

| Controlled substance | Drug code | Schedule |
|---|-----------|----------|
| Beta-hydroxyfentanyl | 9830 | I |
| Beta-hydroxy-3-methylfentanyl | 9831 | I |
| 3-Methylthiofentanyl | 9833 | I |
| Furanyl fentanyl (N-(1-phenethylpiperidin-4-yl)-N-phenylfuran-2-carboxamide) | 9834 | I |
| Thiofentanyl | 9835 | I |
| Beta-hydroxythiofentanyl | 9836 | I |
| Ocfentanil | 9838 | I |
| beta'-Phenyl fentanyl (N-(1-phenethylpiperidin-4-yl)-N,3-diphenylpropanamide; also known as β'-phenyl fentanyl; 3-phenylpropanoyl fentanyl) | 9842 | I |
| N-(1-phenethylpiperidin-4-yl)-N-phenyltetrahydrofuran-2-carboxamide | 9843 | I |
| Crotonyl fentanyl ((E-N-(1-phenethylpiperidin-4-yl)-N-phenylbut-2-enamide) | 9844 | I |
| Cyclopropyl Fentanyl | 9845 | I |
| ortho-Fluorobutryl fentanyl (N-(2-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)butyramide; also known as 2-fluorobutryl fentanyl) | 9846 | I |
| Nabilone | 7379 | II |
| Alphaprodine | 9010 | II |
| Levomethorphan | 9210 | II |
| Racemethorphan | 9732 | II |
| Alfentanil | 9737 | II |

The company plans to import the listed controlled substances for distribution for analytical testing purposes. 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.

Claude Redd,
Acting Deputy Assistant Administrator.
 [FR Doc. 2023-25989 Filed 11-22-23; 8:45 am]
BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1297]

Bulk Manufacturer of Controlled Substances Application: Sigma Aldrich Research Biochemicals, Inc.

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

SUMMARY: Sigma Aldrich Research Biochemicals, Inc. 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 January 23, 2024. Such

persons may also file a written request for a hearing on the application on or before January 23, 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 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.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 27, 2023, Sigma Aldrich Research Biochemicals, Inc., 400-600 Summit Drive, Burlington, Massachusetts 01803, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|------------------------|-----------|----------|
| Lisdexamfetamine | 1205 | I |

The company plans to manufacture the listed controlled substances as reference standards. No other activities

for these drug codes are authorized for this registration.

Claude Redd,
Acting Deputy Assistant Administrator.
 [FR Doc. 2023-25992 Filed 11-22-23; 8:45 am]

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DEPARTMENT OF JUSTICE

[OMB Number 1121-0330]

Agency Information Collection Activities; Proposed eCollection eComments Requested; Law Enforcement Congressional Badge of Bravery

AGENCY: Office of Justice Programs, Department of Justice.
ACTION: 30-Day notice.

SUMMARY: The Office of Justice Programs, Bureau of Justice Assistance (BJA), Department of Justice (DOJ), will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection was previously published in the **Federal Register** on September 15, allowing a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until December 26, 2023.

FOR FURTHER INFORMATION CONTACT: If you have comments especially on the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions or additional information, please contact: Gregory Joy at 202-514-1369, Policy Advisor, Bureau of Justice Assistance, 810 7th Street NW,