

| Drug   | Schedule |
|--|----------|
| Hydrocodone (9193) .....                                 | II       |
| Levomethorphan (9210) .....                              | II       |
| Levorphanol (9220) .....                                 | II       |
| Meperidine (9230) .....                                  | II       |
| Methadone (9250) .....                                   | II       |
| Dextropropoxyphene, bulk (non-dosage forms) (9273) ..... | II       |
| Morphine (9300) .....                                    | II       |
| Thebaine (9333) .....                                    | II       |
| Oxymorphone (9652) .....                                 | II       |
| Alfentanil (9737) .....                                  | II       |
| Sufentanil (9740) .....                                  | II       |

The company plans to import reference standards for sale to researchers and analytical labs.

The company plans to import the listed controlled substances in bulk powder form from foreign sources for the manufacture of analytical reference standards for sale to their customers.

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 United States Pharmacopeial Convention 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 United States Pharmacopeial Convention 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 29, 2013.

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

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration; GE Healthcare

By Notice dated June 7, 2013, and published in the **Federal Register** on June 18, 2013, 78 FR 36594, GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004-1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to import small quantities of ioflupane, in the form of three separate analogues of Cocaine, to validate production and quality control systems, for a reference standard, and for producing material for a future investigational new drug (IND) submission.

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 GE Healthcare 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 GE Healthcare 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 29, 2013.

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

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## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

#### Importer of Controlled Substances; Notice of Registration; Lipomed, Inc.

By Notice dated June 7, 2013, and published in the **Federal Register** on June 18, 2013, 78 FR 36591, Lipomed, Inc., One Broadway, Cambridge, Massachusetts 02142, 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 |
|--|----------|
| Cathinone (1235) .....                     | I        |
| Methcathinone (1237) .....                 | I        |
| 4-Mephedrone (1248) .....                  | I        |
| N-Ethylamphetamine (1475) .....            | I        |
| N,N-Dimethylamphetamine (1480) .....       | I        |
| Fenethylamine (1503) .....                 | I        |
| Aminorex (1585) .....                      | I        |
| 4-Methylaminorex (cis isomer) (1590) ..... | I        |
| Gamma Hydroxybutyric Acid (2010) .....     | I        |
| Methaqualone (2565) .....                  | I        |