On December 16, 2020, the Commission found a violation of section 337 based on the misappropriation of Complainants’ trade secrets (including the Medytox manufacturing processes but not the Medytox bacterial strain). See 85 FR 83610–11 (Dec. 22, 2020). The Commission issued a limited exclusion order (“LEO”) against certain botulinum neurotoxin products that are imported and/or sold by Respondents Daewoong and Evolus and a cease and desist order (“CDO”) against Evolus. Id. The Commission also set a bond during the period of Presidential review in an amount of $441 per 100U vial of Respondents’ accused products. Id.

On February 12, 2021, Complainants filed an appeal from the Commission’s final determination with the Federal Circuit. On the same day, Respondents also filed an appeal from the Commission’s final determination of a violation of section 337. On February 18, 2021, Complainants and Evolus (collectively, “the Settling Parties”) announced that they had reached a settlement agreement to resolve all pending issues between them.

On March 3, 2021, the Settling Parties filed a joint petition to rescind the LEO and CDO (collectively, “the remedial orders”) based on the settlement agreement. On the same day, the Settling Parties also filed a joint motion to limit service of the settlement agreement. On March 16, 2021, Daewoong filed a notice of non-opposition to the joint motion to limit service. On April 1, 2021, the Settling Parties further filed a joint motion to terminate the investigation without prejudice pursuant to 19 CFR 210.21(b).

However, Commissioner Karpel would decline to issue an indicative ruling as to whether Daewoong has established equitable entitlement to the extraordinary remedy of vacatur on the basis of the record before the Commission.

The rescission proceeding is terminated.

The Commission’s vote on this determination took place on May 3, 2021.

The authority for the Commission’s determination is contained in section 337 of the Tariff Act of 1930, as amended (19 U.S.C. 1337), and in part 210 of the Commission’s Rules of Practice and Procedure (19 CFR part 210).


Lisa Barton,
Secretary to the Commission.

[FR Doc. 2021–09652 Filed 5–6–21; 8:45 am]

BILLING CODE 7020–02–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

[Docket No. DEA–814]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Indian Flower 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 July 6, 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–814 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 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</td>
<td>7360</td>
<td>I</td>
</tr>
</tbody>
</table>

William T. McDermott, Assistant Administrator.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration
[Docket No. DEA–832]

Bulk Manufacturer of Controlled Substances Application: Bulk Manufacturer of Marihuana: Green Rush Organic Farms Inc.

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 July 6, 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–832 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 March 31, 2021, Green Rush Organic Farms Inc., 1318 South Kilbourn Avenue, Chicago, Illinois 60623, 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>
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<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols</td>
<td>7370</td>
<td>I</td>
</tr>
</tbody>
</table>

William T. McDermott, Assistant Administrator.

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