The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated each of the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR

1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed companies.

Dated: December 21, 2018.

#### John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01501 Filed 2–6–19; 8:45 am]

BILLING CODE 4410-09-P

#### **DEPARTMENT OF JUSTICE**

# Drug Enforcement Administration

[Docket No. DEA-392]

# Bulk Manufacturer of Controlled Substances Registration

**ACTION:** Notice of registration.

**SUMMARY:** The registrant listed below has applied for and been granted registration by the Drug Enforcement Administration (DEA) as a bulk manufacturer of various classes of schedule II controlled substances.

**SUPPLEMENTARY INFORMATION:** The company listed below applied to be registered as a bulk manufacturer of various basic classes of controlled substances. Information on a previously published notice is listed below. No comments or objections were submitted for the notice.

Company	FR Docket	Published
Janssen Pharmaceuticals, Inc	83 FR 55205	November 2, 2018.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of this registrant to manufacture the applicable basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed company.

Dated: January 7, 2019.

#### John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01502 Filed 2–6–19; 8:45 am]

BILLING CODE 4410-09-P

#### **DEPARTMENT OF JUSTICE**

## **Drug Enforcement Administration**

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Kinetochem, 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 April 8, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 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 17, 2018, Kinetochem, LLC., 111 W Cooperative Way, Ste. 310–B, Georgetown, Texas 78626 applied to be registered as a bulk manufacturer for the basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana	7360	1

Controlled substance	Drug code	Schedule
Tetrahydrocannabino- ls.	7370	I

The company plans to manufacture drug codes 7360 (marihuana) and 7370 (tetrahydrocannabinols), in bulk for distribution and sale to its customers.

The company plans to synthetically manufacture these drugs. No other activities for these drug codes are authorized for this registration.

Dated: December 21, 2018.

#### John J. Martin,

Assistant Administrator.

[FR Doc. 2019–01509 Filed 2–6–19; 8:45 am]

BILLING CODE 4410-09-P

#### **DEPARTMENT OF JUSTICE**

# **Drug Enforcement Administration**

[Docket No. DEA-392]

Bulk Manufacturer of Controlled Substances Application: Johnson Matthey Pharmaceutical Materials, 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 April 8, 2019.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia

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 November 8, 2018, Johnson Matthey Pharmaceutical Materials, Inc., 25 Patton Road, Devens, Massachusetts 01434 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substances	Drug code	Schedule
Amphetamine Methylphenidate Nabilone Hydrocodone Levorphanol Alfentanil Remifentanil	1100 1724 7379 9193 9220 9737 9739 9740	             

The company plans to support its other manufacturing facilities located in West Deptford, New Jersey and Conshohocken, Pennsylvania with manufacturing and analytical testing.

Dated: December 21, 2018.

# John J. Martin,

Assistant Administrator.

[FR Doc. 2019-01507 Filed 2-6-19: 8:45 am]

BILLING CODE 4410-09-P

#### **DEPARTMENT OF JUSTICE**

[OMB Number 1121-0302]

**Agency Information Collection** Activities; Proposed eCollection eComments Requested; Reinstatement, Without Change, of a **Previously Approved Collection for** Which Approval Has Expired: 2019 Supplemental Victimization Survey (SVS) to the National Crime Victimization Survey (NCVS)

**AGENCY:** Bureau of Justice Statistics, Department of Justice.

**ACTION:** 60-Day notice.

**SUMMARY:** The Department of Justice (DOJ), Office of Justice Programs, Bureau of Justice Statistics, will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. DATES: Comments are encouraged and will be accepted for 60 days until April 8, 2019.

FOR FURTHER INFORMATION CONTACT: If

vou have additional comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact Jennifer Truman or Rachel Morgan, Statistician, Bureau of Justice Statistics, 810 Seventh Street NW, Washington, DC 20531 (email: Jennifer.Truman@ usdoj.gov; telephone: 202-514-5083; email: Rachel.Morgan@usdoj.gov; telephone: 202-616-1707).

**SUPPLEMENTARY INFORMATION: Written** comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

- -Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the Bureau of Justice Statistics, including whether the information will have practical utility;
- -Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
- -Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced: and
- -Minimize the burden of the collection of information on those who are to respond, including through the use of

appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

## **Overview of This Information** Collection

(1) Type of Information Collection: Reinstatement of the Supplemental Victimization Survey (SVS), without changes, a previously approved collection for which approval has expired.

(2) The Title of the Form/Collection: 2019 Supplemental Victimization Survey (SVS) to the National Crime Victimization Survey (NCVS).

(3) The agency form number, if any, and the applicable component of the Department sponsoring the collection: The form number for the questionnaire is SVS-1. The applicable component within the Department of Justice is the Bureau of Justice Statistics, in the Office

of Justice Programs.

- (4) Affected public who will be asked or required to respond, as well as a brief abstract: Respondents will be persons 16 years or older living in households located throughout the United States sampled for the National Crime Victimization Survey (NCVS). The SVS will be conducted as a supplement to the NCVS in all sample households for a six (6) month period from July through December 2019. The SVS is primarily an effort to measure the prevalence of stalking victimization among persons, the types of stalking victimization experienced, the characteristics of stalking victims, the nature and consequences of stalking victimization, and patterns of reporting to the police. BJS plans to publish this information in reports and reference it when responding to queries from the U.S. Congress, Executive Office of the President, the U.S. Supreme Court, state officials, international organizations, researchers, students, the media, and others interested in criminal justices
- (5) An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond: An estimate of the total number of respondents is 119,526 persons age 16 or older. About 98.6% (117,879) will have no stalking victimization and will complete the short interview with an average burden of three (3) minutes. Among the 1.4% of respondents (1,647) who experience stalking victimization, the time to ask the detailed questions regarding the aspects of their stalking victimization is estimated to take an average of 18