

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

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 Noramco 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, at this time. DEA has investigated Noramco 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: March 16, 2010.

**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**

By Notice dated December 1, 2009, and published in the **Federal Register** on December 11, 2009 (74 FR 65788), Tocris Cookson Inc., 16144 Westwoods Business Park, Ellisville, Missouri 63021-4500, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedules I and II:

| Drug                                     | Schedule |
|--|----------|
| Marihuana (7360) .....                   | I        |
| Tetrahydrocannabinols (7370) .....       | I        |
| 4-Bromo-2,5-dimethoxyamphetamine (7391). | I        |
| 3,4-Methylenedioxyamphetamine (7405).    | I        |
| Amphetamine (1100) .....                 | II       |
| Phencyclidine (7471) .....               | II       |
| Cocaine (9041) .....                     | II       |

| Drug                       | Schedule |
|----------------------------|----------|
| Diprenorphine (9058) ..... | II       |
| Fentanyl (9801) .....      | II       |

The company plans to import small quantities of the above listed controlled substances for non-clinical, laboratory-based research only.

In reference to drug code 7360 (Marihuana), the company plans to import synthetic cannabinoid agonists. In reference to drug code 7370 (Tetrahydrocannabinols), the company will import a synthetic Delta-9-THC. No other activity for these drug codes is authorized for this registration.

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 Tocris Cookson 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, at this time. DEA has investigated Tocris Cookson 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: March 16, 2010.

**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**

By Notice dated January 6, 2010, and published in the **Federal Register** on January 13, 2010 (75 FR 1812), Mallinckrodt Inc., 3600 North Second Street, St. Louis, Missouri 63147, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule II:

| Drug                           | Schedule |
|--------------------------------|----------|
| Phenylacetone (8501) .....     | II       |
| Coca Leaves (9040) .....       | II       |
| Opium, raw (9600) .....        | II       |
| Poppy Straw Concentrate (9670) | II       |

The company plans to import the listed controlled substances for the manufacture of controlled substances in bulk for distribution to its 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 Mallinckrodt 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, at this time. DEA has investigated Mallinckrodt 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: March 16, 2010.

**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 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 USC 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 of the Code of Federal Regulations (CFR), 1301.34(a), this is notice that on January 22, 2010, Meridian Medical Technologies, 2555 Hermelin Drive, St. Louis, Missouri 63144, made