XIII. NO REACQUISITION

Defendants may not reacquire any part of the Divestiture Assets during the term of this Final Judgment.

XIV. RETENTION OF JURISDICTION

This Court retains jurisdiction to enable any party to this Final Judgment to apply to this Court at any time for further orders and directions as may be necessary or appropriate to carry out or construe this Final Judgment, to modify any of its provisions, to enforce compliance, and to punish violations of its provisions.

XV. EXPIRATION OF FINAL JUDGMENT

Unless this Court grants an extension, this Final Judgment shall expire ten (10) years from the date of its entry.

XVI. PUBLIC INTEREST DETERMINATION

Entry of this Final Judgment is in the public interest. The parties have complied with the requirements of the Antitrust Procedures and Penalties Act, 15 U.S.C. 16, including making copies available to the public of this Final Judgment, the Competitive Impact Statement, and any comments thereon and the United States' responses to comments. Based upon the record before the Court, which includes the Competitive Impact Statement and any comments and response to comments filed with the Court, entry of this Final Judgment is in the public interest.

Dated this __ day of_____, 2015.

Court approval subject to procedures of Antitrust Procedures and Penalties Act, 15 U.S.C. 16

United States District Judge

SCHEDULE A

List of products and functionality included in ''Divested Product,'' as defined in Section II.L of this Final Judgment:

Dealertrack eCarList®;
Dealertrack AAX®;
Inventory+;
InventoryPro;
PriceDriver;
TrueTarget® (including TrueTarget® Appraisal and TrueTarget® Pricing Reports);
TrueTarget® Mobile;
Inventory+Mobile (including Inventory+ for iPhone® and Android);
Inventory Management Stocking and Sourcing;
TrueScore;
Inventory+ Appraisal Workflow;
Inventory+ Merchandising;
AutoInk and eBay Listing and Merchandising Tools (including integrated AutoInk description writer and direct distribution to leading Web sites such as backpage.com, Craigslist, eBay Motors);
Dealer Web sites (eCarList only);
Dealertrack AutoReel® with TruVoice™;
Inventory+ integrated, ''multi-site'' lead Management system (including Email Lead Management);
Dealertrack Interactive Automated Incentives;
OutClick™;
Inventory Health Report;
Lot Services;
PROShots;
Inventory+ New Car Pricing;
Dealertrack Inventory+ integration;
Inventory+ Multiplatform Listing;
Appraisal Central;
GroupTrade;
Software code for Inventory+ Exchange (including Social Trade and OpenTrade) and its predecessor Dealertrack Marketplace;
Ability to enable Dealertrack SmartChat® reporting within Inventory+ for customers who have both Inventory+ and SmartChat®; and
Fully integrated access and interoperability with Broker Connection.

[FR Doc. 2015–26042 Filed 10–9–15; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Importer of Controlled Substances Registration: Unither Manufacturing, LLC

ACTION: Notice of registration.

SUMMARY: Unither Manufacturing, LLC applied to be registered as an importer of a certain basic class of controlled substance. The Drug Enforcement Administration (DEA) grants Unither Manufacturing, LLC registration as an importer of this controlled substance.

SUPPLEMENTARY INFORMATION: By notice dated April 14, 2015, and published in the **Federal Register** on April 22, 2015, 80 FR 22552, Unither Manufacturing, LLC, 331 Clay Road, Rochester, New York 14623 applied to be registered as an importer of a certain basic class of controlled substance. No comments or objections were submitted for this notice.

The DEA has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Unither Manufacturing, LLC 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. The DEA investigated the company's maintenance of effective controls against diversion by inspecting and testing the company's physical security systems, verifying the company's compliance with state and local laws, and reviewing 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 methylphenidate (1724), a basic class of controlled substance listed in schedule II.

The company plans to import the listed substance as a raw material for updated testing purposes for EU customer requirements.

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

Dated: October 2, 2015.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2015–25881 Filed 10–9–15; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: American Radiolabeled Chemicals, 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 December 14, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODXL, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her 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, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on August 10, 2015, American Radiolabeled Chemicals, Inc., 101 Arc Drive, St. Louis, Missouri 63146 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled substance	Schedule
Gamma Hydroxybutyric Acid (2010)	I
Ibogaine (7260)	I
Lysergic Acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Dimethyltryptamine (7435)	I
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470)	I
Dihydromorphine (9145)	I
Heroin (9200)	I
Normorphine (9313)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Amobarbital (2125)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Ecgonine (9180)	II
Hydrocodone (9193)	II
Meperidine (9230)	II
Metazocine (9240)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273)	II
Morphine (9300)	II
Oripavine (9330)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Phenazocine (9715)	II
Carfentanil (9743)	II
Fentanyl (9801)	II

The company plans to manufacture small quantities of the listed controlled substances as radiolabeled compounds for biochemical research.

Dated: October 2, 2015.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2015–25882 Filed 10–9–15; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Cambridge Isotope Lab

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, 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 December 14, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODXL, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION:

The Attorney General has delegated her 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, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on August 7, 2015, Cambridge Isotope Lab, 50 Frontage Road, Andover, Massachusetts 01810 applied to be registered as a bulk manufacturer of morphine (9300), a basic class of controlled substance listed in schedule II.

The company plans to utilize small quantities of the listed controlled substance in the preparation of analytical standards.

Dated: October 2, 2015.

Louis J. Milione,

Deputy Assistant Administrator.

[FR Doc. 2015–25879 Filed 10–9–15; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–392]

Bulk Manufacturer of Controlled Substances Application: Apertus Pharmaceuticals

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 December 14, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/ODXL, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her 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, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control (''Deputy Assistant Administrator'') pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.33(a), this is notice that on August 6, 2015, Apertus Pharmaceuticals, 331 Consort Drive, St. Louis, Missouri 63011 applied to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Controlled Substance	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Remifentanil (9739)	II