

**DEPARTMENT OF JUSTICE**

**Antitrust Division**

**Notice Pursuant to the National Cooperative Research and Production Act of 1993—Electrified Vehicle and Energy Storage Evaluation**

Notice is hereby given that, on September 21, 2021, pursuant to Section 6(a) of the National Cooperative Research and Production Act of 1993, 15 U.S.C. 4301 *et seq.* (“the Act”), Electrified Vehicle and Energy Storage Evaluation (“EVESE”) has filed written notifications simultaneously with the Attorney General and the Federal Trade Commission disclosing changes in its membership. The notifications were filed for the purpose of extending the Act’s provisions limiting the recovery of antitrust plaintiffs to actual damages under specified circumstances. Specifically, LG Energy Solution, Ltd., Seoul, KOREA, has been added as a party to this venture.

No other changes have been made in either the membership or planned activity of the group research project. Membership in this group research project remains open, and EVESE intends to file additional written notifications disclosing all changes in membership.

On September 24, 2020, EVESE filed its original notification pursuant to Section 6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on October 15, 2020 (85 FR 65423).

The last notification was filed with the Department on July 27, 2021. A notice was published in the **Federal Register** pursuant to Section 6(b) of the Act on August 23, 2021 (86 FR 47157).

**Suzanne Morris,**

*Chief, Premerger and Division Statistics, Antitrust Division.*

[FR Doc. 2021-23098 Filed 10-21-21; 8:45 am]

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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-913]

**Bulk Manufacturer of Controlled Substances Application: Chattem Chemicals**

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

**ACTION:** Notice of application.

**SUMMARY:** Chattem Chemicals has applied to be registered as a bulk

manufacturer 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 file written comments on or objections to the issuance of the proposed registration on or before December 21, 2021. Such persons may also file a written request for a hearing on the application on or before December 21, 2021.

**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 July 23, 2021, Chattem Chemicals, 3801 Saint Elmo Avenue, Chattanooga, Tennessee 37409-1237, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance            | Drug code | Schedule |
|---------------------------------|-----------|----------|
| Gamma Hydroxybutyric Acid ..... | 2010      | I        |
| Marihuana .....                 | 7360      | I        |
| Tetrahydrocannabinols .....     | 7370      | I        |
| 4-Methoxyamphetamine .....      | 7411      | I        |
| Noroxymorphone .....            | 9145      | I        |
| Amphetamine .....               | 1100      | II       |
| Methamphetamine .....           | 1105      | II       |
| Lisdexamfetamine .....          | 1205      | II       |
| Methylphenidate .....           | 1724      | II       |
| Cocaine .....                   | 9041      | II       |
| Codeine .....                   | 9050      | II       |
| Dihydrocodeine .....            | 9120      | II       |
| Oxycodone .....                 | 9143      | II       |
| Hydromorphone .....             | 9150      | II       |
| Ecgonine .....                  | 9180      | II       |
| Hydrocodone .....               | 9193      | II       |
| Levorphanol .....               | 9220      | II       |
| Methadone .....                 | 9250      | II       |
| Methadone intermediate .....    | 9254      | II       |
| Morphine .....                  | 9300      | II       |
| Oripavine .....                 | 9330      | II       |
| Thebaine .....                  | 9333      | 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 the listed controlled substances in bulk for distribution and sale to its customers. In reference to drug code 7360 (Marihuana) and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as a synthetic.

No other activities for this drug code are authorized for this registration.

**Brian S. Besser,**

*Acting Assistant Administrator.*

[FR Doc. 2021-23037 Filed 10-21-21; 8:45 am]

**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-919]

**Importer of Controlled Substances Application: Mycrodose Therapeutics**

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

**ACTION:** Notice of application.

**SUMMARY:** Mycrodose Therapeutics 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 file written comments on or objections to the issuance of the proposed registration on or before November 22, 2021. Such persons may also file a written request for a hearing on the application on or before November 22, 2021.

**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 September 17, 2021, Mycrodose Therapeutics, 5940 Pacific Mesa Court, Suite 210, San Diego, California 92121-4317 applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|----------------------|-----------|----------|
| Psilocybin .....     | 7437      | I        |
| Psilocyn .....       | 7438      | I        |