DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-583]

Bulk Manufacturer of Controlled Substances Application: Siemens Healthcare Diagnostics 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 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 December 10, 2019, Siemens Healthcare Diagnostics Inc., 100 GBC Drive, Mailstop 514, Newark, Delaware 19702-2461 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Drug Code	Schedule
Ecgonine	9180	II

The company plans to produce the listed controlled substance in bulk to be used in the manufacture of DEA exempt products.

Dated: February 11, 2020.

William T. McDermott,
Assistant Administrator.

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