

Allen, N.P., for additional registration in Indiana. This Order is effective September 24, 2026.

Signing Authority

This document of the Drug Enforcement Administration was signed on August 18, 2026, by DEA Administrator Terrance C. Cole. That document with the original signature and date is maintained by DEA. For administrative purposes only, and in compliance with requirements of the Office of the Federal Register, the undersigned DEA Federal Register Liaison Officer has been authorized to sign and submit the document in electronic format for publication, as an official document of DEA. This administrative process in no way alters the legal effect of this document upon publication in the **Federal Register**.

Heather Achbach,

Federal Register Liaison Officer, Drug Enforcement Administration.

[FR Doc. 2026-17275 Filed 8-24-26; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1750]

Bulk Manufacturer of Controlled Substances Application: Biopharmaceutical Research Company

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Biopharmaceutical Research Company 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 submit electronic comments on or objections to the issuance of the proposed registration on or before October 26, 2026. Such persons may also file a written request for a hearing on the application on or before October 26, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission

of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 14, 2026, Biopharmaceutical Research Company, 11045 Commercial Parkway, Castroville, California 95012-3209, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols ...	7370	I

The company plans to bulk manufacture the listed controlled substances to provide Pharmaceutical-grade marihuana in order to facilitate research in a manner that complies with local, state, and federal regulations. No other activities for these drug codes are authorized for this registration.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026-17273 Filed 8-24-26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. 1753]

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 submit electronic comments on or objections to the issuance of the proposed registration on or before October 26, 2026. Such persons may also file a written request for a hearing on the application on or before October 26, 2026.

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SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 20, 2026, 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
Dihydromorphine	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 bulk manufacture the listed controlled substances in bulk 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.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026–17287 Filed 8–24–26; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1754]

Bulk Manufacturer of Controlled Substances Application: Kinetochem LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Kinetochem LLC 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 submit electronic comments on or objections to the issuance of the proposed registration on or before October 26, 2026. Such persons may also file a written request for a hearing on the application on or before October 26, 2026.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 7, 2026, Kinetochem LLC, 96 Market Street, Suite 102, Georgetown, Texas 78626–3618, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
3,4-Methylenedioxy-N-methylcathinone.	7540	I

The company plans to bulk manufacture the listed controlled substance as Active Pharmaceutical Ingredients to its customers as well as for research and clinical trials. No other activity for this drug code is authorized for this registration.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026–17288 Filed 8–24–26; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1755]

Bulk Manufacturer of Controlled Substances Application: Curia Missouri Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Curia Missouri Inc. 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 submit electronic comments on or objections to the issuance of the proposed registration on or before October 26, 2026. Such persons may also file a written request for a hearing on the application on or before October 26, 2026.

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SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on July 14, 2026, Curia Missouri Inc., 2460 West Bennett Street, Springfield, Missouri 65807–1229, 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
Amphetamine	1100	II
Lisdexamfetamine	1205	II
Methylphenidate	1724	II
Phenylacetone	8501	II
Tapentadol	9780	II

The company plans to bulk manufacture the listed controlled substances for internal use intermediates or for sale to its customers. No other activities for these drug codes are authorized for this registration.

Justin Wood,

Acting Deputy Assistant Administrator.

[FR Doc. 2026–17280 Filed 8–24–26; 8:45 am]

BILLING CODE:P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–1749]

Importer of Controlled Substances Application: Vici Health Sciences, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Vici Health Sciences, LLC 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 submit electronic comments on or objections to the issuance of the proposed registration on or before September 24, 2026. Such persons may also file a written request for a hearing on the application on or before September 24, 2026.

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