DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Patheon Pharmaceuticals, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class and applicants therefore may file written comments or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before May 26, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 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 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 December 11, 2014, Patheon Pharmaceuticals, Inc., 2110 E. Galbraith Road, Cincinnati, Ohio 45237 applied to be registered as a bulk manufacturer of gamma hydroxybutyric acid (2010), a basic class of nonnarcotic controlled substance in schedule I.

The company plans to manufacture the listed controlled substance for distribution to its customers.

Dated: March 9, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2015–06733 Filed 3–23–15; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Penick Corporation

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 May 26, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODW, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 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 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 February 26, 2014, Penick Corporation, 33 Industrial Park Road, Pennsville, New Jersey 08070, applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Opium tincture (9630)	II
Oxymorphone (9652)	II

The company plans to manufacture the above-listed controlled substances as bulk controlled substance intermediates for distribution to its customers.

Dated: March 9, 2015.

Joseph T. Rannazzisi,

Deputy Assistant Administrator.

[FR Doc. 2015–06734 Filed 3–23–15; 8:45 am]

BILLING CODE 4410–09–P

LIBRARY OF CONGRESS

Copyright Royalty Board

[Docket No. 14–CRB–0006 DART SR (CO/FA) 2013]

Distribution of the 2013 Digital Audio Recording Technology Royalty Funds

AGENCY: Copyright Royalty Board, Library of Congress.

ACTION: Notice announcing commencement of proceeding with request for Petitions to Participate and comments on intention to conduct paper proceeding.

SUMMARY: The Copyright Royalty Board is announcing the commencement of a proceeding to determine the distribution of the digital audio recording technology royalty fees in the 2013 Sound Recordings Fund (Copyright Owners and Featured Recording Artists Subfunds). The Board is also announcing the date by which a party who wishes to participate in this proceeding must file its Petition to Participate and the accompanying $150 filing fee, if applicable. Finally, the Board is announcing the Copyright Royalty Judges' intention to conduct a paper proceeding.

DATES: Petitions to Participate, comments on the intention to conduct a paper proceeding, and applicable filing fee are due no later than April 23, 2015.

ADDRESSES: An original, five copies, and an electronic copy in Portable Document Format (PDF) on a CD of the Petition to Participate, along with the $150 filing fee, if applicable, may be delivered to the Copyright Royalty Board by either mail or hand delivery. Petitions to Participate and the $150 filing fee may not be delivered by an overnight delivery service other than the U.S. Postal Service Express Mail. If by mail (including overnight delivery),