Billing Code 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Bulk Manufacturer of Controlled Substances Application: Siemens Healthcare

Diagnostics Inc.

[Docket No. DEA-392]

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].

ADDRESS: Written comments should be sent to: Drug Enforcement Administration,

Attention: DEA Federal Register Representative/DRW, 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 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 1301.33(a), this is notice that on December 13, 2017, Siemens Healthcare Diagnostics Inc., 100 GBC Drive, Mailstop 514, Newark, DE 19702 applied to be registered as a bulk manufacturer of the following basic class of controlled substance:

Controlled Substance	Drug Code	Schedule
Ecgonine	9180	II

The company plans to bulk manufacture a material used in the manufacture of reagents for a Cocaine in vitro diagnostic test system.

Dated: February 1, 2018

Susan A. Gibson,

Deputy Assistant Administrator.

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