

proposed information collection instrument with instructions, or additional information, please contact: James Chancey, National Firearms Act Division either by mail at 244 Needy Road, Martinsburg, WV 25405, by email at nfaombcomments@atf.gov, or by telephone at 304-616-4500.

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):* Extension without change of a currently approved collection.

2. *The Title of the Form/Collection:* Notice of Firearms Manufactured or Imported.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number (if applicable): ATF Form 2 (5320.2).

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): Federal Government, and State, Local or Tribal Government.

Abstract: The Notice of Firearms Manufactured or Imported—ATF Form 2 (5320.2) is required of (1) a person who is qualified to manufacture National Firearms Act (NFA) firearms, or (2) a person who is qualified to

import NFA firearms to register manufactured or imported NFA firearm(s).

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 4,212 respondents will utilize the form approximately 3.415 times annually, and it will take each respondent 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 7,192 hours, which is equal to 4,212 (total respondents) * 3.415 (# of responses per respondent) * .5 (30 minutes).

7. *An Explanation of the Change in Estimates:* The adjustments associated with this collection includes a decrease in both the number of respondents and responses for this IC by 340 and 1,161 respectively, since the last renewal in 2017. Due to less respondents, both the hourly and total public cost burden have also reduced by 581 hours and \$ 697, since 2017.

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: March 5, 2020.

Melody Braswell,

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

[FR Doc. 2020-04890 Filed 3-9-20; 8:45 am]

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DEPARTMENT OF JUSTICE

Bureau of Alcohol, Tobacco, Firearms and Explosives

[OMB Number 1140-0097]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension Without Change of a Currently Approved Collection; Supplemental Information on Water Quality Considerations—ATF Form 5000.30

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.

DATES: Comments are encouraged and will be accepted for 60 days until May 11, 2020.

FOR FURTHER INFORMATION CONTACT: If you have additional comments, regarding the estimated public burden, response time, suggestions, or need a copy of the proposed information collection instrument with instructions, or additional information, please contact: Shawn Stevens, Federal Explosives Licensing Center, either by mail at 244 Needy Road, Martinsburg, WV 25405, by email at Shawn.Stevens@atf.gov, or by telephone at 304-616-4400.

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):* Extension without Change of a currently approved collection.

2. *The Title of the Form/Collection:* Supplemental Information on Water Quality Considerations.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* Form number (if applicable): ATF Form 5000.30.

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): None.

Abstract: A person engaged in the business of manufacturing explosives is required to have a license under the provisions of 18 U.S.C. 843. The Federal Water Pollution Control Act, 33 U.S.C. 1341, authorizes the execution of the Supplemental Information on Water Quality Considerations—ATF 5000.30, during the application process, in order to ensure compliance with the Act.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 680 respondents will utilize the form 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 340 hours, which is equal to 680 (# of respondents) *.5 (30 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: March 5, 2020.

Melody Braswell,

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

[FR Doc. 2020-04889 Filed 3-9-20; 8:45 am]

BILLING CODE 4410-18-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA 577]

Importer of Controlled Substances Application: Caligor Pharma Services

ACTION: Notice of application.

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 April 9, 2020. Such persons may also file a written request for a hearing on the application on or before April 9, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/DPW, 8701

Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on September 6, 2019, Caligor Coghlan Pharma Services, 1500 Business Park Drive, Unit B, Bastrop, Texas 78602, applied to be registered as an importer of the following basic class(es) of controlled substance:

Controlled substance	Drug code	Schedule
Tapentadol	9780	II

The company plans to import the listed controlled substance in finished dosage form to be used in pediatric clinical trials. No other activity for this drug code is 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 be extended to the import of FDA approved or non-approved finished dosage forms for commercial use.

Dated: February 11, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020-04835 Filed 3-9-20; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-599]

Importer of Controlled Substances Application: SpecGx LLC

ACTION: Notice of application.

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 April 9, 2020. Such persons may also file a written request for a hearing on the application on or before April 9, 2020.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All request for a hearing should also 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on December 30, 2019, SpecGx LLC, 3600 North Second Street, Saint Louis, Missouri 63147 applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	I
Phenylacetone ..	8501	II
Coca Leaves	9040	II
Thebaine	9333	II
Opium, raw	9600	II
Poppy Straw Concentrate.	9670	II
Tapentadol	9780	II

The company plans to import the listed controlled substances for bulk manufacture into Active Pharmaceutical Ingredients (API) for distribution to its customers. In reference to drug code 7360 (marihuana), the company plans to import synthetic cannabiniol. No other activity for this drug is authorized for this registration. Placement of these codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances.

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 the import of FDA approved or non-approved finished forms for commercial sale.

Dated: February 27, 2020.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2020-04834 Filed 3-9-20; 8:45 am]

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