

written submissions, except for confidential business information, will be made available for inspection by interested parties.

The Commission may include some or all of the confidential business information submitted in the course of this investigation in the report it sends to the USTR. Additionally, all information, including confidential business information, submitted in this investigation may be disclosed to and used: (i) By the Commission, its employees and Offices, and contract personnel (a) for developing or maintaining the records of this or a related proceeding, or (b) in internal investigations, audits, reviews, and evaluations relating to the programs, personnel, and operations of the Commission including under 5 U.S.C. Appendix 3; or (ii) by U.S. government employees and contract personnel (a) for cybersecurity purposes or (b) in monitoring user activity on U.S. government classified networks. The Commission will not otherwise disclose any confidential business information in a manner that would reveal the operations of the firm supplying the information.

Summaries of Written Submissions: The Commission intends to publish summaries of the positions of interested persons. Persons wishing to have a summary of their position included in the report should include a summary with their written submission. The summary may not exceed 500 words, should be in MSWord format or a format that can be easily converted to MSWord, and should not include any confidential business information. The summary will be published as provided if it meets these requirements and is germane to the subject matter of the investigation. The Commission will identify the name of the organization furnishing the summary and will include a link to the Commission's Electronic Document Information System (EDIS) where the full written submission can be found.

By order of the Commission.
 Issued: January 17, 2017.

Lisa R. Barton,
Secretary to the Commission.
 [FR Doc. 2017-01401 Filed 1-19-17; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-392]

Importer of Controlled Substances Application: Mylan Technologies, 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 in accordance with 21 CFR 1301.34(a) on or before February 22, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before February 22, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 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 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 1301.34(a), this is notice that on October 31, 2016, Mylan Technologies, Inc., 110 Lake Street, Saint Albans, Vermont 05478 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Methylphenidate	1724	II
Fentanyl	9801	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

Dated: October 22, 2016.

Louis J. Milione,
Assistant Administrator.
 [FR Doc. 2017-01305 Filed 1-19-17; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Consent Decree Under the Resource Conservation and Recovery Act

On January 12, 2017, the Department of Justice and the State of Louisiana on behalf of the Louisiana Department of Environmental Quality ("LDEQ") filed a Complaint and lodged a proposed Consent Decree with the United States District Court for the Middle District of Louisiana in the matter of *United States of America and Louisiana Department of Environmental Quality vs. Innophos, Inc.*, Civil Action No. 17-26-SDD-RLB (M.D. La.).

In the Complaint filed in this action, the United States and LDEQ sought injunctive relief and civil penalties against Innophos, Inc. ("Innophos") for violations of the Resource Conservation and Recovery Act ("RCRA"), 42 U.S.C. 6901-6992k, at Innophos's purified phosphoric acid manufacturing facility near Geismar, Louisiana. The Complaint alleged that Innophos routinely generated two hazardous wastes, Raffinate and RP Pondwater, and sent them to an adjacent facility for disposal; the receiving facility was not authorized to dispose of hazardous waste. LDEQ is a co-plaintiff and has brought its own claims under state law.

The proposed Consent Decree memorializes that Innophos has already corrected the violations related to RP Pondwater. Innophos also agrees in the Consent Decree to handle Raffinate appropriately, either by disposing of it in a permitted hazardous waste Underground Injection Control well system, by treating it on-site, or by shipping it to a permitted hazardous waste treatment, storage, and disposal