

Controlled substance	Drug code	Schedule
Nabilone .....	7379	II
1-Phenylcyclohexylamine .....	7460	II
Phencyclidine .....	7471	II
ANPP (4-Anilino-N-phenethyl-4-piperidine) .....	8333	II
Norfentanyl (N-phenyl-N-(piperidin-4-yl) propionamide) .....	8366	II
Phenylacetone .....	8501	II
1-Piperidinocyclohexanecarbonitrile .....	8603	II
Alphaprodine .....	9010	II
Anileridine .....	9020	II
Coca Leaves .....	9040	II
Cocaine .....	9041	II
Etorphine HCl .....	9059	II
Dihydrocodeine .....	9120	II
Diphenoxylate .....	9170	II
Ecgonine .....	9180	II
Ethylmorphine .....	9190	II
Levomethorphan .....	9210	II
Levorphanol .....	9220	II
Isomethadone .....	9226	II
Meperidine .....	9230	II
Meperidine intermediate-A .....	9232	II
Meperidine intermediate-B .....	9233	II
Meperidine intermediate-C .....	9234	II
Metazocine .....	9240	II
Oliceridine (N-[(3-methoxythiophen-2-yl)methyl] ({2-[9r]-9-(pyridin-2-yl)-6-oxaspiro[4.5] decan-9-yl} ethyl {time})amine fumarate) .....	9245	II
Metopon .....	9260	II
Dextropropoxyphene, bulk (non-dosage forms) .....	9273	II
Dihydroetorphine .....	9334	II
Opium tincture .....	9630	II
Opium, powdered .....	9639	II
Opium, granulated .....	9640	II
Noroxymorphone .....	9668	II
Phenazocine .....	9715	II
Thiafentanil .....	9729	II
Piminodine .....	9730	II
Racemethorphan .....	9732	II
Racemorphan .....	9733	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Bezitramide .....	9800	II
Moramide-intermediate .....	9802	II

The company plans to import small quantities of the listed controlled substances to support research activities funded by the National Institute on Drug Abuse. 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.

**Marsha Ikner,**

*Acting Deputy Assistant Administrator.*

[FR Doc. 2024-07037 Filed 4-2-24; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1339]

#### Importer of Controlled Substances Application: Purisys, LLC

**AGENCY:** Drug Enforcement Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Purisys, 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 May 3, 2024. Such persons may also file a written request for a hearing on the application on or before May 3, 2024.

**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. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on February 8, 2024, Purisys, LLC, 1550 Olympic Drive, Athens, Georgia 30601-1602, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract .....	7350	I
Marihuana .....	7360	I
Tetrahydrocannabinols ..	7370	I
Nabilone .....	7379	II
Phenylacetone .....	8501	II
Ecgonine .....	9180	II
Levorphanol .....	9220	II
Thebaine .....	9333	II
Opium, raw .....	9600	II
Opium, powdered .....	9639	II
Opium, granulated .....	9640	II
Noroxymorphone .....	9668	II
Poppy Straw Con- centrate.	9670	II
Tapentadol .....	9780	II

The company plans to import Opium, Raw (9600), Opium, Powered (9639) and Opium, Granulated (9640) to manufacture an Active Pharmaceutical Ingredient (API) only for distribution to its customers. The company plans to import Phenylacetone (8501) and Poppy Straw Concentrate (9670), to bulk manufacture other Controlled substances for distribution to its customers. The company plans to import impurities of buprenorphine that have been determined by DEA to be captured under Thebaine (9333). In reference to Marihuana Extract (7350), Marihuana (7360) and Tetrahydrocannabinols (7370) the company plans to import as synthetic. No other activity 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.

**Marsha Ikner,**  
*Acting Deputy Assistant Administrator.*  
[FR Doc. 2024-07035 Filed 4-2-24; 8:45 am]

**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**[OMB Number 1110-0073]**

**Agency Information Collection Activities; Proposed eCollection eComments Requested; Crime Data Explorer (CDE) Feedback**

**AGENCY:** Federal Bureau of Investigation, Department of Justice.  
**ACTION:** 30-Day notice.

**SUMMARY:** The Federal Bureau of Investigation (FBI), Department of Justice (DOJ), will be submitting 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. The proposed information collection was previously published in the **Federal Register** on February 20, 2024, allowing a 60-day comment period.

**DATES:** Comments are encouraged and will be accepted for 30 days until May 3, 2024.

**FOR FURTHER INFORMATION CONTACT:** If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Edward L. Abraham, Crime and Law Enforcement Statistics Unit Chief, FBI, CJIS Division, Module D-1, 1000 Custer Hollow Road, Clarksburg, West Virginia 26306; [elabraham@fbi.gov](mailto:elabraham@fbi.gov), 304-625-4830.

**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;

- Enhance the quality, utility, and clarity of the information to be collected; and/or
- 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.

Written comments and recommendations for this information collection should be submitted within 30 days of the publication of this notice on the following website [www.reginfo.gov/public/do/PRAMain](http://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 and entering either the title of the information collection or the OMB Control Number 1110-0073. This information collection request may be viewed at [www.reginfo.gov](http://www.reginfo.gov). Follow the instructions to view Department of Justice, information collections currently under review by OMB.

DOJ seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOJ notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

**Overview of This Information Collection**

1. *Type of Information Collection:* Extension of a previously approved collection.
2. *Title of the Form/Collection:* Crime Data Explorer Feedback Survey.
3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* There is no form number for this collection. The applicable component within DOJ is the Criminal Justice Information Services (CJIS) Division, FBI.
4. *Affected public who will be asked or required to respond, as well as a brief abstract:*
  - Primary:* Law enforcement, academia, and the general public.
  - Abstract:* This survey is needed to collect feedback on the functionality of the CDE in order to make improvements to the application.
5. *Obligation to Respond:* Voluntary.
6. *Total Estimated Number of Respondents:* 200 respondents.
7. *Estimated Time per Respondent:* 2 minutes.