Commission is properly sought will be treated accordingly. A redacted non-confidential version of the document must also be filed simultaneously with the any confidential filing. All non-confidential written submissions will be available for public inspection at the Office of the Secretary and on EDIS.

This action is taken under the authority of section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and of sections 201.10 and 210.50 of the Commission's Rules of Practice and Procedure (19 CFR 201.10, 210.50).

By order of the Commission.

Issued: May 9, 2014.

**Lisa R. Barton,**

*Secretary to the Commission.*

[FR Doc. 2014–11128 Filed 5–14–14; 8:45 am]

**BILLING CODE 7020–02–P**

## INTERNATIONAL TRADE COMMISSION

**[USITC SE–14–014]**

### Government in the Sunshine Act Meeting Notice

**AGENCY HOLDING THE MEETING:** United States International Trade Commission.

**TIME AND DATE:** May 21, 2014 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–1143 (Review) (Small Diameter Graphite Electrodes from China). The Commission is currently scheduled to complete and file its determination and views of the Commission on June 2, 2014.
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: May 12, 2014.

**William R. Bishop,**

*Supervisory Hearings and Information Officer.*

[FR Doc. 2014–11307 Filed 5–13–14; 11:15 am]

**BILLING CODE 7020–02–P**

## DEPARTMENT OF JUSTICE

### Notice of Lodging of an Amendment to Consent Decree Under the Clean Air Act

On May 9, 2014, the Department of Justice lodged with the United States District Court for the Southern District of Illinois a proposed Fourth Amendment to the consent decree in the lawsuit entitled *United States et al* v. *Lafarge North America, et al.,* Civil Action No. 3:10–cv–44–JPG.

The consent decree, which was entered by the Court on March 18, 2010, resolves claims of the United States and twelve states or state agencies against Lafarge North America, Inc., Lafarge Midwest, Inc., and Lafarge Building Materials, Inc. (''Lafarge'') for alleged violations of the Clean Air Act (''CAA'' or ''Act'') at its thirteen Portland cement production facilities in the United States. The proposed Fourth Amendment affects only two of the thirteen cement plants addressed in the Consent Decree: The Sugar Creek, Missouri cement plant and the Tulsa, Oklahoma cement plant. As of November 30, 2012, the Lafarge Companies have transferred ownership and operation of the Sugar Creek and Tulsa plants to Audubon Materials LLC and Tulsa Cement LLC, respectively. If approved by the Court, the Fourth Amendment would substitute Audubon Materials LLC for the Lafarge Companies at the Sugar Creek, Missouri plant, and would substitute Tulsa Cement LLC for the Lafarge Companies at the Tulsa, Oklahoma plant. Audubon Materials LLC, Tulsa Cement LLC, and their parent, Eagle Materials Inc. (collectively, ''the Eagle Companies'') have agreed in writing to assume the obligations, rights, and benefits of, and to be bound by the terms and conditions of, the Consent Decree, to the extent those terms and conditions relate to the Sugar Creek and Tulsa Cement plants.

The publication of this notice opens a period for public comment on the Fourth Amendment. Comments should be addressed to the Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States et al* v. *Lafarge North America, et al.,* Civil Action No. 3:10–cv–44–JPG, D.J. Ref No. 90–5–2–1–08221. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

| *To submit comments:* | *Send them to:* |
|---|---|
| By e-mail ..... | *pubcomment-ees.enrd@usdoj.gov.* |
| By mail ..... | Assistant Attorney General, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611. |

During the public comment period, the proposed Fourth Amendment may be examined and downloaded at this Justice Department Web site: *http://www.usdoj.gov/enrd/Consent_Decrees.html.* We will provide a paper copy of the proposed Fourth Amendment upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611.

Please enclose a check or money order for $12.75 (25 cents per page reproduction cost) payable to the United States Treasury. For a paper copy without the exhibits and signature pages, the cost is $9.00.

**Maureen Katz,**

*Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.*

[FR Doc. 2014–11151 Filed 5–14–14; 8:45 am]

**BILLING CODE 4410–15–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances, Notice of Application, Stepan Company

Pursuant to 21 CFR 1301.34(a), this is notice that on February 18, 2014, Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of Coca Leaves (9040), a basic class of controlled substance listed in schedule II.

The company plans to import the listed controlled substance to manufacture bulk controlled substance for distribution to its customer.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate. 72 FR 3417 (January 25,2007).

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 class of any controlled substance in schedules 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: May 1, 2014.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2014–11240 Filed 5–14–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–392]**

### Bulk Manufacturer of Controlled Substances Application: SIEMENS Healthcare Diagnostics, 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 1301.33(a) on or before July 14, 2014.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 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, and dispensers of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been re-delegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to sec. 7(g) of 28 CFR part 0, subpart R, App.

In accordance with 21 CFR 1301.33(a), this is notice that on November 26, 2013, Siemens Healthcare Diagnostics, Inc., Attn: RA, 100 GBC Drive, Mail Stop 514, Newark, Delaware 19702, applied to be registered as a bulk manufacturer of the following basic classes of narcotic or non-narcotic controlled substances:

| Controlled substance | Schedule |
|---|---|
| Tetrahydrocannabinols (7370) | I |
| Ecgonine (9180) | II |
| Morphine (9300) | II |

The company plans to produce the listed controlled substances in bulk to be used in the manufacture of reagents and drug calibrator controls which are DEA exempt products.

In reference to drug code 7370 the company plans to bulk manufacture a synthetic tetrahydrocannabinol. No other activity for this drug code is authorized for this registration.

Dated: May 2, 2014.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator.*

[FR Doc. 2014–11112 Filed 5–14–14; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–392]**

### Bulk Manufacturer of Controlled Substances Application: Alltech Associates, 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 1301.33(a) on or before July 14, 2014.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODW, 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, and dispensers of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been re-delegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to sec. 7(g) of 28 CFR part 0, subpart R, App.

In accordance with 21 CFR 1301.33(a), this is notice that on March 17, 2014, Alltech Associates, Inc., 2051 Waukegan Road, Deerfield, Illinois 60015, applied to be registered as a bulk manufacturer of the following basic classes of narcotic or non-narcotic controlled substances:

| Controlled substance | Schedule |
|---|---|
| Methcathinone (1237) | I |
| N-Ethylamphetamine (1475) | I |
| N,N-Dimethylamphetamine (1480) | I |
| 4-Methylaminorex (cis isomer) (1590) | I |
| Gamma Hydroxybutyric Acid (2010) | I |
| Alpha-ethyltryptamine (7249) | I |
| Lysergic acid diethylamide (7315) | I |
| 2C-T-7 (2,5-Dimethoxy-4-(n)- | I |
| propylthiophenethylamine) (7348) | I |
| Tetrahydrocannabinols (7370) | I |
| Mescaline (7381) | I |
| 2C-T-2 (2-(4-Ethylthio-2,5-dimethoxyphenyl)ethanamine) (7385) | I |
| 4-Bromo-2,5-dimethoxyamphetamine (7391) | I |
| 4-Bromo-2,5-dimethoxyphenethylamine (7392) | I |
| 4-Methyl-2,5-dimethoxyamphetamine (7395) | I |
| 2,5-Dimethoxyamphetamine (7396) | I |