

exceed fifteen (15) pages of textual material, double-spaced and single-sided, when printed out on pages measuring 8.5 x 11 inches. In addition, the presiding official may permit persons to file answers to questions or requests made by the Commission at the hearing for the injury phase, and at any hearing for the remedy phase, within a specified time. In addition, any person who has not entered an appearance as a party to the investigation may submit a written statement of information pertinent to the consideration of injury on or before January 19, 2021, and pertinent to the consideration of remedy on or before March 3, 2021.

Except as provided above, all written submissions must conform with the provisions of section 201.8 of the Commission's rules; any submissions that contain CBI must also conform with the requirements of sections 201.6 and 206.17 of the Commission's rules. The Commission's *Handbook on E-Filing*, available on the Commission's website at <https://edis.usitc.gov>, further explains the Commission's rules with respect to electronic filing.

Any additional written submission to the Commission, including requests pursuant to section 201.12 of the Commission's rules, will not be accepted unless good cause is shown for accepting such a submission, or unless the submission is pursuant to a specific request by a Commissioner or Commission staff.

In accordance with section 201.16(c) of the Commission's rules, each document filed by a party to the investigation must be served on all other parties to the investigation (as identified by the service list), and a certificate of service must be timely filed. The Secretary will not accept a document for filing without a certificate of service.

For further information concerning the conduct of this investigation and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A and B (19 CFR part 201), and part 206, subparts A and B (19 CFR part 206).

Authority: This investigation is being conducted under authority of Section 202 of the Act; this notice is published pursuant to section 203(b)(3) of the Act.

By order of the Commission.

Issued: October 6, 2020.

Lisa Barton,

Secretary to the Commission.

[FR Doc. 2020-22423 Filed 10-8-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms, and Explosives

[OMB Number 1140-0026]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Revision of a Currently Approved Collection; Report of Theft or Loss—Explosive Materials—ATF Form 5400.5

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 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. The proposed information collection (IC) OMB 1140-0026 (Report of Theft or Loss—Explosive Materials—ATF Form 5400.5), is being revised to include separate categories of loss with descriptions and example scenarios, as well as additional clarifications to improve user experience when completing this form. This IC also being published to obtain comments from the public and affected agencies.

DATES: Comments are encouraged and will be accepted for 60 days until December 8, 2020.

FOR FURTHER INFORMATION CONTACT: If you have additional comments, regarding 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: William O'Brien, Explosives Industry Programs Branch, Firearms and Explosives Industry Division either by mail at 99 New York Ave. NE, Washington, DC 20226, by email at eipb-informationcollection@atf.gov, or by telephone at 202-648-7120.

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* (check justification or form 83): Revision of a currently approved collection.

2. *The Title of the Form/Collection:* Report of Theft or Loss—Explosive Materials.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number (if applicable): ATF Form 5400.5.

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.

Other (if applicable): Individuals or households, Not-for-profit institutions, Farms, Federal Government, and State, Local, or Tribal Government.

Abstract: According to 27 CFR 555.30 (a) entitled *Reporting Theft or Loss of Explosive Materials*, "Any licensee or permittee who has knowledge of the theft or loss of any explosive materials from his stock shall, within 24 hours of discovery, report the theft or loss by telephoning 1-800-461-8841 (nationwide toll free number) and on ATF F 5400.5 [Report of Theft or Loss—Explosive Materials], in accordance with the instructions on the form. Theft or loss of any explosive materials shall also be reported to appropriate local authorities."

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 300 respondents will utilize the form annually, and it will take each respondent approximately 1 hour and 48 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 540 hours, which is equal to 300 (# of respondents) * 1 (# of responses per respondents) * 1.8 (1 hour and 48 minutes).

If additional information is required contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, 3E.405A, Washington, DC 20530.

Dated: October 6, 2020.

Melody Braswell,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2020-22453 Filed 10-8-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-720]

Bulk Manufacturer of Controlled Substances Application: Purisys, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Purisys, LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to Supplemental Information listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before December 8, 2020. Such persons may also file a written request for a hearing on the application on or before December 8, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on August 27, 2020, Purisys, LLC, 1550 Olympic Drive, Athens, Georgia 30601-1602, applied to be registered as an bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
5-Methoxy-N-N-dimethyltryptamine.	7431	I
Norlevorphanol	9634	I

The company plans to manufacture the above-listed controlled substances as clinical trial and starting materials to make compounds for distribution to its customers. No other activity for these drug codes is authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020-22442 Filed 10-8-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Steven A. Holper, M.D.; Decision and Order

On October 22, 2019, the Acting Assistant Administrator, Diversion Control Division, Drug Enforcement Administration (hereinafter, Government or DEA), issued an Order to Show Cause (hereinafter, OSC) to Steven A. Holper, M.D., (hereinafter, Registrant), of Las Vegas, Nevada. Government's Request for Final Agency Action Exhibit (hereinafter, RFAAX) 5 (OSC), at 1. The OSC proposed the revocation of Registrant's Certificate of Registration No. BH2498106. It alleged that Registrant is without "authority to handle controlled substances in Nevada, the state in which [Registrant is] registered with the DEA." *Id.* (citing 21 U.S.C. 823(f) and 824(a)(3)).

Specifically, the OSC alleged that Registrant's state controlled substance license expired on October 21, 2018. *Id.* at 1-2. The OSC also alleged that Registrant's state medical license was revoked by the Board of Medical Examiners of the State of Nevada on September 6, 2019. *Id.* at 2. The OSC further alleged that Registrant is not eligible to obtain or retain a DEA registration because he lacks state authority to handle controlled substances in the state of Nevada. *Id.*

The OSC notified Registrant of the right to either request a hearing on the allegations or submit a written statement in lieu of exercising the right to a hearing, the procedures for electing each option, and the consequences for failing to elect either option. *Id.* (citing 21 CFR 1301.43). The OSC also notified Registrant of the opportunity to submit

a corrective action plan. *Id.* at 3 (citing 21 U.S.C. 824(c)(2)(C)).

A DEA Diversion Investigator personally served Registrant with the OSC on December 16, 2019. RFAAX 12, at 2-3 (Declaration of Diversion Investigator One). I find that more than thirty days have now passed since the Government accomplished service of the OSC. Further, based on the Government's written representations, I find that neither Registrant, nor anyone purporting to represent Registrant, requested a hearing, submitted a written statement while waiving Registrant's right to a hearing, or submitted a corrective action plan. RFAAX 11, at 3-4 (Declaration of Diversion Investigator Two). Accordingly, I find that Registrant has waived the right to a hearing and the right to submit a written statement and corrective action plan. 21 CFR 1301.43(d) and 21 U.S.C. 824(c)(2)(C). I, therefore, issue this Decision and Order based on the record submitted by the Government, which constitutes the entire record before me. 21 CFR 1301.46.

Findings of Fact

Registrant's DEA Registration

Registrant is the holder of DEA Certificate of Registration No. BH2498106 at the registered address of 3233 W. Charleston Blvd. 202, Las Vegas, NV 89102. RFAAX 1 (Registrant's DEA Certificate of Registration). Pursuant to this registration, Registrant is authorized to dispense controlled substances in schedules II through V as a practitioner. *Id.* Registrant's registration will expire on its own terms on October 31, 2020. *Id.*

DEA Investigation and the Status of Registrant's State Licenses

On July 22, 2019, Registrant was sentenced in the United States District Court for the District of Nevada on a matter related to his conviction on one count of unlawful distribution of a controlled substance. RFAAX 11, at 2. On August 12, 2019, a DEA Diversion Investigator (hereinafter, DI Two) asked Registrant, through his legal counsel, to voluntarily surrender his DEA registration. *Id.* Registrant declined. *Id.* The General Counsel for the Nevada State Board of Pharmacy (hereinafter, Pharmacy Board) sent DI Two a letter dated September 17, 2019, stating that Registrant did not renew his Nevada controlled substance license and did not hold an active controlled substance license with the Pharmacy Board. RFAAX 4. According to the online records of the Pharmacy Board, Registrant's controlled substance