

respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on January 18, 2017, Stepan Company, Natural Products Dept., 100 W. Hunter Avenue, Maywood, New Jersey 07607 applied to be registered as an importer of coca leaves (9040), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance in bulk for the manufacture of controlled substance for distribution to its customers.

Dated: August 21, 2017.

Demetra Ashley,

Acting Assistant Administrator.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Cambrex Charles City

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before September 28, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before September 28, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and

(2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. Comments and request for hearing on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Office of Diversion Control (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on June 12, 2017, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Phenylacetone	8501	II
Coca Leaves	9040	II
Opium, raw	9600	II
Poppy Straw Concentrate	9670	II

The company plans to import the listed controlled substances for internal use, and to manufacture bulk intermediates for sale to its customers.

Dated: August 21, 2017.

Demetra Ashley,

Acting Assistant Administrator.

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applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before October 30, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled

substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on July 6, 2017, Noramco, Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801-4417 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Codeine-N-oxide	9053	I
Dihydromorphine	9145	I
Hydromorphanol	9301	I

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Noramco, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and

Controlled substance	Drug code	Schedule
Morphine-N-oxide	9307	I
Amphetamine	1100	II
Methylphenidate	1724	II
Nabilone	7379	II
Phenylacetone	8501	II
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	II
Morphine	9300	II
Oripavine	9330	II
Thebaine	9333	II
Opium extracts	9610	II
Opium fluid extract	9620	II
Opium tincture	9630	II
Opium, powdered	9639	II
Opium, granulated	9640	II
Oxymorphone	9652	II
Noroxymorphone	9668	II
Tapentadol	9780	II

The company plans to manufacture the above-listed controlled substances in bulk for distribution to its customers. In reference to drug code 7360, the company plans to manufacture a synthetic version of cannabidiol in bulk for sale to its customers, who are final dosage form manufacturers. No other activity for this drug code is authorized for this registration.

Dated: August 21, 2017.

Demetra Ashley,

Acting Assistant Administrator.

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DEPARTMENT OF JUSTICE

[OMB Number 1125-0005]

Agency Information Collection Activities; Proposed Collection; Comments Requested; Request To Be Included on the List of Pro Bono Legal Service Providers for Individuals in Immigration Proceedings (Form EOIR-56)

AGENCY: Executive Office for Immigration Review, Department of Justice.

ACTION: 30-day Notice.

SUMMARY: The Department of Justice (DOJ), Executive Office for Immigration Review (EOIR), 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 was previously published in the **Federal Register** on June 26, 2017, allowing for a 60-day comment period.

DATES: Comments are encouraged and will be accepted for additional days until September 28, 2017.

FOR FURTHER INFORMATION CONTACT: 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 Jean King, General Counsel, USDOJ-EOIR-OGC, Suite 2600, 5107 Leesburg Pike, Falls Church, Virginia, 20530; telephone: (703) 305-0470. Written comments and/or suggestions can also be sent to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503 or sent to *OIRA_submissions@omb.eop.gov*.

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;
- Enhance the quality, utility, and clarity of the information to be collected; and/or
- 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 *The Title of the Form/Collection:* Request to be Included on the List of Pro Bono Legal Service Providers for Individuals in Immigration Proceedings.

3 *The agency form number:* EOIR-56 (OMB #1125-0015).

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

Primary: Legal service providers seeking to be included on the List of Pro Bono Legal Service Providers (“List”), a

list of persons who have indicated their availability to represent aliens on a pro bono basis. Abstract: EOIR seeks to replace the current paper version of the EOIR Forms-56, with an electronic system to make an initial application and apply for continued participation in the List. Form EOIR-56 will be mandatory, and is intended to elicit, in a uniform manner, all of the required information for EOIR to determine whether an applicant meets the eligibility requirements for inclusion on the List.

5 *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 161 respondents will complete each form within approximately 30 minutes.

6 *An estimate of the total public burden (in hours) associated with the collection:* 80.5 annual burden hours. If additional information is required contact: Jake Bishop-Green, Acting Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., 3E.405B, Washington, DC 20530.

Dated: August 24, 2017.

Jake Bishop-Green,

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

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DEPARTMENT OF JUSTICE

[OMB Number 1121-0220]

Agency Information Collection Activities; Proposed New e-Collection; Bureau of Justice Assistance Application Form: Public Safety Officers’ Benefits (PSOB) Program Applications Package

AGENCY: Bureau of Justice Assistance, Department of Justice.

ACTION: 30-day notice.

SUMMARY: The Department of Justice, Office of Justice Programs, Bureau of Justice Assistance, will be submitting the following information collection request for review and clearance in accordance with the Paperwork Reduction Act of 1995.

DATES: Comments are encouraged and will be accepted for 30 days until September 28, 2017.

FOR FURTHER INFORMATION CONTACT: If you have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the