

Controlled substance	Drug code	Schedule
Meperidine intermediate-C .....	9234	II
Metazocine .....	9240	II
Methadone .....	9250	II
Methadone intermediate .....	9254	II
Metopon .....	9260	II
Dextropropoxyphene, bulk (non-dosage forms) .....	9273	II
Morphine .....	9300	II
Oripavine .....	9330	II
Thebaine .....	9333	II
Dihydroetorphine .....	9334	II
Levo-alphaacetylmethadol .....	9648	II
Oxymorphone .....	9652	II
Noroxymorphone .....	9668	II
Phenazocine .....	9715	II
Thiafentanil .....	9729	II
Piminodine .....	9730	II
Racemethorphan .....	9732	II
Racemorphan .....	9733	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Bezitramide .....	9800	II
Fentanyl .....	9801	II
Moramide-intermediate .....	9802	II

The company plans to bulk manufacture the listed controlled substances for distribution to its customers. In reference to drug codes 7360 (Marihuana), and 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture these drugs as synthetic. No other activities for these drug codes are authorized for this registration.

**Kristi O'Malley,**

*Assistant Administrator.*

[FR Doc. 2022-17003 Filed 8-8-22; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

[Docket No. DEA-1055]

#### Importer of Controlled Substances Application: Catalent Pharma Solutions, LLC

**AGENCY:** Drug Enforcement  
Administration, Justice.

**ACTION:** Notice of application.

**SUMMARY:** Catalent Pharma Solutions, LLC has applied to be registered as an importer 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 September 8, 2022. Such persons may also file a written request for a hearing on the application on or before September 8, 2022.

**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 submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrisette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on July 1, 2022, Catalent Pharma Solutions, LLC, 3031 Red Lion Road, Philadelphia, Pennsylvania 19114, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Lysergic acid diethylamide .....	7315	I
5-Methoxy-N-N-dimethyltryptamine .....	7431	I
Psilocybin .....	7437	I
Psilocyn .....	7438	I
Tapentadol .....	9780	II

The company plans to import the listed controlled substances as finished dosage unit products for clinical trials, research, and analytical activities. 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.

**Kristi O'Malley,**

*Assistant Administrator.*

[FR Doc. 2022-16998 Filed 8-8-22; 8:45 am]

**BILLING CODE 4410-09-P**