

**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 March 17, 2017, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816 applied to be registered as a bulk manufacturer the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Dihydromorphine .....	9145	I
Methylphenidate .....	1724	II
Codeine .....	9050	II
Dihydrocodeine .....	9120	II
Oxycodone .....	9143	II
Hydromorphone .....	9150	II
Hydrocodone .....	9193	II
Levorphanol .....	9220	II
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Tapentadol .....	9780	II
Fentanyl .....	9801	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to finished dosage form manufacturers.

In reference to drug code 7360 and 7370, the company plans to bulk manufacture a synthetic CBD and tetrahydrocannabinol.

No other activity for drug code 7360 and 7370 are authorized for this registration.

Dated: March 15, 2018.

**Susan A. Gibson,**

*Deputy Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Noramco, Inc.**

**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 on or before April 20, 2018. Such persons may also file a written request for a hearing on the application on or before April 20, 2018.

**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 requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

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Controlled substance	Drug code	Schedule
Marihuana .....	7360	I
Tetrahydrocannabinols .....	7370	I
Nabilone .....	7379	II
Phenylacetone .....	8501	II
Opium, raw .....	9600	II
Poppy Straw Concentrate .....	9670	II
Tapentadol .....	9780	II

The company plans to import phenylacetone (8501), opium, raw (9600), and poppy straw concentrate (9670) to bulk manufacture other controlled substances for distribution to its customers. The company plans to import an intermediate form of tapentadol (9780) to bulk manufacture tapentadol (9780) for distribution to its customers.

In reference to drug codes 7360 and 7370, the company plans to import a synthetic cannabidiol and a synthetic tetrahydrocannabinol. No other activity for these drug codes is authorized for this registration. Placement of these drug codes onto the company’s registration does not translate into automatic approval of subsequent permit applications to import controlled substances. Approval of permit applications will occur only when the registrant’s business activity is consistent with what is authorized under 21 U.S.C 952(a)(2). Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: March 15, 2018.

**Susan A. Gibson,**

*Deputy Assistant Administrator.*

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**DEPARTMENT OF LABOR**

**Employment and Training Administration**

**Agency Information Collection Activities; Comment Request; Senior Community Service Employment Program (SCSEP)**

**ACTION:** Notice of information collection; request for comment.

**SUMMARY:** The Department of Labor (DOL), Employment and Training Administration (ETA) is soliciting comments concerning a proposed extension for the authority to conduct the information collection request (ICR) titled, “Senior Community Service Employment Program (SCSEP).” This comment request is part of continuing Departmental efforts to reduce