## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–831]**

### Importer of Controlled Substances Application: VHG Labs DBA LGC Standards

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

**ACTION:** Notice of application.

**SUMMARY:** VHG Labs DBA LGC Standards 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 June 3, 2021. Such persons may also file a written request for a hearing on the application on or before June 3, 2021.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA **Federal Register** Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on April 21, 2021, VHG Labs DBA LGC Standards, 3 Perimeter Road, Manchester, New Hampshire 03103, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Fentanyl related-compounds as defined in 21 CFR 1308.11(h) | 9850 | I |
| Oxycodone | 9143 | II |
| Hydromorphone | 9150 | II |

The company plans to import the listed controlled substances for sale to research facilities for drug testing and analysis. 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.

**William T. McDermott,**

*Assistant Administrator.*

[FR Doc. 2021–09302 Filed 5–3–21; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–828]**

### Importer of Controlled Substances Application: Wildlife Laboratories, LLC

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

**ACTION:** Notice of application.

**SUMMARY:** Wildlife Laboratories, LLC 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 June 3, 2021. Such persons may also file a written request for a hearing on the application on or before June 3, 2021.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA **Federal Register** Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA **Federal Register** Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.34(a), this is notice that on April 8, 2021, Wildlife Laboratories, LLC, 1230 W Ash Street, Unit D, Windsor, Colorado 80550–4677, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Etorphine HCl | 9059 | II |
| Thiafentanil | 9729 | II |

The company plans to import the listed controlled substances for distribution to its customers. No other activity for these drug codes 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.

**William T. McDermott,**

*Assistant Administrator.*

[FR Doc. 2021–09300 Filed 5–3–21; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Drug Enforcement Administration

**[Docket No. DEA–830]**

### Bulk Manufacturer of Controlled Substances Application: Cargill, Incorporated

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

**ACTION:** Notice of application.

**SUMMARY:** Cargill, Inc., has applied to be registered as a bulk manufacturer 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 July 6, 2021. Such persons may also file a written request for a hearing on the application on or before July 6, 2021.

**ADDRESSES:** Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

**SUPPLEMENTARY INFORMATION:** In accordance with 21 CFR 1301.33(a), this is notice that on March 31, 2021, Cargill, Incorporated, 17540 Monroe Wapello Road, Eddyville, Iowa 52553, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

| Controlled substance | Drug code | Schedule |
|---|---|---|
| Gamma Hydroxybutyric Acid | 2010 | I |

The company plans to bulk manufacture butanediol as a raw material for industrial and consumer products. Gamma Hydroxybutyric Acid will be manufactured as a byproduct and an impurity waste of butanediol. The company does not plan to bulk manufacture this drug. No other activities for this drug code are authorized for this registration.

**William T. McDermott,**

*Assistant Administrator.*

[FR Doc. 2021–09301 Filed 5–3–21; 8:45 am]

**BILLING CODE P**

## DEPARTMENT OF JUSTICE

### Executive Office for Immigration Review

### Fee Waiver Request; Correction

**AGENCY:** Executive Office for Immigration Review, Department of Justice.

**ACTION:** 60-Day notice; correction.

**SUMMARY:** The Executive Office for Immigration Review, Department of Justice, submitted a 60-day notice for publishing in the **Federal Register** on March 4, 2021 soliciting comments to an information collection request *Fee Waiver Request,* to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. Please disregard the duplicate 60-day notice, which was inadvertently published on April 28, 2020.

**DATES:** April 29, 2020.

**FOR FURTHER INFORMATION CONTACT:** If you have additional 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 Lauren Alder Reid, Assistant Director, Office of Policy, Executive Office for Immigration Review, 5107 Leesburg Pike, Suite 2500, Falls Church, VA 22041, telephone: (703) 305–0289.

**SUPPLEMENTARY INFORMATION:**

### Correction

The 60-day notice for the *Fee Waiver Request* was published in the **Federal Register** of March 4, 2021 in FR Doc. 2021–04418, on page 12713. Please disregard the duplicate published on April 28, 2021 in FR Doc. 2021–08807, on page 22457.

Dated: April 29, 2021.

**Melody Braswell,**

*Departmental Clearance Officer.*

[FR Doc. 2021–09370 Filed 5–3–21; 8:45 am]

**BILLING CODE 4410–30–P**

## DEPARTMENT OF LABOR

### Employment and Training Administration

### Notice of Intent To Reestablish the Advisory Committee on Apprenticeship (ACA) Charter and Request for Member Nominations

**AGENCY:** Employment and Training Administration (ETA), Labor.

**ACTION:** Notice.

**SUMMARY:** The Secretary of Labor (Secretary) has determined that the reestablishment of the Advisory Committee on Apprenticeship (ACA or Committee) is necessary and in the public interest. The Department of Labor (DOL) intends to reestablish the ACA charter with revisions which are not intended to change the Committee's purpose or original intent. The revisions update the charter to ensure its closer alignment with the Department's current apprenticeship priorities. Additionally, DOL is requesting nominations of qualified candidates to be considered for appointment to the ACA.

**DATES:** The reestablishing ACA charter will be filed May 19, 2021. ACA member nominations must be received by June 3, 2021.

**ADDRESSES:** DOL has adopted a maximum telework posture in response to the COVID–19 pandemic. As such, nominations for individuals to serve on the ACA should be submitted electronically. Interested persons may submit ACA nominations, including relevant attachments, through any of the following methods:

*Electronically:* Send to: *AdvisoryCommitteeonApprenticeship@dol.gov* (and please specify in the email subject line, ''Nominations for Advisory Committee on Apprenticeship (ACA).''

*If you do not have access to an electronic means of submission*: please call the Office of Apprenticeship on (202) 693–3795, and leave a message and someone will coordinate a mail submission; however, the Department highly encourages electronic submissions as provided above.

*Mail, express delivery, messenger service, or courier service:* Submit one copy of the documents listed above to the following address: U.S. Department of Labor, Employment and Training Administration, Office of Apprenticeship, ACA, Room C– 5321, 200 Constitution Avenue NW, Washington, DC 20210.

**FOR FURTHER INFORMATION CONTACT:** For any questions concerning the ACA nomination process, please contact Ms. Kenya Huckaby, Executive Assistant, Employment and Training Administration, Office of Apprenticeship, at *Huckaby.Kenya@dol.gov,* telephone (202) 693–3795 (this is not a toll-free number).

**SUPPLEMENTARY INFORMATION:** Registered Apprenticeship is a unique public private partnership that is highly dependent on the engagement and involvement of its stakeholders and partners for its ongoing operational effectiveness. Apart from the ACA, there is no single organization or group with the broad representation of labor, employers, and the public available to consider the complexities and relationship of apprenticeship activities to other training efforts or to provide advice on such matters to the Secretary. It is particularly important to have such perspectives as DOL considers the expansion of registered apprenticeship, fundamentally instilling a permanent culture of inclusion in our workforce, and supports our Nation's economic recovery in the aftermath of the COVID–19 pandemic. The ACA's insight and recommendations on the best ways to address critical apprenticeship issues to meet the emerging needs of industry, labor, and the public is critical. For these reasons, the Secretary has determined that the reestablishment of a national advisory committee on apprenticeship is necessary and in the public interest.

There is currently no active charter for the ACA as the previous ACA charter expired on December 19, 2018. The pending charter has been revised to ensure alignment with current DOL priorities in the following four sections: (1) Objectives and Scope of Activities; (2) Description of Duties; (3) Designated Federal Officer (DFO); and (4) Membership and Designation.

Summary of the Charter Changes:

1. The Objectives and Scope of Activities section has been updated to reflect the current priorities of the Administration and charge the ACA with providing advice and recommendations on ways to better utilize the apprenticeship training model in order to provide equitable career pathways that advance the dignity of work for everyone.