INTERNATIONAL TRADE COMMISSION

[USITC SE–17–012]

Government in the Sunshine Act Meeting Notice

AGENCY HOLDING THE MEETING: United States International Trade Commission.

TIME AND DATE: March 30, 2017 at 11:00 a.m.

PLACE: Room 101, 500 E Street SW., Washington, DC 20436, Telephone: (202) 205–2000.

STATUS: Open to the public.

Matters To Be Considered

1. *Agendas for future meetings:* None.
2. Minutes.
3. Ratification List.
4. Vote in Inv. No. 731–TA–1314 (Final) (Phosphor Copper from Korea). The Commission is currently scheduled to complete and file its determination and views of the Commission by April 17, 2017.
5. *Outstanding action jackets:* None.

In accordance with Commission policy, subject matter listed above, not disposed of at the scheduled meeting, may be carried over to the agenda of the following meeting.

By order of the Commission.

Issued: March 21, 2017.

William R. Bishop,

Supervisory Hearings and Information Officer.

[FR Doc. 2017–05893 Filed 3–21–17; 4:15 pm]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as bulk manufacturers of various classes of controlled substances.

SUPPLEMENTARY INFORMATION:

The companies listed below applied to be registered as manufacturers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted for these notices.

Company	FR docket	Published
Noramco, Inc	81 FR 57936	August 24, 2016.
Isosciences	81 FR 61249	September 6, 2016.
AMRI Rensselaer, Inc	81 FR 61250	September 6, 2016.
Halo Pharmaceutical, Inc	81 FR 63220	September 14, 2016.
AMPAC Fine Chemicals LLC	81 FR 63222	September 14, 2016.
Insys Manufacturing LLC	81 FR 63221	September 14, 2016.
Patheon API Manufacturing, Inc	81 FR 64509	September 20, 2016.
Euticals Inc	81 FR 64510	September 20, 2016.
Nanosyn, Inc	81 FR 64949	September 21, 2016.
Cerilliant Corporation	81 FR 66079	September 26, 2016.
Research Triangle Institute	81 FR 91948	December 19, 2016.
Synthcon LLC	81 FR 95641	December 28, 2016.
Navinta LLC	81 FR 95640	December 28, 2016.
Johnson Matthey, Inc	81 FR 95647	December 28, 2016.
AMRI Rensselaer, Inc	81 FR 95639	December 28, 2016.
Cayman Chemical Company	81 FR 95644	December 28, 2016.
Cambridge Isotope Laboratories	81 FR 95644	December 28, 2016.
Janssen Pharmaceutical, Inc	81 FR 96045	December 29, 2016.

The DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of these registrants to manufacture the applicable basic classes of 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 of the 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. 823(a), and in accordance with 21 CFR 1301.33, the DEA has granted a registration as a bulk manufacturer to the above listed persons.

Dated: March 17, 2017.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2017–05727 Filed 3–22–17; 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: Registrants 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 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
Fisher Clinical Services, Inc	81 FR 61248	September 6, 2016.
Sigma-Aldrich International:		
GMBH-Sigma Aldrich Company LLC	81 FR 63223	September 14, 2016.
Fisher Clinical Services, Inc	81 FR 68455	October 4, 2016.
Anderson Brecon, Inc	81 FR 71766	October 18, 2016.
Johnson Matthey Inc	81 FR 71766	October 18, 2016.
Wildlife Laboratories, Inc	81 FR 95644	December 28, 2016.
Noramco, Inc	81 FR 95640	December 28, 2016.
Mylan Technologies, Inc	82 FR 7859	January 23, 2017.

The 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: March 17, 2017.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2017–05730 Filed 3–22–17; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Application: Wildlife Laboratories, 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 in accordance with 21 CFR 1301.34(a) on or before April 24, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before April 24, 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.

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 February 2, 2017, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550–8055 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Drug code	Schedule
Etorphine (except HCl)	9056	I
Etorphine HCl	9059	II
Thiafentanil	9729	II

The company plans to import the listed controlled substances for sale to its customers.

Dated: March 17, 2017.

Louis J. Milione,

Assistant Administrator.

[FR Doc. 2017–05729 Filed 3–22–17; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Office of the Secretary

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Federal-State Unemployment Insurance Program Data Exchange Standardization

ACTION: Notice.

SUMMARY: The Department of Labor (DOL) is submitting the Employment and Training Administration (ETA) sponsored information collection request (ICR) titled, ''Federal-State Unemployment Insurance Program Data Exchange Standardization,'' to the Office of Management and Budget (OMB) for review and approval for continued use, without change, in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that agency receives on or before April 24, 2017.