Controlled substance	Drug code	Schedule
Thebaine	9333	II
Oxymorphone	9652	II
Phenazocine	9715	II
Carfentanil	9743	II
Fentanyl	9801	II

The company plans to bulk manufacture the listed controlled substances for internal use or for sale to its customers. The company plans to manufacture small quantities of the above-listed controlled substances as radiolabeled compounds for biochemical research. No other activities for these drug codes are authorized for this registration.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–10996 Filed 5–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–838]

Importer of Controlled Substances Application: SpecGX, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: SpecGX, LLC. has applied to be registered as an importer 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 June 24, 2021. Such persons may also file a written request for a hearing on the application on or before June 24, 2021.

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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on February 5, 2021, SpecGX LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

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 Tapentadol (9780) and Thebaine (9333), the company plans to import intermediate forms of these controlled substances for further manufacturing prior to distribution to its customers. In reference to drug code 7360 (Marihuana), the company plans to import synthetic cannabinol. 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 Food and Drug Administration-approved or non-approved finished forms for commercial sale.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–10995 Filed 5–24–21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

[OMB Number 1105–0008]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Extension of a Currently Approved Collection; Claim for Damage, Injury, or Death

AGENCY: Civil Division, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Civil Division, 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.

DATES: Comments are encouraged and will be accepted for 60 days until July 26, 2021.

FOR FURTHER INFORMATION CONTACT: Comments are encouraged and all comments should reference the 8 digit OMB number for the collection or the title of the collection. If you have questions concerning the collection, please contact James G. Touhey, Jr., Director, Torts Branch, Civil Division, U.S. Department of Justice, P.O. Box 888, Benjamin Franklin Station, Washington, DC 20044, Telephone: (202) 616–4400.

Written comments and/or suggestions can also be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Room 10235, 725 17th Street NW, Washington, DC 20503 or sent to *OIRA_submissions@omb.eop.gov.*

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

—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 of a currently approved collection.

2. *The Title of the Form/Collection:* Claim for Damage, Injury, or Death.

3. *The agency form number, if any, and the applicable component of the*

Department sponsoring the collection: The form number is CIV SF 95. The applicable component within the Department of Justice is the Civil Division.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Individuals or households. Other: Businesses or other for-profit, Non-for-profit institutions, and State, Local, or Tribal Governments. Abstract: This form is used by those persons making a claim against the United States Government under the Federal Tort Claims Act.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that there will be 100,000 respondents who will each require 6 hours to respond.

6. *An estimate of the total public burden (in hours) associated with the collection:* The total estimated annual burden hours to complete the certification form is 600,000 hours.

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: May 19, 2021.

Melody Braswell,

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

[FR Doc. 2021–10966 Filed 5–24–21; 8:45 am]

BILLING CODE 4410–12–P

DEPARTMENT OF LABOR

Employee Benefits Security Administration

State All Payer Claims Databases Advisory Committee—Notice of Virtual Meeting

AGENCY: Employee Benefits Security Administration (EBSA), Department of Labor (DOL).

ACTION: Notice.

SUMMARY: This notice announces the second meeting of the State All Payer Claims Databases Advisory Committee (hereinafter the Committee). This notice provides information to members of the public who may be interested in attending the meeting or providing written comments related to the work of the Committee. Notice of this meeting is required under the Federal Advisory Committee Act (FACA).

DATES: The second meeting of the State All Payer Claims Databases Advisory Committee will be held virtually on June 11, 2021.

1. *Deadline for Registration without Oral Presentation: June 9, 2021.* Individuals can register for the meeting by visiting the Committee website: *https://www.dol.gov/agencies/ebsa/about-ebsa/about-us/state-all-payer-claims-databases-advisory-committee.*

2. *Deadline for Registration of Oral Presentations: June 7, 2021.* Requests should be submitted by email to *SAPCDAC@dol.gov.*

3. *Deadline for Submission of Oral Remarks and Written Comments: June 7, 2021.* Remarks and comments should be submitted by email to *SAPCDAC@dol.gov.*

4. *Deadline for Requesting Special Accommodations: June 7, 2021.* Requests should be submitted by email to *SAPCDAC@dol.gov.*

ADDRESSES: The meeting will be held via webinar. The webinar link and log-in information will be available at DOL's Committee website: *https://www.dol.gov/agencies/ebsa/about-ebsa/about-us/state-all-payer-claims-databases-advisory-committee.*

FOR FURTHER INFORMATION CONTACT: Elizabeth Schumacher, Designated Federal Officer, EBSA, DOL, by sending an email to *SAPCDAC@dol.gov.* For press inquiries please contact Grant Vaught, Office of Public Affairs, DOL at 202–693–4672.

SUPPLEMENTARY INFORMATION: The Committee is mandated by section 735 of the Employee Retirement Income Security Act of 1974 as added by section 115(b) of the No Surprises Act, enacted as part of the Consolidated Appropriations Act, 2021, div. BB, tit. I, Public Law 116–260 (Dec. 27, 2020). The Committee is governed by the provisions of the FACA, as amended, 5 U.S.C. App. 2.

The Committee will advise the Secretary of Labor on the standardized reporting format for the voluntary reporting by group health plans to State All Payer Claims Databases. Reporting will include medical claims, pharmacy claims, dental claims, and eligibility and provider files collected from private and public payers. The Committee will also advise the Secretary on what guidance is necessary to provide to States on the process by which States may collect such data in the standardized reporting format.

The Committee will be responsible for issuing a report that includes recommendations on the establishment of the format and guidance to the Secretary of Labor and certain congressional committees no later than 180 days after the date of enactment of the Consolidated Appropriations Act, 2021.

The second meeting of the Committee will be held on June 11, 2021 via webinar. The meeting will begin at 9:30 a.m. and end at approximately 5:00 p.m., with a one hour break for lunch. The meeting will focus on the various issues related to all payer claims databases as well as a general discussion of the work plan for the report that must be submitted by the committee. Additional details about the agenda items and topics, as well as agenda updates, will be available at on the Committee's website: *https://www.dol.gov/agencies/ebsa/about-ebsa/about-us/state-all-payer-claims-databases-advisory-committee.*

Dated: May 19, 2021.

Ali Khawar,

Acting Assistant Secretary, Employee Benefits Security Administration, U.S. Department of Labor.

[FR Doc. 2021–10930 Filed 5–24–21; 8:45 am]

BILLING CODE 4510–29–P

NATIONAL CAPITAL PLANNING COMMISSION

Revised Adopted Submission Guidelines

AGENCY: National Capital Planning Commission.

ACTION: Notice of final adoption and effective date.

SUMMARY: At its May 6, 2021 monthly meeting, the National Capital Planning Commission (NCPC or Commission) adopted revised Submission Guidelines related to concept review of Master Plans and the purpose, need, and timing of an Information Presentations. The amended guidelines recommend an early concept review of complex master plans to ensure timely input from the Commission before a master plan advances to draft and final review. The amendments regarding Information Presentations identify the types of projects for which an Information Presentation is advisable and establish the appropriate timing for the presentation. Federal and non-federal agency applicants whose development proposals and plans are subject to statutory mandated Commission plan and project review must submit their proposals to the Commission following a process laid out in the Submission Guidelines.

DATES: The revised Submission Guidelines will become effective June 24, 2021.