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Rebecca Curtiss,

Land Law Examiner, Adjudication Section.

[FR Doc. 2022–23561 Filed 10–28–22; 8:45 am]

BILLING CODE 4331–10–P

DEPARTMENT OF THE INTERIOR

National Park Service

[NPS–FOVA–NPS33107; PPPWFOVAAO
PPMPSAS1Z.Y00000]

Fort Vancouver National Historic Site; Relinquishment of Exclusive Jurisdiction

AGENCY: National Park Service, Interior.

ACTION: Notice.

SUMMARY: On behalf of the United States, the National Park Service has relinquished jurisdiction to the State of Washington in order to establish concurrent legislative jurisdiction over certain lands owned and administered by the National Park Service, known as the East and South Barracks, at Fort Vancouver National Historic Site.

DATES: Concurrent legislative jurisdiction within the East and South Barracks at Fort Vancouver National Historic Site became effective on October 5, 2022.

FOR FURTHER INFORMATION CONTACT:

Tracy Fortmann, Superintendent, Fort Vancouver National Historic Site, 612 E Reserve Street, Vancouver, WA 98661–3811; *telephone:* 360–816–6205; *email:* Tracy_Fortmann@nps.gov.

SUPPLEMENTARY INFORMATION: On June 8, 1981, the Director of the National Park Service (NPS) relinquished jurisdiction and the Governor of the State of Washington accepted concurrent jurisdiction at Fort Vancouver National Historic Site. On May 22, 2012, the Department of Army transferred the East and South Barracks (“Barracks”) to the NPS for inclusion in the Fort Vancouver National Historic Site. On October 5, 2022, acting in accordance with the provisions of 54 U.S.C. 100754 and Revised Code of Washington section 37.04.050, the Director of the NPS relinquished jurisdiction and the Governor of the State of Washington accepted concurrent jurisdiction over the Barracks, in order to enable the United States and State of Washington to exercise concurrent jurisdiction over the Barracks consistent with the

remainder of the Fort Vancouver National Historical Site.

Jennifer Flynn,

Associate Director, Visitor and Resource Protection, National Park Service.

[FR Doc. 2022–23659 Filed 10–28–22; 8:45 am]

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DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms and Explosives

[OMB Number 1140–0107]

Agency Information Collection Activities; Proposed eCollection of eComments Requested; Extension of a Currently Approved Collection; National Firearms Act (NFA) Responsible Person Questionnaire— ATF Form 5320.23

AGENCY: Bureau of Alcohol, Tobacco, Firearms and Explosives, Department of Justice.

ACTION: 30-Day notice.

SUMMARY: The Bureau of Alcohol, Tobacco, Firearms and Explosives (ATF), Department of Justice (DOJ) will submit the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995.

DATES: Comments are encouraged and will be accepted for an additional 30 days until November 30, 2022.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to www.reginfo.gov/public/do/PRAMain. Find this particular information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function.

SUPPLEMENTARY INFORMATION: Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
- Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

- Evaluate whether and, if so, how the quality, utility, and clarity of the information to be collected can be enhanced; and
- Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

1. *Type of Information Collection:* Extension with Revision of a Currently Approved Collection.

2. *The Title of the Form/Collection:* National Firearms Act (NFA) Responsible Person Questionnaire.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number: ATF Form 5320.23.

Component: Bureau of Alcohol, Tobacco, Firearms and Explosives, U.S. Department of Justice.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:*

Primary: Business or other for-profit, Federal Government, State Local or Tribal Government.

Other: Not-for-profit institutions and Farms.

Abstract: The National Firearms Act (NFA) Responsible Person Questionnaire—ATF Form 5320.23 (ATF Form 5320.23) must be completed by a responsible person (RP), identified as part of a trust or legal entity on the Application to Make and Register a Firearm—ATF Form 1 (5320.1) (ATF Form 1). This form must also be completed by a RP who is the identified as the firearm maker or the transferee on the Application for Tax Paid Transfer and Registration of Firearm—ATF Form 4 (5320.4) (ATF Form 4), or the Application for Tax Exempt Transfer of Firearm—ATF Form 5 (5320.5) ATF Form 5.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 115,829 respondents will respond to this collection once annually, and it will take each respondent approximately 30 minutes to complete their responses.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated annual public burden associated with this collection is 57,914.5 or 57,915 hours, which is equal to 115,829 (total respondents) * 1 (# of response per respondent) * .5 (30

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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