DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-602]

Bulk Manufacturer of Controlled Substances Application: Navinta LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers 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 60 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on September 23, 2019, Navinta LLC, 1499 Lower Ferry Road, Ewing, New Jersey 08618-1414 registered as a bulk manufacturer of the following basic class(es), of controlled substances:

Controlled Substance	Drug Code	Schedule
4-Anilino-N-phenethyl-4-piperidine (ANPP)	8333	II
Levomethorphan	9210	II
Levorphanol	9220	II
Remifentanil	9739	II
Fentanyl	9801	II

The company plans to bulk manufacture API quantities of the listed controlled substances for validation purposes and FDA approval.

Dated: March 5, 2020.

William T. McDermott,

Assistant Administrator.

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