include any confidential business information in the report that the Commission sends to the Committee. 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.

Issued: June 15, 2011.

By order of the Commission.

James R. Holbein,
Secretary to the Commission.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[OMB Number 1117–0010]

Agency Information Collection Activities: Proposed Collection; Comments Requested: U.S. Official Order Forms for Schedule I and II Controlled Substances (Accountable Forms); Order Form Requisition; DEA Form 222, 222a, Controlled Substances Order System

ACTION: 30-Day notice of information collection under review.

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 at Volume 76, Number 71, Page 20710, April 13, 2011, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until July 20, 2011. This process is conducted in accordance with 5 CFR 1320.10.

If you have comments, especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Cathy A. Gallagher, Acting Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152; (202) 307–7297.

Written comments concerning this information collection should be sent to the Office of Information and Regulatory Affairs, Office of Management and Budget, Attn: DOJ Desk Officer. The best way to ensure your comments are received is to e-mail them to oira_submission@omb.eop.gov or fax them to (202) 395–7285. All comments should reference the eight-digit OMB number for the collection or the title of the collection. If you have questions concerning the collection, please contact Cathy A. Gallagher, Acting Chief, Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, 8701 Morrissette Drive, Springfield, VA 22152; (202) 307–7297, or the DOJ Desk Officer at (202) 395–3176.

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 1117–0010

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

(2) Title of the Form/Collection: U.S. Official Order Forms for Schedule I and II Controlled Substances (Accountable Forms); Order Form Requisition.

(3) Agency form number, if any, and the applicable component of the Department sponsoring the collection: Form number: DEA Forms 222 and 222a.

Component: Office of Diversion Control, Drug Enforcement Administration, Department of Justice.

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

Primary: Business or other for-profit.

Other: Not-for-profit; State, local or Tribal government.

Abstract: DEA–222 is used to transfer or purchase Schedule I and II controlled substances and data are needed to provide an audit of transfer and purchase. DEA–222a Requisition Form is used to obtain the DEA–222 Order Form. Persons may also digitally sign and transmit orders for controlled substances electronically, using a digital certificate. Orders for Schedule I and II controlled substances are archived and transmitted to DEA.

(5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: DEA estimates that 109,632