stationery. Requests should identify the name, title, and company or other organizational affiliation (if any), address, telephone number, email address, and industry or main line of business of the company, if any, of the person signing the request letter and of the persons who plan to appear at one or both hearings. Requests to appear may be made by mail or delivered in person to the Commission’s Office of the Secretary [(see ADDRESSES), or may be filed by email sent to SMEHearing@usitc.gov. The Commission does not accept requests filed by fax.

Written Submissions: In lieu of or in addition to participating in the hearing, interested parties are invited to file written submissions concerning this investigation. Such submissions should be addressed to the secretary, and should be received no later than 5:15 p.m., October 15, 2013. All written submissions must conform to 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 interested parties file documents electronically on or before the filing deadline and submit eight (8) true paper copies by 12:00 p.m. eastern time on the next business day. In the event that confidential treatment of a document is requested, interested parties must file, at the same time as the eight paper copies, at least four (4) additional true paper copies in which the confidential information must be deleted (see the following paragraph for further information regarding confidential business information). Persons with questions regarding electronic filing should contact the Secretary (202–205–2000).

Any submissions that contain confidential business information (CBI) must also conform to 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 is clearly identified by means of brackets. All written submissions, except for confidential business information, will be made available for inspection by interested parties.

In the request letter, the USTR stated that the Office of the USTR 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 or national security classified information in the report that the Commission sends to the USTR. 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: August 16, 2013
Lisa R. Barton.
Acting Secretary to the Commission.

[FR Doc. 2013–20388 Filed 8–20–13; 8:45 am]
BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; NORAMCO, Inc.

Pursuant to Title 21 Code of Federal Regulations 1301.34 (a), this is notice that on June 27, 2013, Noramco, Inc., 1440 Olympic Drive, Athens, Georgia 30601, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the basic classes of controlled substances:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Phenylacetone (8501)</td>
<td>II</td>
</tr>
<tr>
<td>Thebaine (9333)</td>
<td>II</td>
</tr>
<tr>
<td>Poppy Straw Concentrate (9670)</td>
<td>II</td>
</tr>
<tr>
<td>Tapentadol (9780)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import Thebaine (9333) analytical standards for distribution to its customers. The company plans to import an intermediate form of Tapentadol (9780) to bulk manufacture Tapentadol for distribution to its customers. The company plans to import the Phenylacetone (8501) in bulk for the manufacture of a controlled substance.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (2007).

In reference to the non-narcotic raw material, any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedules I or II, which fall under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), 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 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 September 20, 2013.

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: August 14, 2013.
Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–20285 Filed 8–20–13; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Office of Justice Programs

[OJP (BJA) Docket No. 1629]

Meeting of the Department of Justice’s (DOJ’s) National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee

AGENCY: Office of Justice Programs (OJP), Justice.
ACTION: Notice of meeting.

SUMMARY: This is an announcement of a meeting of DOJ’s National Motor Vehicle Title Information System (NMVTIS) Federal Advisory Committee to discuss various issues relating to the operation and implementation of NMVTIS.

DATES: The meeting will take place on Tuesday October 8, 2013, from 10:00 a.m. to 4:00 p.m. ET.

ADDRESSES: The meeting will take place at the Office of Justice Programs (OJP), 810 7th Street NW., Washington, DC 20531.

FOR FURTHER INFORMATION CONTACT: Todd Brighton, Designated Federal Employee (DFE), Bureau of Justice
I. Background

Under Section 103(g) of the Federal Mine Safety and Health Act of 1977, as amended (Mine Act), a representative of miners, or any individual miner where there is no representative of miners, may submit a written or oral notification of an alleged violation of the Mine Act or a mandatory standard that an imminent danger exists. The notifier has the right to obtain an immediate inspection by the Mine Safety and Health Administration (MSHA). A copy of the notice must be provided to the operator, with individual miner names redacted.

MSHA regulations at 30 CFR Part 43 implement Section 103(g) of the Mine Act. These regulations provide the procedures for submitting notification of the alleged violation or imminent danger and the actions that MSHA must take after receiving the notice. Although the regulations contain a review procedure (required by Section 103(g)(2) of the Mine Act) whereby a miner or a representative of miners may in writing request a review if no citation or order is issued as a result of the original notice, the option is so rarely used that it was not considered in the burden estimates.

II. Desired Focus of Comments

MSHA is soliciting comments concerning the proposed extension of the information collection related to Hazardous Conditions Complaints in 30 CFR 43.4 and 43.7. MSHA is particularly interested in comments that:

- Evaluate whether the collection of information is necessary for the proper performance of the functions of the agency, including whether the information has practical utility;
- Evaluate the accuracy of the MSHA’s estimate of the burden of the collection of information, including the validity of the methodology and assumptions used;
- Suggest methods to 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.

This proposed information collection request is available on MSHA’s Web site at http://www.msha.gov under “Federal Register Documents” on the right side of the screen by selecting “New and Existing Information Collections and Supporting Statements”. This proposed information collection request will be available on MSHA’s Web site for 60 days after the publication date of this notice, and on http://www.regulations.gov. Because comments will not be edited to remove any identifying or contact information, MSHA cautions the commenter against including any information in the submission that should not be publicly disclosed.

The public may also examine the proposed information collection at MSHA, Office of Standards, Regulations, and Variances, 1100 Wilson Boulevard, Room 2350, Arlington, VA 22209–3939 by signing in at the receptionist’s desk on the 21st floor.