Controlled substance	Drug code	Schedule
Codeine	9050	II
Ecgonine	9180	II
Levorphanol	9220	II
Meperidine	9230	II
Methadone	9250	II
Morphine	9300	II
Thebaine	9333	II
Levo-Alphacetylmethadol (LAAM)	9648	II
Noroxymorphone	9668	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Fentanyl	9801	II

The company plans to manufacture reference standards.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–03760 Filed 2–23–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–796]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Livwell Michigan, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: The Drug Enforcement Administration (DEA) is providing notice of an application it has received from an entity applying to be registered to manufacture in bulk basic class(es) of controlled substances listed in schedule I. DEA intends to evaluate this and other pending applications according to its regulations governing the program of growing marihuana for scientific and medical research under DEA registration.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections to the issuance of the proposed registration on or before April 26, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW 8701 Morrissette Drive, Springfield, Virginia 22152. To ensure proper handling of comments, please reference Docket No. DEA–796 in all correspondence, including attachments.

SUPPLEMENTARY INFORMATION: The Controlled Substances Act (CSA) prohibits the cultivation and distribution of marihuana except by persons who are registered under the CSA to do so for lawful purposes. In accordance with the purposes specified in 21 CFR 1301.33(a), DEA is providing notice that the entity identified below has applied for registration as a bulk manufacturer of schedule I controlled substances. In response, registered bulk manufacturers of the affected basic class(es), and applicants therefor, may file written comments on or objections of the requested registration, as provided in this notice. This notice does not constitute any evaluation or determination of the merits of the application submitted.

The applicant plans to manufacture bulk active pharmaceutical ingredients (APIs) for product development and distribution to DEA-registered researchers. If the application for registration is granted, the registrant would not be authorized to conduct other activity under this registration aside from those coincident activities specifically authorized by DEA regulations. DEA will evaluate the application for registration as a bulk manufacturer for compliance with all applicable laws, treaties, and regulations and to ensure adequate safeguards against diversion are in place.

As this applicant has applied to become registered as a bulk manufacturer of marihuana, the application will be evaluated under the criteria of 21 U.S.C. 823(a). DEA will conduct this evaluation in the manner described in the rule published at 85 FR 82333 on December 18, 2020, and reflected in DEA regulations at 21 CFR part 1318.

In accordance with 21 CFR 1301.33(a), DEA is providing notice that on January 25, 2021, Livwell Michigan, LLC, 21550 Hoover Road, Warren, Michigan 48089, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

Controlled substance	Drug code	Schedule
Marihuana Extract	7350	I
Marihuana	7360	I
Tetrahydrocannabinols	7370	I

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021–03759 Filed 2–23–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–792]

Bulk Manufacturer of Controlled Substances Application: Synthcon LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Synthcon LLC, 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 April 26, 2021. Such persons may also file a written request for a hearing on the application on or before April 26, 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 January 5, 2021, Synthcon LLC., 770 Wooten Road, Suite 101, Colorado Springs, Colorado 80915–3538, applied to be registered as a bulk

manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
3-FMC	1233	I
Cathinone	1235	I
Methcathinone	1237	I
4-FMC	1238	I
Pentedrone	1246	I
Mephedrone (4-Methyl-N-methylcathinone)	1248	I
4-MEC	1249	I
Naphyrone	1258	I
N-Ethylamphetamine	1475	I
N,N-Dimethylamphetamine	1480	I
Aminorex	1585	I
Cis-4-Methylaminorex	1590	I
GHB	2010	I
Methaqualone	2565	I
Mecloqualone	2572	I
JWH-250	6250	I
ADB-PINACA	7035	I
JWH-018	7118	I
JWH-073	7173	I
JWH-200	7200	I
JWH-203	7203	I
4-Methyl-alpha-ethylaminopentiophenone	7245	I
N-Ethyhexedrone	7246	I
AET	7249	I
Ibogaine	7260	I
CP-47,497	7297	I
CP-47,497 C8 HOMOLOG	7298	I
LSD	7315	I
2C-T-7	7348	I
Tetrahydrocannabinols	7370	I
Mescaline	7381	I
2C-T-2	7385	I
3,4,5-TMA	7390	I
DOB	7391	I
2CB	7392	I
DOM	7395	I
2,5-DMA	7396	I
JWH-398	7398	I
DOE	7399	I
MDA	7400	I
5-METHOXY-MDA	7401	I
N-HYDROXY-MDA	7402	I
MDEA	7404	I
MDMA	7405	I
PMA	7411	I
5-MeO-DMT	7431	I
AMT	7432	I
Bufotenine	7433	I
DET	7434	I
DMT	7435	I
Psilocybin	7437	I
Psilocin	7438	I
5-Methoxy-N,N-diisopropyltryptamine	7439	I
4-Methyl-alpha-pyrrolidinohexiophenone	7446	I
PCE	7455	I
PCPy	7458	I
TCP	7470	I
TCPy	7473	I
JB323	7482	I
JB336	7484	I
BZP	7493	I
4-MePPP	7498	I
2C-D	7508	I
2C-E	7509	I
2C-H	7517	I
2C-I	7518	I
2C-C	7519	I
2C-N	7521	I
2C-P	7524	I
2C-T-4	7532	I
MDPV	7535	I

Controlled substance	Drug code	Schedule
25B-NBOME	7536	I
25C-NBOME	7537	I
25I-NBOME	7538	I
Methylone	7540	I
Butylone	7541	I
Pentylone	7542	I
N-Ethylpentyloe	7543	I
Alpha-Pyrrolidinohexanophenone	7544	I
Alpha-PVP	7545	I
Alpha-PBP	7546	I
Ethylone	7547	I
AM-694	7694	I
Pseudoephedrine	8112	I
Ephedrine	8113	I
Heroin	9200	I
Normorphine	9313	I
Acetorphine	9319	I
U-47700	9547	I
AH-7921	9551	I
MT-45	9560	I
Acetylmethadol	9601	I
Allylprodine	9602	I
Alphacetylmethadol	9603	I
Alphameprodine	9604	I
Alphamethadol	9605	I
Benzethidine	9606	I
Betacetylmethadol	9607	I
Clonitazine	9612	I
Isontonitazene	9614	I
Diampromide	9615	I
Diethylthiambutene	9616	I
Dimethylthiambutene	9619	I
Etonitazene	9624	I
Ketobemidone	9628	I
MPPP	9661	I
PEPAP	9663	I
Tilidine	9750	I
Acryl Fentanyl	9811	I
Para-fluorofentanyl	9812	I
3-Methylfentanyl	9813	I
Alpha-methylfentanyl	9814	I
Acetyl-alpha-methylfentanyl	9815	I
Ortho-fluorofentanyl	9816	I
Acetylfentanyl	9821	I
Butyrylfentanyl	9822	I
Para-fluorofentanyl	9823	I
Methoxyacetyl fentanyl	9825	I
4-Fluoroisobutyryl fentanyl (N-(4-fluorophenyl)-N-(1-phenethylpiperidin-4-yl)isobutyramide)	9824	I
Para-chloroisobutyryl fentanyl	9826	I
Isobutyrylfentanyl	9827	I
Beta-Hydroxyfentanyl	9830	I
Beta-Hydroxy-3-methylfentanyl	9831	I
Alpha-Methylthiofentanyl	9832	I
3-Methylthiofentanyl	9833	I
Furanylfentanyl	9834	I
Thiofentanyl	9835	I
Beta-Hydroxythiofentanyl	9836	I
Para-Methoxybutyryl Fetnanyl	9837	I
Ocfentanil	9838	I
Valeryl Fentanyl	9840	I
Tetrahydrofuranyl Fentanyl	9843	I
Crotonyl Fentanyl	9844	I
Cyclopropyl Fentanyl	9845	I
Cyclopentyl Fentanyl	9847	I
Fentanyl Related Compounds	9850	I
Amphetamine	1100	II
Methamphetamine	1105	II
1-Phenylcyclohexylamine	7460	II
PCP	7471	II
ANPP	8333	II
Norfentanyl	8366	II
P2P	8501	II
PCC	8603	II
Alphaprodine	9010	II

Controlled substance	Drug code	Schedule
Anileridine	9020	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
Levo-alphacetylmethadol	9648	II
Alfentanil	9737	II
Remifentanil	9739	II
Sufentanil	9740	II
Carfentanil	9743	II
Tapentadol	9780	II
Fentanyl	9801	II

The company plans to manufacture the above listed controlled substances as analytical reference 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.

William T. McDermott,

Assistant Administrator.

[FR Doc. 2021-03758 Filed 2-21; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–790]

Importer of Controlled Substances Application: Globyz Pharma, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Globyz Pharma, 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 March 26, 2021. Such persons may also file a written request for a hearing on the application on or before March 26, 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 September 30, 2020, Globyz Pharma, LLC, 2101 Market Street, Suite 5, Upper Chichester, Pennsylvania 19061–4001, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Oxycodone	9143	II

The company plans to import the listed controlled substance to complete analytical testing. 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. 2021–03834 Filed 2–23–21; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Mine Safety and Health Administration

Petitions for Modification of Application of Existing Mandatory Safety Standards

AGENCY: Mine Safety and Health Administration, Labor.

ACTION: Notice.

SUMMARY: This notice is a summary of a petition for modification submitted to the Mine Safety and Health Administration (MSHA) by the party listed below.

DATES: All comments on the petition must be received by MSHA's Office of Standards, Regulations, and Variances on or before March 26, 2021.

ADDRESSES: You may submit your comments, identified by ''docket number'' on the subject line, by any of the following methods:

1. *Electronic Mail: zzMSHA-comments@dol.gov.* Include the docket number of the petition in the subject line of the message.

2. *Facsimile:* 202–693–9441.

3. *Regular Mail or Hand Delivery:* MSHA, Office of Standards, Regulations, and Variances, 201 12th Street South, Suite 4E401, Arlington, Virginia 22202–5452, Attention: S. Aromie Noe, Acting Deputy Director, Office of Standards, Regulations, and Variances. Persons delivering documents are required to check in at the receptionist's desk in Suite 4E401. Individuals may inspect copies of the petition and comments during normal business hours at the address listed above.

MSHA will consider only comments postmarked by the U.S. Postal Service or proof of delivery from another delivery service such as UPS or Federal Express on or before the deadline for comments.