



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 class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) 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/ODW, 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 Deputy Assistant Administrator of the DEA Office of Diversion Control (“Deputy 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 14, 2015, Siemens Healthcare Diagnostics, Inc., 100 GBC Drive, Mailstop 514, Newark, Delaware 19702 applied to be registered as a bulk manufacturer of ecgonine (9180) a basic class of controlled substance listed in schedule II.

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

Dated: March 14, 2016

Louis J. Milione,
Deputy Assistant Administrator.