United States and Rhode Island v. *City of Newport, Rhode Island,* Civil Action No. 08–265S, was filed with the United States District Court for Rhode Island.

In this action, the United States and the other plaintiffs sought penalties and injunctive relief for the Defendant's violations of the Clean Water Act, 33 U.S.C. 1251 *et seq.,* at its sewer system and water pollution control plant. To resolve the United States' claims, the Defendants will pay a penalty of $170,000, and will undertake extensive work to its sewer system and water pollution control plant to eliminate violations of the Clean Water Act.

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. 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 either: *Environment Rhode Island et al. and the United States and Rhode Island* v. *City of Newport, Rhode Island,* Civil Action No. 08–265S, or D.J. Ref. 90–5–1–1–09855. The Consent Decree may be examined at the Office of the United States Attorney, District of Rhode Island, Fleet Center, 50 Kennedy Plaza, 8th Floor, Providence, Rhode Island 02903, and at the United States Environmental Protection Agency, 5 Post Office Square, Suite 100, Boston, Massachusetts 02109. During the public comment period, the 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, P.O. Box 7611, U.S. Department of Justice, 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 enclose a check, payable to the U.S. Treasury, in the amount of $21.50 (25 cents per page reproduction cost), or, if by e-mail or fax, forward a check in the applicable amount to the Consent Decree Library at the stated address.

Ronald Gluck,

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

[FR Doc. 2011–20996 Filed 8–17–11; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF JUSTICE

Notice of Proposed Partial Consent Decree Under the Comprehensive Environmental Response, Compensation, and Liability Act and the Clean Air Act

Notice is hereby given that on August 12, 2011, a proposed Partial Consent Decree in *United States* v. *C.A.I., Inc., et al.,* Civil Action No. 1:10–cv–10390–GAO, was lodged with the United States District Court for the District of Massachusetts.

The proposed Partial Consent Decree will settle the United States' claims on behalf of the U.S. Environmental Protection Agency (''EPA'') against Defendants C.A.I., Inc. (''CAI''), Sartorelli Realty, LLC (''SRLLC''), and Roy A. Nelson as Trustee of Nelson Danvers Realty Trust (''NDRT''), pursuant to Section 107 of the Comprehensive Environmental Response, Compensation, and Liability Act (''CERCLA''), 42 U.S.C. 9607, and Sections 112(r) and 114(a) of the Clean Air Act (''CAA''), 42 U.S.C. 7412(r), 7414(a), with respect to the Danversport Superfund Site, a former inks and paint products manufacturing facility, in Danvers, Massachusetts (''Site''). Pursuant to the Partial Consent Decree, based on demonstrations of limited financial resources: CAI will pay $400,000, including $300,000 in response costs under CERCLA and $100,000 as a civil penalty under the CAA; SRLLC will pay $150,000 in response costs; NDRT will pay $140,000 in response costs; and the settling defendants will transfer to the United States funds from an escrow account totaling approximately $27,000 as of March 2011. In addition, SRLLC and NDRT will make best efforts to sell the Site property and will transfer all net sales proceeds to the United States. Finally, the settling defendants will pay the United States 90% of any net proceeds from the resolution of other Site-related proceedings, up to the total amount of the United States' unreimbursed response costs. The proposed Partial Consent Decree, together with a Partial Consent Decree between the United States and Defendant Arnel Company, Inc. entered on July 1, 2011, will resolve this action in its entirety.

The Department of Justice will receive comments relating to the proposed Partial Consent Decree for a period of 30 days from the date of this publication. Comments on the Partial Consent Decree 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* v. *C.A.I., Inc., et al.,* Civil Action No. 1:10–cv–10390–GAO, D.J. Ref. 90–11–2–09184 & 90–11–2–09184/1.

During the public comment period, the proposed Partial Consent Decree may be examined at the following Department of Justice Web site: *http://www.usdoj.gov/enrd/Consent_Decrees.html.* A copy of the proposed Partial Consent Decree may also be obtained by mail from the Consent Decree Library, P.O. Box 7611, U.S. Department of Justice, Washington, DC 20044–7611 or by faxing or e-mailing a request to Tonia Fleetwood (*tonia.fleetwood@usdoj.gov*), fax number (202) 514–0097, phone confirmation number (202) 514–1547. If requesting a copy by mail from the Consent Decree Library, please enclose a check in the amount of $15.75 ($0.25 per page reproduction cost) payable to the U.S. Treasury or, if requesting by e-mail or fax, forward a check in that amount to the Consent Decree Library at the above-referenced address.

Ronald G. Gluck,

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

[FR Doc. 2011–21002 Filed 8–17–11; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to Title 21 Code of Federal Regulations 1301.34(a), this is notice that on May 4, 2011, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616–3466, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Drug	Schedule
Phenylacetone (8501)	II
Opium, raw (9600)	II
Poppy Straw Concentrate (9670)	II

The company plans to import the listed controlled substances to manufacture a bulk intermediate for sale to its customers. With regards to the phenylacetone, the company plans to use it as a base material in the bulk manufacture of another controlled substance.

No comments, objections, or requests for any hearings will be accepted on any application for registration or re-registration to import crude opium, poppy straw, concentrate of poppy straw, and coca leaves. As explained in the Correction to Notice of Application pertaining to Rhodes Technologies, 72 FR 3417 (2007), comments and requests for hearings on applications to import narcotic raw material are not appropriate.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act 21 U.S.C. 952(a)(2)(B) may, in the circumstances set forth in 21 U.S.C. 958(i), 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 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 September 19, 2011.

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: August 11, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011–21117 Filed 8–17–11; 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 21 CFR 1301.34(a), this is notice that on May 4, 2011, Cambrex Charles City, 1205 11th Street, Charles City, Iowa 50616–3466, 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 basic class of controlled substance in bulk for sale to its customers.

Any bulk manufacturers who are presently, or are applying to be, registered with DEA to manufacture such basic class of controlled substance 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 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 September 19, 2011.

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: August 10, 2011.

Joseph T. Rannazzisi,

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

[FR Doc. 2011–21121 Filed 8–17–11; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to Title 21, of the CFR 1301.34(a), this is notice that on June 8, 2011, Aptuit, 10245 Hickman Mills Drive, Kansas City, Missouri 64137, made application by renewal to the Drug Enforcement Administration (DEA) for registration as an importer of the following basic classes of controlled substances:

Drug	Schedule
Marihuana (7360)	I
Poppy Straw Concentrate (9670)	II

The company plans to import a finished pharmaceutical product containing cannabis extracts in dosage form for packaging for a clinical trial study. In addition, the company also plans to import an ointment for the treatment of wounds which contain trace amounts of the controlled substances normally found in poppy straw concentrate for packaging and labeling for clinical trials.

No comments, objections, or requests for any hearings will be accepted on any application for registration or re-registration to import crude opium, poppy straw, concentrate of poppy straw or coca leaves. As explained in the Correction to Notice of Application pertaining to Rhodes Technologies, 72 FR 3417(2007), comments and requests for hearings on applications to import narcotic raw material are not appropriate.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), 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.