

incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on February 25, 2019, Shertech Laboratories, LLC, 1185 Woods Chapel Road, Duncan, South Carolina 29334 applied to be registered as an importer of the following basic class of controlled substance:

| Controlled substance | Drug code | Schedule |
|----------------------|-----------|----------|
| Cocaine .....        | 9041      | II       |

The company plans to import synthetic derivatives of the listed

controlled substance in bulk form to conduct clinical trials.

Approval of permit applications will occur only when the registrant’s activity is consistent with what is authorized under to 21 U.S.C.952 (a)(2).

Authorization will not extend to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: May 17, 2019.

**John J. Martin,**  
Assistant Administrator.

[FR Doc. 2019–11876 Filed 6–5–19; 8:45 am]

**BILLING CODE 4410–09–P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA–392]

**Bulk Manufacturer of Controlled Substances Application: SpecGx LLC**

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before August 5, 2019.

**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 February 1, 2019, SpecGx LLC, 3600 North 2nd Street, Saint Louis, Missouri 63147 applied to be registered as a bulk manufacturer of the following basic class of controlled substances:

| Controlled substance  | Drug code | Schedule |
|---|-----------|----------|
| Gamma Hydroxybutyric Acid .....   | 2010      | I        |
| Tetrahydrocannabinols .....   | 7370      | I        |
| Codeine-N-oxide .....   | 9053      | I        |
| Dihydromorphine .....   | 9145      | I        |
| Difenoxin .....   | 9168      | I        |
| Morphine-N-oxide .....  | 9307      | I        |
| Normorphine .....   | 9313      | I        |
| Norlevorphanol .....  | 9634      | I        |
| Acetyl Fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylacetamide) ..... | 9821      | I        |
| Butyryl Fentanyl .....  | 9822      | I        |
| Fentanyl related-compounds as defined in 21 CFR 1308.11(h) .....        | 9850      | I        |
| Amphetamine .....   | 1100      | II       |
| Methamphetamine .....   | 1105      | II       |
| Lisdexamfetamine .....  | 1205      | II       |
| Methylphenidate .....   | 1724      | II       |
| Nabilone .....  | 7379      | II       |
| 4-Anilino-N-phenethyl-4-piperidine (ANPP) .....                         | 8333      | II       |
| Codeine .....   | 9050      | II       |
| Dihydrocodeine .....  | 9120      | II       |
| Oxycodone .....   | 9143      | II       |
| Hydromorphone .....   | 9150      | II       |
| Diphenoxylate .....   | 9170      | II       |
| Ecgonine .....  | 9180      | II       |
| Hydrocodone .....   | 9193      | II       |
| Levorphanol .....   | 9220      | II       |
| Meperidine .....  | 9230      | II       |
| Methadone .....   | 9250      | II       |
| Methadone intermediate .....  | 9254      | II       |
| Dextropropoxyphene, bulk (non-dosage forms) .....                       | 9273      | II       |
| Morphine .....  | 9300      | II       |
| Oripavine .....   | 9330      | II       |
| Thebaine .....  | 9333      | II       |
| Opium tincture .....  | 9630      | II       |
| Opium, powdered .....   | 9639      | II       |
| Oxymorphone .....   | 9652      | II       |
| Noroxymorphone .....  | 9668      | II       |
| Alfentanil .....  | 9737      | II       |
| Remifentanil .....  | 9739      | II       |
| Sufentanil .....  | 9740      | II       |
| Tapentadol .....  | 9780      | II       |
| Fentanyl .....  | 9801      | II       |

The company plans to manufacture bulk active pharmaceutical ingredients (APIs) for distribution to its customers.

Dated: May 23, 2019.

**John J. Martin,**

*Assistant Administrator.*

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Registration**

**ACTION:** Notice of registration.

**SUMMARY:** The registrant listed below has applied for and been granted a registration by the Drug Enforcement Administration (DEA) as a bulk manufacturer of various classes of schedule I and II controlled substances.

**SUPPLEMENTARY INFORMATION:** The company listed below applied to be registered as a bulk manufacturer of various basic classes of controlled substances. Information on a previously published notice is listed below. No comments or objections were submitted for the notice.

| Company           | FR docket   | Published          |
|-------------------|-------------|--------------------|
| Noramco, Inc .... | 84 FR 5499. | February 21, 2019. |

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of this registrant to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed company.

Dated: May 21, 2019.

**John J. Martin,**

*Assistant Administrator.*

[FR Doc. 2019-11881 Filed 6-5-19; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Notice of Lodging of Proposed Consent Decree Under the Clean Air Act**

On May 22, 2019, the Department of Justice lodged a proposed Consent Decree ("Consent Decree") with the United States District Court for the District of Connecticut in the lawsuit entitled *United States and the State of New Jersey, Department of Environmental Protection v. Gloucester County Utilities Authority*, Civil Action No. 2:19-cv-12818.

In a Complaint, the United States, on behalf of the U.S. Environmental Protection Agency ("EPA") and the State of New Jersey, on behalf of the Department of Environmental Protection, alleges that the Gloucester County Utilities Authority ("GCUA") violated the Clean Air Act (the "Act"), 42 U.S.C. 7413, by violating: (1) The Solid Waste Combustion provisions in Section 129 of the Clean Air Act, 42 U.S.C. 7429, and (2) the Federal Plan Requirements for Sewage Sludge Incineration Units Constructed on or Before October 14, 2010, 40 CFR part 62, subpart LLL ("Subpart LLL"). The proposed Consent Decree in this case, among other things, requires that GCUA pay a civil penalty of \$132,500 in two installments. In addition, the Consent Decree requires a New Jersey-sponsored supplemental project, to be overseen by the state, involving the purchase and installation of four electric vehicle-charging stations within Gloucester County by September 31, 2019.

The publication of this notice opens a period for public comment on the proposed Consent Decree. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States and the State of New Jersey, Department of Environmental Protection v. Gloucester County Utilities Authority*, D.J. Ref. No. 90-5-2-1-11877. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

| To submit comments: | Send them to:   |
|---------------------|---|
| By email .....      | <a href="mailto:pubcomment-ees.enrd@usdoj.gov">pubcomment-ees.enrd@usdoj.gov</a>              |
| By mail .....       | Assistant Attorney General,<br>U.S. DOJ—ENRD, P.O.<br>Box 7611, Washington, DC<br>20044-7611. |

During the public comment period, the proposed Consent Decree may be

examined and downloaded at this Justice Department website: <https://www.justice.gov/enrd/consent-decrees>. We will provide a paper copy of the proposed Consent Decree upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044-7611.

Please enclose a check or money order for \$10.00 (25 cents per page reproduction cost), payable to the United States Treasury.

**Jeffrey Sands,**

*Assistant Chief, Environmental Enforcement Section, Environment & Natural Resources Division.*

[FR Doc. 2019-11863 Filed 6-5-19; 8:45 am]

**BILLING CODE 4410-15-P**

**DEPARTMENT OF LABOR**

**Employee Benefits Security Administration**

**196th Meeting of the Advisory Council on Employee Welfare and Pension Benefit Plans; Notice of Meeting**

Pursuant to the authority contained in Section 512 of the Employee Retirement Income Security Act of 1974 (ERISA), 29 U.S.C. 1142, the 196<sup>th</sup> meeting of the Advisory Council on Employee Welfare and Pension Benefit Plans (also known as the ERISA Advisory Council) will be held on June 25-27, 2019.

The three-day meeting will take place at the U.S. Department of Labor, 200 Constitution Avenue NW, Washington, DC 20210 in N5437 A-C. The meeting will run from 9:00 a.m. to approximately 5:30 p.m. on June 25 and 26 with a one hour break for lunch, and from 8:30 a.m. to 11:00 a.m. on June 27. The purpose of the open meeting is for Advisory Council members to hear testimony from invited witnesses and to receive an update from the Employee Benefits Security Administration (EBSA). The EBSA update is scheduled for the morning of June 25, subject to change.

The Advisory Council will study the following topics: (1) Beyond Plan Audit Compliance: Improving the Financial Statement Audit Process (on June 25); and, (2) Permissive Transfers of Uncashed Checks from ERISA Plans to State Unclaimed Property Funds (on June 26). The Advisory Council will continue with discussions of its topics on June 27. Descriptions of these topics are available on the Advisory Council page of the EBSA website, at <https://www.dol.gov/agencies/ebsa/about-ebsa/about-us/erisa-advisory-council>.