

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 September 30, 2015. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before September 30, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODXL, 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. Comments and request for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

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 June 3, 2015, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616–3466 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
4-Anilino-N-phenethyl-4-piperidine (8333).	II
Phenylacetone (8501)	II
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II

The company plans to import the listed controlled substances for internal use, and to manufacture bulk intermediates for sale to its customers.

Dated: August 21, 2015.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
 [FR Doc. 2015–21557 Filed 8–28–15; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Manufacturer of Controlled Substances Registration: Johnson Matthey Pharmaceutical Materials, Inc.

ACTION: Notice of registration.

SUMMARY: Johnson Matthey Pharmaceutical Materials, Inc. applied to be registered as a manufacturer of certain basic classes of controlled substances. The Drug Enforcement Administration (DEA) grants Johnson Matthey Pharmaceutical Materials, Inc. registration as a manufacturer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated April 14, 2015, and published in the **Federal Register** on April 22, 2015, 80 FR 22559, Johnson Matthey Pharmaceutical Materials, Inc., 25 Patton Road, Devens, Massachusetts 01434 applied to be registered as a manufacturer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Johnson Matthey Pharmaceutical Materials, Inc. to manufacture the 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 above-named company is granted registration as a bulk manufacturer of the basic classes of controlled substances:

Controlled substance	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Hydrocodone (9193)	II
Alfentanil (9737)	II

Controlled substance	Schedule
Remifentanil (9739)	II
Sufentanil (9740)	II

The company plans to utilize this facility to manufacture small quantities of the listed controlled substances in bulk and to conduct analytical testing in support of the company’s primary manufacturing facility in West Deptford, New Jersey. The controlled substances manufactured in bulk at this facility will be distributed to its company’s customers.

Dated: August 21, 2015.
Joseph T. Rannazzisi,
Deputy Assistant Administrator.
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Rhodes Technologies

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 1301.33(a) on or before October 30, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODXL, 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 1301.33(a), this is notice that on July 8, 2015, Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Tetrahydrocannabinols (7370)	I
Dihydromorphine (9145)	I
Methylphenidate (1724)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Tapentadol (9780)	II
Fentanyl (9801)	II

The company plans to manufacture the listed controlled substances in bulk for conversion and sale to dosage form manufacturers.

In reference to drug code 7370 the company plans to bulk manufacture a synthetic tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

Dated: August 21, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

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DEPARTMENT OF JUSTICE

[OMB Number 1121-NEW]

Agency Information Collection Activities; Proposed eCollection Activities Requested; New Collection: Body Worn Camera Supplement (BWCS) to the Law Enforcement Management and Administrative Statistics (LEMAS) Survey

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 October 30, 2015.

FOR FURTHER INFORMATION CONTACT: If you 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 Alexia Cooper, Statistician, Law Enforcement Statistics, Bureau of Justice Statistics, 810 Seventh Street NW., Washington, DC 20531 (email: *Alexia.Cooper@usdoj.gov*; telephone: 202-307-0582).

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:* New collection.

(2) *The Title of the Form/Collection:* Body Worn Camera Supplement (BWCS) to the Law Enforcement Management and Administrative Statistics (LEMAS) Survey

(3) *The agency form number, if any, and the applicable component of the Department sponsoring the collection:* No agency form number at this time. 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 general purpose state and local law enforcement agencies (LEAs), including police departments, sheriff’s offices, and state law enforcement agencies. *Abstract:* Since 1987, BJS has collected information about the personnel, policies, and practices of law enforcement agencies via the Law Enforcement Management and Administrative Statistics (LEMAS) survey. This core survey, which has been administered every 4 to 6 years, has been used to produce nationally representative estimates of the functions and responsibilities of law enforcement agencies and the staff serving in those organizations. In addition to core management and administrative information, BJS will also begin using the LEMAS platform for topical supplemental surveys, fielded periodically, to collect data on key issues in contemporary policing. The body worn camera supplement (BWC) is the first of these topical supplements. Specifically, the BWCS survey will focus on LEAs use of body-worn media and will ask agencies about their experiences with body-worn cameras, factors that influence the choice to acquire the technology, and considerations that guide policies for the use of these technologies. This survey will build on the existing LEMAS program and provide key information on an issue that is of particular interest to the law enforcement community and the communities they serve.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An agency-level survey will be sent to approximately 3,336 LEA respondents. The expected burden placed on these respondents is about 23 minutes per respondent.

(6) *An estimate of the total public burden (in hours) associated with the collection:* The total respondent burden is approximately 1,278.8 burden hours.

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE., 3E.405B, Washington, DC 20530.

Dated: August 25, 2015.

Jerri Murray,

Department Clearance Officer for PRA, U.S. Department of Justice.

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