recommendation by the Coast Guard Commandant. To be eligible, applicants should have particular expertise, knowledge, and experience regarding the regulations and policies on the pilotage of vessels on the Great Lakes, and at least 5 years of practical experience in maritime operations.

We will consider applicants for two positions that expire or become vacant on September 30, 2013.

- One member representing the interests of Great Lakes ports.
- One member representing the interests of shippers whose cargoes are transported through Great Lakes ports.

Members shall serve terms of office of up to three years and may be reappointed. All members serve at their own expense but may receive reimbursement for travel and per diem from the Federal Government.

Registered lobbyists are not eligible to serve on Federal Advisory Committees. Registered lobbyists are lobbyists required to comply with provisions contained in the *Lobbying Disclosure Act,* Title 2, United States Code, Section 1603.

The Department of Homeland Security (DHS) does not discriminate in employment on the basis of race, color, religion, sex, national origin, political affiliation, sexual orientation, gender identity, marital status, disability and genetic information, age, membership in an employee organization, or other non-merit factor. DHS strives to achieve a widely diverse candidate pool for all of its recruitment actions.

To visit our online docket, go to *http://www.regulations.gov* enter the docket number for this notice (USCG–2013–0702) in the Search box, and click ''Search''. Please do not post your resume on this site. Note, during the vetting process, applicants may be asked to provide date of birth and social security number.

Dated: August 9, 2013.

Mike M. Sollosi,

Acting Director, Marine Transportation Systems, U.S. Coast Guard.

[FR Doc. 2013–19742 Filed 8–13–13; 8:45 am]

BILLING CODE 9110–04–P

INTERNATIONAL TRADE COMMISSION

[USITC SE–13–019]

Government in the Sunshine Act Meeting Notice

AGENCY HOLDING THE MEETING: United States International Trade Commission.

TIME AND DATE: August 19, 2013 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. Nos. 701–TA–365–366 and 731–TA–734–735 (Third Review) (Certain Pasta from Italy and Turkey). The Commission is currently scheduled to complete and file its determinations and views of the Commission on or before August 30, 2013.
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: August 12, 2013.

William R. Bishop,

Supervisory Hearings and Information Officer.

[FR Doc. 2013–19857 Filed 8–12–13; 4:15 pm]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration; Alltech Associates, Inc.

By Notice dated May 14, 2013, and published in the **Federal Register** on May 22, 2013, 78 FR 30330, Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Lysergic acid diethylamide (7315)	I
Heroin (9200)	I
Cocaine (9041)	II
Codeine (9050)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Methadone (9250)	II
Morphine (9300)	II

The company plans to import these controlled substances for the manufacture of reference standards.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Alltech Associates, Inc. to import the 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. DEA has investigated Alltech Associates, Inc. to ensure that the company's registration is consistent with the public interest.

The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of 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 above named company is granted registration as an importer of the basic classes of controlled substances listed.

Dated: August 5, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–19634 Filed 8–13–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated May 14, 2013, and published in the **Federal Register** on May 22, 2013, 78 FR 30330, Arizona Department of Corrections, ASPC-Florence, 1305 E. Butte Avenue, Florence, Arizona 85132, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The facility intends to import the above listed controlled substance for legitimate use. Supplies of this particular controlled substance are inadequate and are not available in the form needed within the current domestic supply of the United States.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Arizona Department of Corrections to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties,

conventions, or protocols in effect on May 1, 1971.

DEA has investigated Arizona Department of Corrections, ASPC-Florence to ensure that its registration is consistent with the public interest. The investigation has included inspection and testing of the Arizona Department of Corrections, ASPC-Florence facility's physical security systems, verification of its compliance with state and local laws, and a review of its background and history.

Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR § 1301.34, the above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: August 5, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–19637 Filed 8–13–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances, Notice of Registration, Wildlife Laboratories Inc.

By Notice dated May 14, 2013, and published in the **Federal Register** on May 22, 2013, 78 FR 30329, Wildlife Laboratories Inc., 1401 Duff Drive, Suite 400, Fort Collins, Colorado 80524, made application by letter to the Drug Enforcement Administration (DEA) to be registered as an importer of Etorphine (except HCI) (9056), a basic class of controlled substance listed in schedule I.

The company plans to import the listed controlled substance for sale to its customer.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and 952(a) and determined that the registration of Wildlife Laboratories Inc. to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. DEA has investigated Wildlife Laboratories Inc. to ensure that the company's registration is consistent with the public interest. The investigation has included inspection and testing of the company's physical security systems, verification of the company's compliance with state and local laws, and a review of 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 above named company is granted registration as an importer of the basic class of controlled substance listed.

Dated: August 5, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–19621 Filed 8–13–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; Siegfried USA, LLC

This is notice that on June 10, 2013, Siegfried USA, LLC., 33 Industrial Park Road, Pennsville, New Jersey 08070, made application by letter to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Drug	Schedule
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II

The company plans to import the listed controlled substances to bulk manufacture API'S for distribution to its customer.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (2007).

As noted in a previous notice published in the **Federal Register** on September 23, 1975, 40 FR 43745, all applicants for registration to import a basic classes of any controlled substances in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: August 2, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–19745 Filed 8–13–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application; IRIX Manufacturing, Inc.

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on January 18, 2013, IRIX Manufacturing, Inc., 309 Delaware Street, Greenville, South Carolina 29605, made application to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed controlled substance as API for clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than October 15, 2013.

Dated: August 5, 2013.

Joseph T. Rannazzisi,

Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

[FR Doc. 2013–19616 Filed 8–13–13; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration; Wildlife Laboratories, Inc.

By Notice dated April 16, 2013, and published in the **Federal Register** on April 23, 2013, 78 FR 23958, Wildlife Laboratories, Inc., 1230 W. Ash Street, Suite D, Windsor, Colorado 80550, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Carfentanil (9743), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the above listed controlled substance for sale to veterinary pharmacies, zoos, and for other animal and wildlife applications.