DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-629]

Importer of Controlled Substances Application: Mylan Pharmaceuticals Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturer of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER]. Such persons may also file a written request for a hearing on the application on or before [INSERT 30 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on March 31, 2020, Mylan Pharmaceuticals Inc., 2898 Manufacturers Road, Greensboro, North Carolina, 27406, applied to be registered as an importer of the following basic class(es) of controlled substances:

Controlled Substance	Drug Code	Schedule
Remifentanil	9739	II

The company plans to import the above-controlled substance as the FDA-approved drug product in finished dosage form for commercial distribution to its customers. Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under **21 U.S.C. 952**(a)(2).

William T. McDermott,
Assistant Administrator.

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