DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated August 11, 2011, and published in the Federal Register on August 18, 2011, 76 FR 51398, Cambrex Charles City, Inc., 1205 11th Street, Charles City, Iowa 50616–3466, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Schedule</th>
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</thead>
<tbody>
<tr>
<td>Phenyacetone (8501)</td>
<td>II</td>
</tr>
<tr>
<td>Opium, raw (9600)</td>
<td>II</td>
</tr>
<tr>
<td>Poppy Straw Concentrate (9670)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances to manufacture a bulk intermediate for sale to its customers. With regard to the Phenyacetone, the company plans to use it as a base material in the bulk manufacture of another controlled substance.

No comments or objections have been received regarding Phenyacetone.

Comments and requests for hearings on applications to import narcotic raw material are not appropriate, 72 FR 3417 (2007). DEA has considered the factors in 21 U.S.C. 823(a) and 952(e) and determined that the registration of Cambrex Charles City, Inc. to import the basic classes of controlled substances is consistent with the public interest.

The investigation has included inspection and testing of the company’s physical security systems, verification of the company’s compliance with state and local laws, and a review of the company’s background and history. Therefore, pursuant to 21 U.S.C. 952(a) and 958(a), and in accordance with 21 CFR 1301.34, the above named company is granted registration as an importer of controlled substances listed.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.

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DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a), Title 21 of the Code of Federal Regulations (CFR), this is notice that on December 15, 2011, Pharmagra Labs, Inc., 158 McLean Road, Brevard, North Carolina 28712, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Pentobarbital (2270), a basic class of controlled substance listed in schedule II.

The company plans to manufacture the listed substance for analytical research and clinical trials.

Any other such applicant, and any person who is presently registered with DEA to manufacture such substance, may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODL), 8701 Morrissette Drive, Springfield, Virginia 22152; and must be filed no later than April 6, 2012.