



DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Bulk Manufacturer of Controlled Substances Application: Noramco Inc.

[Docket No. DEA-526]

ACTION: Notice of application.

DATES: Registered bulk manufacturer 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 [INSERT 60 DAYS AFTER PUBLICATION IN THE *FEDERAL REGISTER*].

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

SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.33(a), this is notice that on August 6, 2019, 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
Hydromorphenol	9301	I
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 listed controlled substances as an Active Pharmaceutical Ingredient (API) for supply to its customers. In reference to drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetics. No other activities for these drug codes are authorized for this registration.

Dated: October 22, 2019 .

William T. McDermott,
Assistant Administrator.

[FR Doc. 2019-24106 Filed: 11/4/2019 8:45 am; Publication Date: 11/5/2019]