comments on or objections to the issuance of the proposed registration on or before April 9, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette 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 July 21, 2017, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Chattanooga, Tennessee 37409 applied to be registered as a bulk manufacturer for the basic classes of controlled substances:

Controlled substance	Drug code	Schedul
Gamma Hydroxybutyric Acid.	2010	1
Marihuana	7360	1
Tetrahydrocannabinols	7370	1
4-Methoxyamphetamine	7411	1
Dihydromorphine	9145	1
Amphetamine	1100	П
Methamphetamine	1105	II
Lisdexamfetamine	1205	П
Methylphenidate	1724	II
Pentobarbital	2270	П
Codeine	9050	II
Dihydrocodeine	9120	II
Oxycodone	9143	II
Hydromorphone	9150	II
Hydrocodone	9193	П
Meperidine	9230	II
Meperidine intermediate- A.	9232	II
Meperidine intermediate- B.	9233	II
Meperidine intermediate- C.	9234	II
Methadone	9250	П
Methadone intermediate	9254	П
Morphine	9300	П
Oripavine	9330	П
Thebaine	9333	П
Opium tincture	9630	П
Opium, powdered	9639	П
Opium, granulated	9640	II
Oxymorphone	9652	II

Controlled substance	Drug code	Schedule
Noroxymorphone Alfentanil Remifentanil Sufentanil Tapentadol Fentanyl	9668 9737 9739 9740 9780 9801	= = = =

The company plans to manufacture the listed controlled substances in bulk for distribution and sale to its customers.

In reference to drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

Dated: January 30, 2018.

## Susan A. Gibson,

Deputy Assistant Administrator. [FR Doc. 2018–02343 Filed 2–5–18; 8:45 am]

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## **DEPARTMENT OF JUSTICE**

# **Drug Enforcement Administration**

[Docket No. DEA-392]

## Bulk Manufacturer of Controlled Substances Application: Cedarburg Pharmaceuticals

**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 April 9, 2018.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrissette 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 August 29, 2017, Cedarburg Pharmaceuticals, Inc., A Division of Albany Molecular Research Inc. (AMRI), 870 Badger Circle, Grafton, Wisconsin 53024 applied to be registered as a bulk manufacturer the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360 7370 1205 2270 7379 8333 9739 9801	  -  -  -  -  -  -

The company plans to manufacture the above listed controlled substances in bulk for distribution to its customers. In reference to drug codes 7360 marihuana, the company plans to bulk manufacture cannabidiol as a synthetic intermediate. The company plans to manufacture bulk active pharmaceutical ingredients (API) for distribution to its customers. This controlled substance will be further synthesized to bulk manufacture a synthetic tetrahydrocannabinols 7370. No other activity for this drug code is authorized for this registration.

Dated: January 30, 2018.

#### Susan A. Gibson,

 $\label{eq:continuity} Deputy Assistant Administrator. \\ [FR Doc. 2018–02341 Filed 2–5–18; 8:45 am]$ 

BILLING CODE 4410-09-P

# **DEPARTMENT OF JUSTICE**

## Notice of Proposed Settlement Agreement and Draft Restoration Plan Under The Oil Pollution Act of 1990, and The Clean Water Act

Notice is hereby given that the United States of America, on behalf of the Department of the Interior ("DOI") acting through the Fish and Wildlife Service and the National Park Service, the District of Columbia, on behalf of the Department of Energy and Environment, and the Commonwealth of Virginia, acting through the Virginia Department of Environmental Quality (collectively "Trustees"), are providing an opportunity for public comment on a proposed Settlement Agreement ("Settlement Agreement") among the DOI, the District of Columbia, the Commonwealth of Virginia, and