SUMMARY: Mylan Technologies, Inc., applied to be registered as an importer of certain basic classes of controlled substances. The DEA grants Mylan Technologies, Inc., registration as an importer of those controlled substances.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the Federal Register on June 4, 2014, 79 FR 32316, Mylan Technologies, Inc., 110 Lake Street, Saint Albans, Vermont 05478, applied to be registered as an importer of certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Mylan Technologies, Inc., to import the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing 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 the basic classes of controlled substances listed:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methylphenidate (1724)</td>
<td>II</td>
</tr>
<tr>
<td>Fentanyl (9801)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company’s own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

DEPARTMENT OF JUSTICE
Drug Enforcement Administration
[Docket No. DEA–392]
Importer of Controlled Substances
Registration: Arizona Department of Corrections, Aspc-Florence

ACTION: Notice of registration.

SUMMARY: Arizona Department of Corrections, ASPC-Florence applied to be registered as an importer of a certain basic class of controlled substance. The DEA grants Arizona Department of Corrections, ASPC-Florence registration as an importer of this controlled substance.

SUPPLEMENTARY INFORMATION: By notice dated May 28, 2014, and published in the Federal Register on June 4, 2014, 79 FR 32317, Arizona Department of Corrections, ASPC-Florence, 1305 E. Butte Avenue, Florence, Arizona 85132, applied to be registered as an importer of a certain basic class of controlled substance. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Arizona Department of Corrections, ASPC-Florence to import the basic class of controlled substance is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing 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 certain basic classes of controlled substances. No comments or objections were submitted for this notice.

The Drug Enforcement Administration (DEA) has considered the factors in 21 U.S.C. 823, 952(a) and 958(a) and determined that the registration of Wildlife Laboratories, Inc., to import the basic classes of controlled substances is consistent with the public interest and with United States obligations under international treaties, conventions, or protocols in effect on May 1, 1971. The DEA investigated the company’s maintenance of effective controls against diversion by inspecting and testing the company’s physical security systems, verifying the company’s compliance with state and local laws, and reviewing 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 the basic classes of controlled substances:

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Etorphine (except HCl) (9056)</td>
<td>I</td>
</tr>
<tr>
<td>Etorphine HCl (9059)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances for sale to its customers.

Dated: August 27, 2014.

Joseph T. Rannazzisi,
Deputy Assistant Administrator.

[FR Doc. 2014–21087 Filed 9–3–14; 8:45 am]