

**SOUTH DAKOTA**

**Miner County**

Wheeler Hotel, 101 N. Main St.,

Additional documentation has been received for the following resource:

**MINNESOTA**

**Ramsey County**

Church of St. Casimir—Catholic, 937 E. Jessamine Ave., St. Paul, AD83000939

**Authority:** 60.13 of 36 CFR part 60

**Dated:** December 13, 2017.

**J. Paul Loether,**

*Chief, National Register of Historic Places/ National Historic Landmarks Program and Keeper, National Register of Historic Places.*

[FR Doc. 2017-28467 Filed 1-3-18; 8:45 am]

**BILLING CODE 4312-52-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Bulk Manufacturer of Controlled Substances Application: Organix, Inc.**

**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 March 5, 2018.

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

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his 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 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 1301.33(a), this is notice that on September 6, 2017, Organix, Inc., 240 Salem Street, Woburn, Massachusetts 01801 applied to be registered as a bulk

manufacturer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid.	2010	I
Lysergic acid diethylamide.	7315	I
Marihuana .....	7360	I
Tetrahydrocannabinols ...	7370	I
Dimethyltryptamine .....	7435	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I
Heroin .....	9200	I
Morphine .....	9300	II

The company plans to manufacture reference standards for distribution to its research and forensic customers. In reference to drug code 7360 (marihuana) and 7370 (THC) the company plans to manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

**Dated:** December 20, 2017.

**Demetra Ashley,**

*Acting Assistant Administrator.*

DC: \_\_\_\_\_

DCX: \_\_\_\_\_

**Return to ODW—Mike Lewis**

DCX: \_\_\_\_\_

DR: \_\_\_\_\_

DRX: \_\_\_\_\_

DRW: \_\_\_\_\_

DRW Policy Analyst: \_\_\_\_\_

DRG: \_\_\_\_\_

DRG/A: \_\_\_\_\_

DRGR Unit Chief: \_\_\_\_\_

DRQ: \_\_\_\_\_

DRGR Staff Coordinator \_\_\_\_\_

DRGR—L.Mckoy 10/25/2017

NOA—ORGANIX INC

Document#: DRGR-17-0306

DFN: 010.02.A1 General

Correspondence

DFN: 301-04 **Federal Register** Files

[FR Doc. 2017-28180 Filed 1-3-18; 8:45 am]

**BILLING CODE 4410-09-P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-392]

**Importer of Controlled Substances Application: Sharp (Bethlehem), 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 February 5, 2018. Such persons may also file a written request for a hearing on the application pursuant on or before February 5, 2018.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA **Federal Register** Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** The Attorney General has delegated his 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 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 June 15, 2017, Sharp (Bethlehem), LLC, 2400 Baglyos Circle, Bethlehem, Pennsylvania 18020 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
3,4-Methylenedioxy methamphetamine.	7405	I
Psilocybin .....	7437	I

The company plans to import the listed controlled substances for clinical trials. No other activity for these drug codes 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-