Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

The company plans to bulk manufacture the listed controlled substance for the internal use intermediates or for sale to its customers. In reference to drug code 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture this drug as synthetic. No other activity for this drug code is authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–21724 Filed 11–28–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1628]

Importer of Controlled Substances Application: Blue Rabbit Veterinary LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Blue Rabbit Veterinary LLC has applied to be registered as an importer 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 31, 2025.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal e rulemaking 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. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on October 28, 2025, Blue Rabbit Veterinary LLC, 1680 East Northrop Boulevard, Unit 1 Chandler, Arizona 85286, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Etorphine HCl	9059	II
Thiafentanil	9729	II

The company plans to import the listed controlled substance for the purpose of distribution in final dosage form to prospective zoo and wildlife customers. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration approved or non-approved finished dosage forms for commercial sale.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–21719 Filed 11–28–25; 8:45 am]

BILLING CODE P

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 29, 2025, Invizyne Technologies, Inc., 750 Royal Oaks Drive, Suite 106, Monrovia, California 91016–6357, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Tetrahydrocannabinols	7370	I

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1629]

Bulk Manufacturer of Controlled Substances Application: Benuvia Operations, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Benuvia Operations, 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 submit electronic comments on or objections to the issuance of the proposed registration on or before January 30, 2026. Such persons may also file a written request for a hearing on the application on or before January 30, 2026.

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 October 27, 2025, Benuvia Operations, LLC., 3950 North Mays Street, Round Rock, Texas 78665, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Lysergic Acid Diethylamide.	7315	I
Codeine	9050	II
Hydromorphone	9150	II
Sufentanil	9740	II

The company plans to bulk manufacture the listed controlled substances for internal research and dosage formulation development. No other activities for these drug codes are authorized for this registration.

Thomas Prevoznik,

Deputy Assistant Administrator.

[FR Doc. 2025–21720 Filed 11–28–25; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

[OMB 1140–0005]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Previously Approved Collection Application/Permit To Import Firearms, Ammunition, and Defense Articles—ATF Form 5330.3A (Form 6, Part I)

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

ACTION: 60-Day notice.

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

DATES: ATF encourages comments on this information collection. You may submit written comments until midnight on January 30, 2026.

ADDRESSES: Submit written comments and recommendations for this information collection, especially on the estimated public burden or associated response time, to Justine Hall, Firearms and Explosives Imports Branch, by email to *justine.hall@atf.gov,* or by mail to 244 Needy Road; Martinsburg, WV 25405. Identify comments by the OMB control number 1140–0005. You may view the proposed information collection instrument online at *https://www.atf.gov/rules-and-regulations/federal-register-actions/forms-and-information-collection.*

FOR FURTHER INFORMATION CONTACT: If you have questions, or need a copy of the proposed information collection instrument with instructions or additional information, contact: Justine Hall, Firearms and Explosives Imports Branch, either by mail at 244 Needy Road; Martinsburg, WV 25405, by email at *justine.hall@atf.gov,* or by telephone at 304–616–4593.

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

—Evaluate whether the proposed information collection is necessary to properly perform ATF's functions, including whether the information will have practical utility;

—Evaluate the agency's estimate of the proposed information collection's burden for accuracy, including validity of the methodology and assumptions used;

—Evaluate whether, and if so, how, the quality, utility, and clarity of the collected information can be enhanced; and

—Minimize the information collection's burden on those who are to respond, including using appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting people to submit electronic responses.

Overview of This Information Collection

1. *Abstract:* The Gun Control Act and other statutes prohibit persons from importing firearms, ammunition, and defense articles into the US unless the articles meet certain criteria, the importer is not a prohibited person, and the Attorney General (delegated to Director, ATF) approves that person to import those items. ATF uses this form to collect the information necessary to determine the above facts and approve a person to import the articles; the form then serves as their permit to do so.

2. *Type of information collection:* revision of a previously approved collection.

3. *Title of the form/collection:* Application/Permit to Import Firearms, Ammunition, and Defense Articles.

4. *Agency form number, if any, and the applicable component of the Department sponsoring the collection:*

Form number: ATF Form 5330.3A (''Form 6, part I'').

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

5. *Affected public who will be asked or required to respond, as well as the obligation to respond:*

Affected public: individuals or households, state, local and tribal governments, private sector-for-profit institutions, and federal government.

Obligation to respond: required to obtain or retain a benefit.

6. *Estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 13,777 respondents will respond to this information collection once annually, and it will take each respondent an average of 17 minutes (0.281 hours) to complete their responses (approximately 30 minutes for persons submitting a paper form, or 15 minutes for persons submitting electronically).

7. *Estimate of the total annual burden (in hours) associated with the collection:* The estimated annual public burden associated with this collection is 3,869 total hours, which is equal to 13,777 (total respondents) * 1 (# of responses per respondent) * 0.281 (17 minutes).

8. *Estimate of the total annual other cost burden associated with the collection, if applicable:* $1,241.

ESTIMATED TOTAL HOURLY BURDEN

Activity	Number of respondents	Frequency	Total annual responses	Time per response (hours)	Total annual burden (hours)
Paper form	1,700	1	1,700	.5	850
Electronic form	12,077	1	12,077	.25	3,019
Totals	13,777	1	13,777	*.281	3,869

* (Combined average; roughly 17 minutes.)