42696, 42697 (2018). ''Moreover, in the absence of an application (whether timely filed or not), there is nothing to act upon.'' *Reece,* 77 FR at 35055, *Rampey,* 83 FR at 42697. Accordingly, because Registrant has allowed his registration to expire and has not filed an application to renew his registration or for any other registration in Alabama, this case is now moot and will be dismissed.

Order

Pursuant to 28 CFR 0.100(b) and the authority thus vested in me by 21 U.S.C. 824(a), I order that the Order to Show Cause issued to Phillip O. Rawlings, Jr., M.D., be, and it hereby is, dismissed. This Order is effective immediately.

Dated: September 26, 2018.

Uttam Dhillon,

Acting Administrator.

[FR Doc. 2018–22421 Filed 10–12–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrant listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various classes of schedule I or II controlled substances.

SUPPLEMENTARY INFORMATION: The company listed below applied to be registered as an importer of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

Company	FR Docket	Published
Galephar Pharmaceutical Research Inc	83 FR 37525	August 1, 2018.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of the listed registrant to import the applicable basic classes of schedule I or II controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying the company's compliance with state and local laws, and reviewing the company's background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the DEA has granted a registration as an importer for schedule I or II controlled substance to the above listed company.

Dated: September 24, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–22420 Filed 10–12–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Cambrex High Point, 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 November 14, 2018. Such persons may also file a written request for a hearing on the application on or before November 14, 2018.

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 hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 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: 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 1301.34(a), this is notice that on July 16, 2018, Cambrex High Point Inc., 4180 Mendenhall Oaks Parkway, High Point, North Carolina 27265–8017 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Amphetamine	1100	II
Codeine	9050	II
Oxymorphone	9652	II
Noroxymorphone	9668	II

The company plans to manufacture the above listed controlled substances in bulk for distribution to its customers.

Dated: October 1, 2018.

John J. Martin,

Assistant Administrator.

[FR Doc. 2018–22416 Filed 10–12–18; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Specgx, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written