

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
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 April 21, 2021, VH
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-com- pounds 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
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 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
Morrisette 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):