6(a) of the Act. The Department of Justice published a notice in the **Federal Register** pursuant to Section 6(b) of the Act on February 27, 2017 (82 FR 11942).

Patricia A. Brink,

Director of Civil Enforcement, Antitrust Division.

[FR Doc. 2017–20880 Filed 9–28–17; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration

ACTION: Notice of registration.

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

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as importers 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
United States Pharmacopeial.		
Convention	82 FR 34694	July 26, 2017.
AMRI Rensselaer, Inc	82 FR 34696	July 26, 2017.
R & D Systems, Inc	82 FR 35546	July 31, 2017.
Sigma-Aldrich International.		
GMBH	82 FR 35547	July 31, 2017.
Cambrex High Point, Inc	82 FR 35992	August 2, 2017.

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 registrants 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 each company's maintenance of effective controls against diversion by inspecting and testing each company's physical security systems, verifying each company's compliance with state and local laws, and reviewing each 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 substances to the above listed persons.

Dated: September 21, 2017.

Demetra Ashley,

Acting Assistant Administrator.

[FR Doc. 2017–20943 Filed 9–28–17; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Cody Laboratories, Inc.

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefor, may file written comments on or objections to the issuance of the proposed registration on or before October 30, 2017. Such persons may also file a written request for a hearing on the application on or before October 30, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW, 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/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152. Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25, 2007).

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 June 16, 2017, Cody Laboratories, Inc., Steve Hartman, 601 Yellowstone Avenue, Cody, Wyoming 82414–9321 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Phenylacetone	8501	II
Poppy Straw Concentrate.	9670	II
Tapentadol	9780	II

The company plans to import narcotic raw materials to manufacture bulk controlled substances for distribution to its customers. The company plans to import an intermediate form of tapentadol (9870), to bulk manufacture tapentadol for distribution to its customers.

Dated: September 21, 2017.

Demetra Ashley,

Acting Assistant Administrator.

[FR Doc. 2017–20942 Filed 9–28–17; 8:45 am]

BILLING CODE 4410–09–P