for a hearing on the application on or before December 21, 2020.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 6, 2020, Noramco Inc. 500 Swedes Landing Road, Wilmington, Delaware 19801–4417, applied to be registered as an bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I
Codeine-N-oxide	9053	I
Dihydromorphine	9145	I
Hydromorphinol	9301	I
Morphine-N-oxide	9307	I
Amphetamine	1100	II
Lisdexamfetamine	1205	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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–23396 Filed 10–21–20; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–733]

Bulk Manufacturer of Controlled Substances Application: Kinetochem LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Kinetochem LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before December 21, 2020. Such persons may also file a written request for a hearing on the application on or before December 21, 2020.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 5, 2020, Kinetochem LLC, 111 W Cooperative Way, Suite 310–B, Georgetown, Texas 78626, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

The company plans to synthetically manufacture drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinols), in bulk for distribution and sale to its customers. No other activities for these drugs are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020–23395 Filed 10–21–20; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Office of Federal Contract Compliance Programs

RIN 1250–ZA01

Request for Information; Race and Sex Stereotyping and Scapegoating

AGENCY: Office of Federal Contract Compliance Programs

ACTION: Request for information

SUMMARY: The Office of Federal Contract Compliance Programs (OFCCP) at the Department of Labor seeks comments, information, and materials from the public relating to workplace trainings that involve race or sex stereotyping or