DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Bulk Manufacturer of Controlled Substances Application: Chemtos, LLC

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic class, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before July 31, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA) to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on April 6, 2017, Chemtos Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665–2402 applied to be registered as an importer of U–47700 (3,4-dichloro-N-[2-dimethylamino)cyclohexyl]-N-methylbenzamide) (9547), a basic class of controlled substance listed in schedule I. The company plans to import small quantities of the listed controlled substance for the manufacture of analytical reference standards and distribution to their research and forensic customers.


Louis J. Milione,
Assistant Administrator.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances Application: Cerilliant Corporation

ACTION: Notice of application.

DATES: Registered bulk manufacturers of the affected basic classes, and applicants therefore, may file written comments on or objections to the issuance of the proposed registration in accordance with 21 CFR 1301.33(a) on or before July 3, 2017. Such persons may also file a written request for a hearing on the application pursuant to 21 CFR 1301.43 on or before July 3, 2017.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DRW 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/LJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DRW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: The Attorney General has delegated his authority under the Controlled Substances Act to the Administrator of the Drug Enforcement Administration (DEA) to exercise all necessary functions with respect to the promulgation and implementation of 21 CFR part 1301, incident to the registration of manufacturers, distributors, dispensers, importers, and exporters of controlled substances (other than final orders in connection with suspension, denial, or revocation of registration) has been redelegated to the Assistant Administrator of the DEA Diversion Control Division (“Assistant Administrator”) pursuant to section 7 of 28 CFR part 0, appendix to subpart R.

In accordance with 21 CFR 1301.34(a), this is notice that on April 6, 2017, Cerilliant Corporation, 811 Paloma Drive, Suite A, Round Rock, Texas 78665–2402 applied to be registered as an importer of U–47700 (3,4-dichloro-N-[2- dimethylamino)cyclohexyl]-N-methylbenzamide) (9547), a basic class of controlled substance listed in schedule I. The company plans to import small quantities of the listed controlled substance for the manufacture of analytical reference standards and distribution to their research and forensic customers.


Louis J. Milione,
Assistant Administrator.

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances Registration

ACTION: Notice of registration.

SUMMARY: Registrants listed below have applied for and been granted registration by the Drug Enforcement Administration (DEA) as importers of various basic classes of controlled substances.

SUPPLEMENTARY INFORMATION: The companies listed below applied to be registered as importers of various basic classes of controlled substances. Information on previously published notices is listed in the table below. No comments or objections were submitted and no requests for hearing were submitted for these notices.

<table>
<thead>
<tr>
<th>Company</th>
<th>FR docket</th>
<th>Published</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chattem Chemicals, Inc</td>
<td>82 FR 13135</td>
<td>March 9, 2017.</td>
</tr>
<tr>
<td>Anderson Brecon, Inc</td>
<td>82 FR 13134</td>
<td>March 9, 2017.</td>
</tr>
<tr>
<td>Hospira</td>
<td>82 FR 11241</td>
<td>February 21, 2017.</td>
</tr>
<tr>
<td>Myoderm</td>
<td>82 FR 11241</td>
<td>February 21, 2017.</td>
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<tr>
<td>Meridian Medical Technologies</td>
<td>82 FR 11241</td>
<td>February 21, 2017.</td>
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