of the Secretary at 202–205–2000. General information concerning the Commission may also be obtained by accessing its Internet server (http://www.usitc.gov). The public record for this review may be viewed on the Commission's electronic docket (EDIS) at http://edis.usitc.gov.

For further information concerning the conduct of this review and rules of general application, consult the Commission's Rules of Practice and Procedure, part 201, subparts A through E (19 CFR part 201), and part 207, subparts A, D, E, and F (19 CFR part 207).

SUPPLEMENTARY INFORMATION: On November 6, 2015, the Commission determined that it should proceed to full reviews in the subject five-year reviews pursuant to section 751(c) of the Tariff Act of 1930 (19 U.S.C. 1675(c)). The Commission found that the domestic interested party group response to its notice of institution (80 FR 46048, August 3, 2015) was adequate and that the respondent interested party group with respect to the review on subject imports from Taiwan was adequate, and decided to conduct a full review of the antidumping duty order on narrow woven ribbons with woven selvedge from Taiwan. Although the Commission received a response to its notice of institution from Chinese respondent interested parties, the Commission found that the respondent interested party group response with respect to the reviews on subject imports from China was inadequate.1 However, the Commission determined

to conduct full reviews concerning the

orders on narrow woven ribbons with

woven selvedge from China to promote administrative efficiency in light of its decision to conduct a full review with respect to the review on subject imports from Taiwan. A record of the Commissioners' votes, the Commission's statement on adequacy, and any individual Commissioner's statements will be available from the Office of the Secretary and at the Commission's Web site.

Authority: These reviews are being conducted under authority of title VII of the Tariff Act of 1930; this notice is published pursuant to section 207.62 of the Commission's rules.

By order of the Commission. Issued: November 19, 2015.

#### Lisa R. Barton.

Secretary to the Commission.

[FR Doc. 2015–29938 Filed 11–24–15; 8:45 am]

BILLING CODE 7020-02-P

#### **DEPARTMENT OF JUSTICE**

# Drug Enforcement Administration [Docket No. DEA-392]

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

**ACTION:** Notice of application.

**DATES:** Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.34(a) on or before December 28, 2015. Such persons may also file a written request

for a hearing on the application pursuant to 21 CFR 1301.43 on or before December 28, 2015.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODXL, 8701 Morrissette Drive, Springfield, Virginia 22152. Request for hearings should be sent to: Drug Enforcement Administration, Attention: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated her authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA), 28 CFR 0.100(b). Authority to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Deputy Assistant Administrator of the DEA Office of Diversion Control ("Deputy Assistant Administrator") pursuant to section. 7 of 28 CFR part 0, appendix to subpart

In accordance with 21 CFR 1301.34(a), this is notice that on December 03, 2014, VHG Labs DBA LGC Standards Warehouse, 3 Perimeter Road, Manchester, New Hampshire 03103 applied to be registered as an importer of the following basic classes of controlled substances:

Controlled substance	Schedule
3-Fluoro-N-methylcathinone (3-FMC) (1233) Cathinone (1235) Methcathinone (1237)	!
Cathinone (1235)	
4-Fluoro-N-methylcathinone (4-FMC) (1238)	1
Pentedrone (a-methylaminovalerophenone) (1246)	i
Mephedrone (4-Methyl-N-methylcathinone) (1248)	li
4-Methyl-N-ethylcathinone (4-MEC) (1249)	i
Naphyrone (1258)	1
N-Ethylamphetamine (1475)	1
N,N-Dimethylamphetamine (1480)	!
Fenethylline (1503)	!
4-Methylaminorex (cls isomer) (1990)	
Mathagualone (2555)	1
Methaqualone (2565)	i
JWH-250 (1-Pentyl-3-(2-methoxyphenylacetyl) indole) (6250)	li
SR-18 (Also known as RCS-8) (1-Cyclohexylethyl-3-(2-methoxyphenylacetyl) Indole) (7008)	l
5-Fluorò-UR-144 and XLR11 [1-(5-Flouro-pentyl)-1H-indol-3-yl] (2,2,3,3-tetramethylcyclopropyl) methanone (7011)	1
AB-FUBINACA (N-(1-amino-3-methyl-1-oxobutan-2-yl) -1-(4-fluorobenzyl)-1H-indazole-3-carboxamide) (7012)  JWH-019 (1-Hexyl-3-(1-naphthoyl)indole) (7019)	1
ADB-PINACA (N-1-amino-3,3-dimethyl-1-oxobutan-2-yl)-1-pentyl-1H-indazole-3-carboxamide) (7035)	i
APINACA and AKB48 N-(1-Adamantyl)-1-pentyl-1H-indazole-3-carboxamide (7048)	

Controlled substance	Schedule
	22000
JWH-081 (1-Pentyl-3-(1-(4-methoxynaphthoyl) Indole) (7081)	
SR-19 (Also known as RCS-4) (1-Pentyl-3-[(4-methoxy)-benzoyl] indole (7104)	!
JWH-018 (also known as AM678) (1-Pentyl-3-(1-naphthoyl)indole) (7118)	
UR-144 (1-Pentyl-1H-indol-3-yl) (2,2,3,3-tetramethylcyclopropyl) methanone (7144)	
JWH-073 (1-Butyl-3-(1-naphthoyl)indole) (7173)	li
JWH-200 (1-[2-(4-Morpholinyl)ethyl]-3-(1-naphthoyl)indole) (7200)	Ì
AM-2201 (1-(5-Fluoropentyl)-3-(1-naphthoyl) Indole) (7201)	1
JWH-203 (1-Pentyl-3-(2-chlorophenylacetyl)Indole) (7203)	!
PB-22 (Quinolin-8-yl 1-pentyl-1H-indole-3-carboxylate) (7222)	
5F-PB-22 (Quinolin-8-yl 1-(5-fluoropentyl)-1H-indole-3-carboxylate) (7225)	
lbogaine (7260)	li
CP-47,497 (5-(1,1-Dimethylheptyl)-2-[(1R,3S)-3-hydroxycyclohexyl-phenol) (7297)	i
CP-47497 C8 Homologue (5-(1,1-Dimethyloctyl)-2-[(1R,3S)3-hydroxycyclohexyl-phenol) (7298)	1
Lysergic acid diethylamide (7315)	1
2,5-Dimethoxy-4-(n)-propylthiophenethylamine (2C-T-7) (7348)	!
Marihuana (7360) Tetrahydrocannabinols (7370)	
Parahexyl (7374)	li
Mescaline (7381)	1
2-(4-Elthylthio-2,5-dimethoxyphenyl) ethanamine (2C-T-2) (7385)	1
3,4,5-Trimethoxyamphetamine (7390)	1
4-Bromo-2,5-dimethoxyamphetamine (7391)	
4-Bromo-2,5-dimethoxyphenethylamine (7392)	
2,5-Dimethoxyamphetamine (7396)	
JWH-398 (1-Pentyl-3-(4-chloro-1-naphthoyl)Indole (7398)	
2,5-Dimethoxy-4-ethylamphetamine (7399)	Ì
3,4-Methylenedioxyamphetamine (7400)	
5-Methoxy-3,4-methylenedioxyamphetamine (7401)	!
N-Hydroxy-3,4-methylenedioxyamphetaimine (7402)	
3,4-Methylenedioxy-N-ethylamphetamine (7404)	i
4-Methoxyamphetamine (7411)	li
Peyote (7415)	i
5-Methoxy-N-N-dimethyltryptamine (7431)	1
Alpha-methyltryptamine (7432)	<u> </u>
Bufotenine (7433)	
Dimethyltryptamine (7435)	i
Psilocybin (7437)	li
Psilocyn (7438)	1
5-Methoxy-N,N-diisopropyltryptamine (7439)	1
N-Ethyl-1-phenylcyclohexylamine (7455)	
1-(1-Phenylcyclohexyl)pyrrolidine (7458)	
1-[1-(2-Thienyl)cyclohexyl]piperidine (7470)	
N-Ethyl-3-piperidyl benzilate (7482)	
N-Methyl-3-piperidyl benzilate (7484)	i
N-Benzylpiperazine (7493)	1
4-Methyl-alphapyrrolidinopropiophenone (4-MePPP) (7498)	
2-(2,5-Dimethoxy-4-methylphenyl) ethanamine (2C-D) (7508)	
2-(2,5-Dimethoxy-4-ethylphenyl) ethanamine (2C-E) (7509)	
2-(2,5-bilinethoxyphenyl) ethanamine (2C-I) (7517)  2-(4-lodo-2,5-dimethoxyphenyl) ethanamine (2C-I) (7518)	li
2-(4-Chloro-2,5-dimethoxyphenyl) ethanamine (2C-C) (7519)	i i
2-(2,5-Dimethoxy-4-nitro-phenyl) ethanamine (2C-N) (7521)	1
2-(4-Isopropylthio)-2,5-dimethoxyphenyl) ethanamine (2C-T-4) (7532)	!
MDPV (3,4-Methylenedioxypyrovalerone) (7535)	
2-(4-bromo-2,5-dimethoxyphenyl)-N-(2-methoxybenzyl) ethanamine (25B-NBOMe) (7536)	
Methylone (3,4-Methylenedioxy-N-methylcathinone) (7540)	li
Butylone (7541)	li
Pentylone (7542)	1
alpha-pyrrolidinopentiophenone (a-PVP) (7545)	1
alpha-pyrrolidinobutiophenone (a-PBP) (7546)	1
AM-694 (1-(5-Fluropentyl)-3-(2-iodobenzoyl)Indole) (7694)	
Acetyldihydrocodeine (9051)	
Benzylmorphine (9052)	
Cyprenorphine (9054)	i
Desomorphine (9055)	1
Etorphine (except HCI) (9056)	11

Controlled substance	Schedule
Codeine methylbromide (9070)	1
Dihydromorphine (9145)	i
Difenoxin (9168)	I
Heroin (9200)	
Methyldesorphine (9302)	i
Methyldihydromorphine (9304)	I
Morphine methylbromide (9305)	
Morphine-N-oxide (9307)	i
Myrophine (9308)	I
Nicocodeine (9309)	
Normorphine (9313)	i
Pholocodine (9314)	
Thebacon (9315)	l i
Drotebanol (9335)	İ
Acetylmethadol (9601)	
Allylprodine (9602)	
Alphameprodine (9604)	İ
Alphamethadol (9605)	
Betacetylmethadol (9607)	İ
Betameprodine (9608)	I
Betamethadol (9609)	
Betaprodine (9611)	
Dextromoramide (9613)	I
Diampromide (9615)	
Dimenoxadol (9617)	i
Dimethylthiambutene (9619)	Į.
Dioxaphetyl butyrate (9621)	
Ethylmethylthiambutene (9623)	i
Etonitazene (9624)	I I
Etoxeridine (9625)	
Ketobemidone (9628)	1
Levomoramide (9629)	
Morpheridine (9632)	i
Noracymethadol (9633)	1
Normethadone (9635)  Norpipanone (9636)	
Phenadoxone (9637)	i
Phenoperidine (9641)	!
Piritramide (9642)	
Properidine (9644)	i
Racemoramide (9645)	Į.
Trimeperidine (9646)	
Propiram (9649)`	i
1-Methyl-4-phenyl-4-propionoxypiperdine (9661)	
1-(2-Phenylethyl)-4-phenyl-4-acetoxypiperdine (9663) Tilidine (9750)	l i
Para-Fluorofentanyl (9812)	ĺ
3-Methylfentanyl (9813)	
Alpha-methylfentanyl (9814)	i
Beta-hydroxyfentanyl (9830)`	į.
Beta-hydroxy-3-methylfentanyl (9831)	
Alpha-methylthiofentanyl (9832)	li
Thiofentanyl (9835)	I.
Methamphetamine (1105)	II II
Phenmetrazine (1631)	II   II
Pentobarbital (2270)	II
Secobarbital (2315)	
GIGCOTIITIIQO (2000)	- 11

Controlled substance	Schedu
	II
	II
	II
Phenylacetone (8501)	II
	II
	II
	II
torphine HCI (9059)	II
Dihydrocodeine (9120)	II
	II
cgonine (9180)	II
	II
Neperidine intermediate-C (9234)	II
	II
	II
	II
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	ii

The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes. Placement of these drug codes onto the company's registration does not translate into automatic approval of subsequent permit applications to import controlled substances.

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 FDA approved or non-approved finished dosage forms for commercial sale.

Dated: November 19, 2015.

#### Louis J. Milione,

Deputy Assistant Administrator. [FR Doc. 2015–29934 Filed 11–24–15; 8:45 am]

BILLING CODE 4410-09-P

### DEPARTMENT OF JUSTICE

## **Drug Enforcement Administration**

# William Mikaitis, M.D.; Decision and Order

On July 23, 2015, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to William Mikaitis, M.D. (Registrant), of Lockport, Illinois. The Show Cause Order proposed the revocation of Registrant's DEA Certificate of Registration AM1585770, and the denial of any applications to renew or modify the registration as well as any other applications for a DEA registration, on the ground that he "do[es] not have authority to practice medicine or handle controlled substances in Illinois, the [S]tate in which [he] is registered with the DEA." Show Cause Order at 1 (citing 21 U.S.C. 823(f) and 824(a)(3)).

The Show Cause Order alleged that Registrant is registered with the DEA as a practitioner authorized to handle controlled substances in Schedules II through V at the registered address of 1206 E. 9th St., Suite 210, Lockport, Illinois. *Id.* The Order alleged that Registrant's registration expires by its terms on January 31, 2018. *Id.* 

The Order further alleged that "[e]ffective March 5, 2015, the State of Illinois, Department of Financial and Professional Regulation (IDFPR), Division of Professional Regulation, issued an Order in which the IDFPR temporarily suspended [his] Illinois [P]hysician and [S]urgeon [L]icense and [his] controlled substance licenses" and that "[t]his Order remains in effect." Id. The Show Cause Order thus asserted that "DEA must revoke [his] registration based upon [his] lack of authority to handle controlled substances in the State of Illinois." Id. (citing 21 U.S.C. 802(21), 823(f) and 824(a)(3)).

The Show Cause Order also notified Registrant of his right to request a hearing on the allegations or to submit a written statement in lieu of a hearing, the procedure for electing either option, and the consequence for failing to elect either option. *Id.* at 2 (citing 21 CFR 1301.43).

On July 30, 2015, a Diversion Investigator met Registrant at the office of his attorney and personally served the Show Cause Order on him. See GX 5 (Affidavit of DI). The Government