DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-735]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Contract Pharmacal Corp

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 proposed regulations that, if finalized, would govern 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 [INSERT 60 DAYS AFTER PUBLICATION IN THE FEDERAL REGISTER].

ADDRESS: 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-735 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 proposes to conduct this evaluation in the manner described in the rule proposed at 85 FR 16292, published on March 23, 2020, if finalized.

In accordance with 21 CFR 1301.33(a), DEA is providing notice that on August 21, 2020, Contract Pharmacal Corp., 165 Oser Avenue, Hauppauge, New York 11788, 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>7360</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols</td>
<td>7370</td>
<td>I</td>
</tr>
</tbody>
</table>
The applicants notice above applied to become registered with DEA to grow marihuana as a bulk manufacturer subsequent to a 2020 DEA notice of proposed rulemaking that provided information on how DEA intends to expand the number of registrations and described the way it would oversee those additional growers. If finalized, the proposed rule would govern persons seeking to become registered with DEA to grow marihuana as a bulk manufacturer, consistent with applicable law. The notice of proposed rulemaking is available at 85 FR 16292.

William T. McDermott,
Assistant Administrator.
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