

Controlled substance	Drug code	Schedule
Anileridine .....	9020	II
Cocaine .....	9041	II
Diphenoxylate .....	9170	II
Ecgonine .....	9180	II
Levorphanol .....	9220	II
Meperidine .....	9230	II
Meperidine Intermediate-A .....	9232	II
Meperidine Intermediate-B .....	9233	II
Meperidine Intermediate-C .....	9234	II
Dextropropoxyphene .....	9273	II
Morphine .....	9300	II
Levo-alphaacetylmethadol .....	9648	II
Alfentanil .....	9737	II
Remifentanil .....	9739	II
Sufentanil .....	9740	II
Carfentanil .....	9743	II
Tapentadol .....	9780	II
Fentanyl .....	9801	II

The company plans to bulk manufacture the listed controlled substances as analytical materials, proficiency test materials, and academic research materials for distribution to its customers. No other activities for these drug codes are authorized for this registration.

**Matthew J. Strait,**  
*Deputy Assistant Administrator.*  
 [FR Doc. 2022-04057 Filed 2-24-22; 8:45 am]  
**BILLING CODE P**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. DEA-965]

**Importer of Controlled Substances Application: Lyndra Therapeutics**

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

**SUMMARY:** Lyndra Therapeutics has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTAL INFORMATION** listed below for further drug information.

**DATES:** Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before March 28, 2022. Such persons may also file a written request for a hearing on the application on or before March 28, 2022.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing must

be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152. All requests for a hearing should also 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.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on January 13, 2022, Lyndra Therapeutics, 65 Grove Street, Suite 301, Watertown, Massachusetts 02472, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

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

The company plans to develop the formulation and process, and then manufacture the finished oral dosage form for use in preclinical and human clinical trials and analysis. No other activity for this drug code is 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.

**Matthew J. Strait,**  
*Deputy Assistant Administrator.*  
 [FR Doc. 2022-04061 Filed 2-24-22; 8:45 am]  
**BILLING CODE 4410-09-P**

**DEPARTMENT OF LABOR**

**Agency Information Collection Activities; Submission for OMB Review; Comment Request; Miner's Claim for Benefits and Employment History**

**ACTION:** Notice of availability; request for comments.

**SUMMARY:** The Department of Labor (DOL) is submitting this Office of Workers' Compensation Programs (OWCP)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

**DATES:** The OMB will consider all written comments that the agency receives on or before March 28, 2022.

**ADDRESSES:** Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find this particular information collection by selecting "Currently under 30-day Review—Open for Public Comments" or by using the search function.

Comments are invited on: (1) Whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) if the information will be processed and used in a timely manner; (3) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (4) ways to enhance the quality, utility and clarity of the information collection; and