## SUMMARY:

Unither Manufacturing LLC has applied to be registered as an importer of basic class(es) of controlled substances. Refer to Supplemental 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 July 5, 2022. Such persons may also file a written request for a hearing on the application on or before July 5, 2022.

## 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](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](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. All requests for a hearing must be sent to: Federal Register Representative, Office of Management and Budget, Room 1016, 410 First Street, NE, Washington, DC 20503, Attention: Drug Enforcement Administration, Attn: DEA, Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative, Office of Management and Budget, Room 1016, 410 First Street, NE, Washington, DC 20503, Attention: DEA, Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152.

## SUPPLEMENTARY INFORMATION:

In accordance with 21 CFR 1301.27(a), this is notice that on April 8, 2022, Unither Manufacturing LLC, 331 Clay Road, Orange, New York 14623–3226, has applied to be registered as an importer of the following basic class(es) of controlled substance(s):

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methylphenidate</td>
<td>1724</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substance solely for commercial sale. Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Kristi O’Malley, 
Assistant Administrator.

[FR Doc. 2022–11824 Filed 6–1–22; 8:45 am]
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: 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 submit electronic 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 (API) 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 November 18, 2021, FPC Group LLC, 1601 South East 1st Street, Lawton, Oklahoma 73501, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substances:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Marihuana Extract</td>
<td>7350</td>
<td>I</td>
</tr>
<tr>
<td>Marihuana</td>
<td>7350</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols</td>
<td>7370</td>
<td>I</td>
</tr>
</tbody>
</table>

Kristi O’Malley, Assistant Administrator.

[FR Doc. 2022–11825 Filed 6–1–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Construction Standards on Posting Emergency Telephone Numbers and Floor Load Limits

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety & Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3506(c)(2)(A)). This program ensures that information is in the desired format, reporting burden (time and costs) is minimal, the collection instruments are clearly understood, and OSHA’s estimate of the information collection burden is accurate. The Occupational Safety and Health Act of 1970 (OSH Act) (29 U.S.C. 651 et seq.) authorizes information collection by employers as necessary or appropriate for enforcement of the OSH Act or for developing information regarding the causes and prevention of occupational injuries, illnesses, and accidents (29 U.S.C. 657). The OSH Act also requires that OSHA obtain such information with minimum burden upon employers, especially those operating small businesses, and to reduce to the maximum extent feasible unnecessary duplication of effort in obtaining information (29 U.S.C. 657).

Two construction standards, “Medical Services and First Aid” (§ 1926.50), and “General Requirements for Storage” (§ 1926.250), contain posting provisions. Paragraph (f) of § 1926.50 requires employers to conspicuously post emergency telephone numbers for physicians, hospitals, or ambulances at their worksites if 911 emergency telephone service is not locally available; in the event that a worker has a serious injury at a worksite, this posting requirement helps expedite emergency medical treatment of the worker. Paragraph (a)(2) of § 1926.250 specifies that employers must post the maximum safe load limits of floors located in storage areas inside buildings or other structures under construction, unless the floors or slabs are on grade (sitting on the ground). This provision prohibits employers from overloading floors in areas used to store material and equipment where a structure’s floors are not supported directly by the ground. This requirement is intended to prevent floor collapses which could seriously injure or kill workers. For additional substantive information about this ICR, see the related notice published in the Federal Register on March 2, 2022 (87 FR 11736).

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