

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]

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**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:** Nanoyns 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 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 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	II
Fentanyl .....	9801	II

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]

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**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	I
Tetrahydrocannabinols	7370	I
Codeine-N-oxide .....	9053	I
Dihydromorphine .....	9145	I
Hydromorphanol .....	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 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.*

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