minutes or the time taken to prepare each response).

If additional information is required contact: Robert Houser, Department Clearance Officer, Policy and Planning Staff, Office of the Chief Information Officer, Justice Management Division, United States Department of Justice, Two Constitution Square, 145 N Street NE, 3.E–206, Washington, DC 20530.

Dated: October 25, 2022.

Robert Houser,

Department Clearance Officer, Policy and Planning Staff, Office of the Chief Information Officer, U.S. Department of Justice. [FR Doc. 2022–23591 Filed 10–28–22; 8:45 am]

BILLING CODE 4410-FY-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1107]

Bulk Manufacturer of Controlled Substances Application: Nanosyn Inc.

AGENCY: Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

SUMMARY: Nanoysn Inc. 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 submit electronic comments on or objections to the issuance of the proposed registration on or before December 30, 2022. Such persons may also file a written request for a hearing on the application on or before December 30, 2022.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to https://www.regulations.gov and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on https://www.regulations.gov. If vou have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this

is notice that on September 2, 2022, Nanosyn Inc., 3331 Industrial Drive, Suite B, Santa Rosa, California 95403– 2062, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Oxymorphone	9652	
Fentanyl	9801	

The company is a contract manufacturer. At the request of the company's customers, it manufactures derivatives of the above controlled substances in bulk form. No other activities for these drug codes are authorized for this registration.

Kristi O'Malley,

Assistant Administrator. [FR Doc. 2022–23609 Filed 10–28–22; 8:45 am] BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1109]

Bulk Manufacturer of Controlled Substances Application: Noramco

AGENCY: Drug Enforcement Administration, Justice. **ACTION:** Notice of application.

SUMMARY: Noramco 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 submit electronic comments on or objections to the issuance of the proposed registration on or before December 30, 2022. Such persons may also file a written request for a hearing on the application on or before December 30, 2022.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to https://www.regulations.gov and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on https://www.regulations.gov. If

you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 29, 2022, Noramco, 500 Swedes Landing Road, Wilmington, Delaware 19801–4417, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
Marihuana	7360	1
Tetrahydrocannabinols	7370	I
Codeine-N-oxide	9053	I
Dihydromorphine	9145	I
Hydromorphinol	9301	1
Morphine-N-oxide	9307	1
Amphetamine	1100	II
Lisdexamfetamine	1205	П
Methylphenidate	1724	П
Nabilone	7379	П
Phenylacetone	8501	П
Codeine	9050	П
Dihydrocodeine	9120	П
Oxycodone	9143	П
Hydromorphone	9150	П
Hydrocodone	9193	П
Morphine	9300	П
Oripavine	9330	II
Thebaine	9333	П
Opium extracts	9610	П
Opium fluid extract	9620	П
Opium, tincture	9630	П
Opium, powdered	9639	П
Opium, granulated	9640	II
Oxymorphone	9652	II
Noroxymorphone	9668	П
Tapentadol	9780	II

The company plans to bulk 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 synthetic. No other activities for these drug codes are authorized for this registration.

Kristi O'Malley,

Assistant Administrator. [FR Doc. 2022–23614 Filed 10–28–22; 8:45 am] BILLING CODE 4410–09–P