confidential business information received by the Commission in this investigation and used in preparing its report will not be published in a manner that would reveal the operations of the firm supplying the information.

By order of the Commission.

Issued: October 10, 2008.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E8–24607 Filed 10–16–08; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[Investigation No. 701–TA–431 (Review)]

Drams and Dram Modules From Korea

AGENCY: United States International Trade Commission.

ACTION: Termination of five-year review.

SUMMARY: The subject five-year review was initiated in July 2008 to determine whether revocation of the countervailing duty order on DRAMs and DRAM modules from Korea would be likely to lead to continuation or recurrence of material injury. On October 3, 2008, the Department of Commerce published notice that it was revoking the order effective August 11, 2008, ''{b}ecause the domestic interested party did not file a substantive response by the applicable deadline and has withdrawn its notice of intent to participate in this sunset review * * *'' (73 FR 57594). Accordingly, pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)), the subject review is terminated.

DATES: *Effective Date:* August 11, 2008.

FOR FURTHER INFORMATION CONTACT: Mary Messer (202–205–3193), Office of Investigations, U.S. International Trade Commission, 500 E Street, SW., Washington, DC 20436. Hearing-impaired individuals are advised that information on this matter can be obtained by contacting the Commission's TDD terminal on 202–205–1810. Persons with mobility impairments who will need special assistance in gaining access to the Commission should contact the Office of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (*http://www.usitc.gov*).

Authority: This review is being terminated under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.69 of the Commission's rules (19 CFR 207.69).

By order of the Commission.

Issued: October 10, 2008.

Marilyn R. Abbott,

Secretary to the Commission.

[FR Doc. E8–24601 Filed 10–16–08; 8:45 am]

BILLING CODE 7020–02–P

INTERNATIONAL TRADE COMMISSION

[USITC SE–08–028]

Government in the Sunshine Act Meeting Notice

AGENCY HOLDING THE MEETING: United States International Trade Commission.

TIME AND DATE: October 21, 2008 at 11 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. *Agenda for future meetings:* none.
2. Minutes.
3. Ratification List.
4. Inv. Nos. 731–TA–1131–1134 (Final)(Polyethylene Terephthalate Film, Sheet, and Strip from Brazil, China, Thailand, and the United Arab Emirates)—briefing and vote. (The Commission is currently scheduled to transmit its determinations and Commissioners' opinions to the Secretary of Commerce on or before October 31, 2008.)
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: October 14, 2008.

William R. Bishop,

Hearings and Meetings Coordinator.

[FR Doc. E8–24769 Filed 10–16–08; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE

Notice of Lodging of Amended Consent Decree; Under the Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (''CERCLA'')

Consistent with Section 122(d) of the Comprehensive Environmental Response, Compensation, and Liability Act of 1980, as amended (''CERCLA''), 42 U.S.C. 9622(d), and 28 CFR 50.7, notice is hereby given that on October 7, 2008, the United States lodged an Amended Consent in *United States of America* v. *Lockheed Martin Corporation, et al.,* Civil No. 4:02–cv–146 (USDC W.D. Ky.) for the Green River Landfill Superfund Site, located in Maceo, Daviess County, Kentucky (the ''Site''). This Court originally approved a Consent Decree in this matter on September 27, 2002. Since the time the original Consent Decree was approved by the Court, the ''Settling Defendants'' as defined therein, and the United States Environmental Protection Agency (''EPA'') have been unable to implement the institutional controls required at the Site by Section IX of the Consent Decree. Under the proposed Amended Consent Decree, one ''Settling Defendant,'' Browning-Ferris Industries of Kentucky, Inc. (''BFIKY'') has or will acquire the property needed to institute the necessary institutional controls and, after entry of the Amended Consent Decree, will transfer such property to de maximus inc., defined in the proposed Amended Consent Decree as the ''Owner Settling Defendant.'' In addition, BFIKY will donate another parcel to Daviess County, which desires to keep it as open space. These property transfers will permit the remaining defendants to institute the required institutional controls and the open space will be an important buffer around the Site.

Under the proposed Amended Consent Decree, in exchange for the property transfers referenced above, BFIKY will have no further obligations under the Amended Consent Decree and will receive from the United States a covenant not to sue or to take administrative action pursuant to Sections 106 or 107 of Comprehensive Environmental Response, Compensation, and Liability Act of 1980 (''CERCLA''), 42 U.S.C. 9606 and 9607 as amended, and Section 7003 of the Resource Conservation and Recovery Act (''RCRA''), 42 U.S.C. 6973, for the United States' past and future costs at the Site. The remaining Settling Defendants will receive from the United States a covenant not to sue or to take administrative action pursuant to Sections 106 and 107 of CERCLA, 42 U.S.C. 9606 and 9607 as amended, and Section 7003 of RCRA, in exchange for implementing the remedy and required institutional controls at the Site and paying EPA's remaining costs under the terms of the proposed Amended Consent Decree.

The Department of Justice will receive for a period of thirty (30) days from the date of this publication comments relating to the Consent Decree Amendments. Comments should be addressed to the Assistant Attorney General, Environment and Natural

Resources Division, and either e-mailed to *pubcomment-ees.enrd@usdoj.gov* or mailed to P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611, and should refer to *United States of America* v. *Lockheed Martin Corporation, et al.,* Civil No. 4:02–cv–146 (USDC W.D. Ky.) (DOJ Ref. No. 90–11–2–1098).

The Amended Consent Decree may be examined at U.S. EPA Region 4, 61 Forsyth Street, Atlanta, GA 30303 (contact Kevin Beswick, Esq. (404) 562–9580). During the public comment period, the Amended Consent Decree may also be examined on the following Department of Justice Web site, *http://www.usdoj.gov/enrd/Consent_Decrees.html.* A copy of the Consent Decree may also be obtained by mail from the Consent Decree Library, U.S. Department of Justice, P.O. Box 7611, Washington, DC 20044–7611 or by faxing or e-mailing a request to Tonia Fleetwood *(tonia.fleetwood@usdoj.gov),* fax no. (202) 514–0097, phone confirmation number (202) 514–1547. In requesting a copy from the Consent Decree Library, please refer to *United States of America* v. *Lockheed Martin Corporation, et al.,* Civil No. 4:02–cv–146 (USDC W.D. Ky.) (DOJ Ref. No. 90–11–2–1098), and enclose a check in the amount of $29.00 (25 cents per page reproduction cost) payable to the U.S. Treasury or, if by e-mail or fax, forward a check in that amount to the Consent Decree Library at the stated address.

Henry S. Friedman,

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

[FR Doc. E8–24711 Filed 10–16–08; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a regulation under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on August 8, 2008, Fisher Clinical Services, Inc., 7554 Schantz Road, Allentown, Pennsylvania 18106, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Noroxymorphone (9668), a basic class of controlled substance listed in schedule II.

The company plans to import the listed substance for analytical research and clinical trials.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such 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 November 17, 2008.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). 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 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: October 9, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8–24774 Filed 10–16–08; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(i), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a regulation under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on August 5, 2008, Noramco Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801–4417, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Opium, Raw (9600) and Concentrate of Poppy Straw (9670), basic classes of controlled substances listed in schedule II.

The company plans to import the listed controlled substances to manufacture other controlled substances.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such 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 November 17, 2008.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). 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: October 9, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8–24780 Filed 10–16–08; 8:45 am]

BILLING CODE 4410–09–P