*respond:* It is estimated that 2,000 respondents will complete the form annually within approximately 2.5 hours.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 5,000 total annual burden hours associated with this collection.

If additional information is required contact: Jerri Murray, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street, NE., Room 2E–508, Washington, DC 20530.

**Jerri Murray,**

*Department Clearance Officer, PRA, United States Department of Justice.*

[FR Doc. 2011–25988 Filed 10–6–11; 8:45 am]

**BILLING CODE 4410–12–P**

## DEPARTMENT OF JUSTICE

### Notice of Lodging of Settlement Agreement Under the Comprehensive Environmental Response, Compensation, and Liability Act

Notice is hereby given that on October 4, 2011, a proposed Settlement Agreement in the bankruptcy matter *In re DPH Holdings Corp., et al.,* Jointly Administered Case No. 05–44481 (RDD), was filed with the United States Bankruptcy Court for the Southern District of New York. The Settlement Agreement between the United States and DPH Holdings Corp., f/k/a Delphi Corp., and its affiliated reorganized debtors (''Reorganized Debtors'') resolves claims and causes of action of the United States on behalf of the Environmental Protection Agency (''EPA'') against debtor Delphi Automotive Systems LLC n/k/a DPH–DAS LLC under the Comprehensive Environmental Response, Compensation, and Liability Act of 1980, as amended, 42 U.S.C 9601–75 (''CERCLA''), and Section 7003 of the Resource Conservation and Recovery Act (''RCRA''), 42 U.S.C. 6973, with respect to the Tremont City Landfill Superfund Site in Tremont City, Ohio (''Tremont Site''), and the South Dayton Dump & Landfill Superfund Site in Moraine, Ohio (''South Dayton Site'').

Under the Settlement Agreement, the United States, on behalf of EPA, will have an allowed claim of $857,582.52. The allowed claim shall be allocated as an allowed claim of $559,292.95 for the Tremont Site and an allowed claim of $298,289.57 for the South Dayton Site. The effectiveness of the settlement is subject to the approval of a potential settlement of a tax refund action, *Delphi Corp., et al.* v. *United States,* Case No. 08 Civ. 4487 (PKC) (the ''Tax Refund Action''), pending in the United States District Court for the Southern District of New York. If the Tax Refund Action settlement is approved, the allowed claim of $857,582.52 shall be applied as a setoff against the refund that would be owed to the Reorganized Debtors. Pursuant to the Settlement Agreement, the Debtors and Reorganized Debtors will receive a covenant not to sue from the United States on behalf of EPA for the sites identified in this Notice, i.e., the Tremont Site and South Dayton Site.

Comments relating to the Settlement Agreement must be received by the Department of Justice no later than fourteen (14) days from the date of this publication. 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, D.C. 20044–7611, and should refer to *In re DPH Holdings Corp.,* D.J. Ref. _90–11–3–08913. Commenters may request an opportunity for a public meeting in the affected area, in accordance with Section 7003(d) of RCRA, 42 U.S.C. 6973(d).

The Settlement Agreement may be examined at the Office of the United States Attorney, 86 Chambers Street, 3rd Floor, New York, New York 10007, and at the U.S. Environmental Protection Agency, Ariel Rios Building, 1200 Pennsylvania Avenue, NW., Washington, DC 20460. During the public comment period, the Settlement Agreement may also be examined on the following Department of Justice Web site, *http://www.usdoj.gov/enrd/Consent_Decrees.html.* A copy of the Settlement Agreement also may 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 in the amount of $2.50 (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.

**Maureen M. Katz,**

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

[FR Doc. 2011–26037 Filed 10–6–11; 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 21 CFR 1301.34(a), this is notice that on August 11, 2011, 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 the following basic classes of controlled substances:

| Drug | Schedule |
|---|---|
| Noroxymorphone (9668) | II |
| Sufentanil (9740) | II |
| Tapentadol (9780) | II |

The company plans to import the listed substances 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 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 November 7, 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 43745–46, 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: September 27, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011–25989 Filed 10–6–11; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances; Notice of Registration

By Notice dated June 7, 2011, and published in the **Federal Register** on June 16, 2011, 76 FR 35241, Chattem Chemicals, Inc., 3801 St. Elmo Avenue, Building 18, Chattanooga, Tennessee 37409, 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 |
|---|---|
| Methamphetamine (1105) | II |
| 4-Anilino-N-phenethyl-4-piperidine (8333). | II |
| Phenylacetone (8501) | II |
| Opium, raw (9600) | II |
| Poppy Straw Concentrate (9670) | II |

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

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. With regard to all non-Narcotic Raw Material drugs on this application 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 Chattem Chemicals, 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 Chattem Chemicals, 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: September 28, 2011.

**Joseph T. Rannazzisi,**

*Deputy Assistant Administrator, Office of Diversion Control,*

Drug Enforcement Administration.

[FR Doc. 2011–26066 Filed 10–6–11; 8:45 am]

**BILLING CODE 4410–09–P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

### Importer of Controlled Substances; Notice of Application

This is notice that on July 19, 2011, Cody Laboratories Inc., 601 Yellowstone Avenue, Cody, Wyoming 82414–9321, 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 |
|---|---|
| Opium, raw (9600) | II |
| Poppy Straw Concentrate (9670) | II |

The company plans to import narcotic raw materials for manufacturing and further distribution to its customers. The company is registered with DEA as a manufacturer of several controlled substances that are manufactured from raw opium, poppy straw, and concentrate of poppy straw.

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.

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: September 28, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011–26068 Filed 10–6–11; 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 13, 2011, and published in the **Federal Register** on May 27, 2011, 76 FR 30969, Akorn, Inc., 1222 W. Grand Avenue, Decatur, Illinois 62522, made application to the Drug Enforcement Administration (DEA) to be registered as an importer of Remifentanil (9739), a basic class of controlled substance listed in schedule II.

The company plans to import Remifentanil in bulk for use in dosage-form manufacturing.

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 Akorn, 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 Akorn 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: September 28, 2011.

**Joseph T. Rannazzisi,**

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

[FR Doc. 2011–25992 Filed 10–6–11; 8:45 am]

**BILLING CODE 4410–09–P**