

Controlled substance	Drug code	Schedule
Hydrocodone .....	9193	II
Levorphanol .....	9220	II
Isomethadone .....	9226	II
Meperidine .....	9230	II
Meperidine-intermediate-A .....	9232	II
Meperidine intermediate-B .....	9233	II
Meperidine intermediate-C .....	9234	II
Methadone .....	9250	II
Methadone intermediate .....	9254	II
Dextropropoxyphene, bulk (non-dosage forms) .....	9273	II
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Opium, raw .....	9600	II
Opium extracts .....	9610	II
Opium fluid extract .....	9620	II
Opium tincture .....	9630	II
Opium, powdered .....	9639	II
Opium, granulated .....	9640	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Bezitramide .....	9800	II
Fentanyl .....	9801	II

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis. In reference to drug codes 7360 (Marihuana) and 7370 (Tetrahydrocannabinols) the company plans to import a synthetic cannabidiol and a synthetic tetrahydrocannabinol. No other activities for these drug codes are authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

**Marsha Ikner,**  
*Acting Deputy Assistant Administrator.*  
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**DEPARTMENT OF JUSTICE**  
**Drug Enforcement Administration**  
**[Docket No. 1324]**

**Importer of Controlled Substances**  
**Application: AndersonBrecon dba PCI**  
**Pharma Services; Correction**

**AGENCY:** Drug Enforcement Administration, Justice.  
**ACTION:** Notice of correction.

**SUMMARY:** The Drug Enforcement Administration (DEA) published a document in the **Federal Register** on March 6, 2024, concerning an application for an Importer of Controlled Substances. The document request removal of Dimethyltryptamine.

**SUPPLEMENTARY INFORMATION:**  
**Correction**

In the **Federal Register** on March 6, 2024, in FR Doc No: 89 FR 16029, FR No. 2024-04753, on pages 16029-16030 (2 pages), in the first column, remove the controlled substance Dimethyltryptamine from the list to read as follows:

Controlled substance	Drug code	Schedule
Cocaine .....	9041	II
Methadone .....	9250	II

**Marsha Ikner,**  
*Acting Deputy Assistant Administrator.*  
 [FR Doc. 2024-09785 Filed 5-3-24; 8:45 am]  
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**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**  
**[Docket No. DEA-1357]**

**Bulk Manufacturer of Controlled Substances Application: Pharmaron Manufacturing Services (US) LLC**

**AGENCY:** Drug Enforcement Administration, Justice.  
**ACTION:** Notice of application.

**SUMMARY:** Pharmaron Manufacturing Services (US) LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before July 5, 2024. Such persons may also file a written request for a hearing on the application on or before July 5, 2024.

**ADDRESSES:** The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for