

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 TOG intends to file additional written notifications disclosing all changes in membership.

On April 21, 1997, TOG 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 June 13, 1997 (62 FR 32371).

The last notification was filed with the Department on September 9, 2015. A notice was published in the **Federal Register** pursuant to section 6(b) of the Act on October 2, 2015 (80 FR 59816).

Patricia A. Brink,

Director of Civil Enforcement, Antitrust Division.

[FR Doc. 2016-00325 Filed 1-8-16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Application: Sharp Clinical Services,
Inc.**

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before February 10, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before February 10, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, 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 Deputy Assistant Administrator of the DEA Office of Diversion Control ("Deputy 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 July 29, 2015, Sharp Clinical Services, Inc., 300 Kimberton Road, Phoenixville, Pennsylvania 19460 applied to be registered as an importer of marihuana (7360), a basic class of controlled substance listed in schedule I.

The company plans to import finished pharmaceutical products containing cannabis extracts in dosage form for clinical trial studies.

This compound is listed under drug code 7360. No other activity for this drug code is authorized for this registration. Approval of permits 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 to the import of FDA approved or non-approved finished dosage forms for commercial sale.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2016-00214 Filed 1-8-16; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

**Importer of Controlled Substances
Application: Myoderm**

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 in accordance with 21 CFR 1301.34(a) on or before February 10, 2016. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before February 10, 2016.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrisette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing

Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, 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 Deputy Assistant Administrator of the DEA Office of Diversion Control ("Deputy 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 October 9, 2015, Myoderm, 48 East Main Street, Norristown, Pennsylvania 19401 applied to be registered as an importer of the following basic classes of controlled substances:

| Controlled substance | Schedule |
|-----------------------------------|----------|
| Amphetamine (1100) | II |
| Lisdexamfetamine (1205) | II |
| Methylphenidate (1724) | II |
| Pentobarbital (2270) | II |
| Nabilone (7379) | II |
| Codeine (9050) | II |
| Oxycodone (9143) | II |
| Hydromorphone (9150) | II |
| Hydrocodone (9193) | II |
| Levomethorphan (9210) | II |
| Meperidine (9230) | II |
| Methadone (9250) | II |
| Methadone intermediate (9254) ... | II |
| Morphine (9300) | II |
| Oxymorphone (9652) | II |
| Fentanyl (9801) | II |

The company plans to import the listed controlled substances in finished dosage form for clinical trials, research, and analytical purposes.

The import of the above listed basic classes of controlled substances will be granted only for analytical testing, research, and clinical trials. This authorization does not extend to the import of a finished FDA approved or non-approved dosage form for commercial sale.

Dated: January 4, 2016.

Louis J. Milione,

Deputy Assistant Administrator.

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