Pursuant to 21 U.S.C. 958(j), the Attorney General shall, prior to issuing a registration under this Section to a bulk manufacturer of a controlled substance in schedule I or II, and prior to issuing a registration under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for the bulk manufacture of the substance an opportunity for a hearing.

Therefore, in accordance with 21 CFR 1301.34(a), this is notice that on November 5, 2009, Mylan Technologies Inc., 110 Lake Street, Saint Albans, Vermont 05478, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the basic classes of controlled substances listed in schedule II:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Methylphenidate</td>
<td>II</td>
</tr>
<tr>
<td>Fentanyl</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances for analytical research and clinical trials.

Any bulk manufacturer who is present in the area of the DEA to manufacture such a basic class of controlled substance may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43, and in such form as prescribed by 21 CFR 1316.47.

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 February 3, 2010.

This procedure is to be conducted simultaneously with, and independent of, the procedures described in 21 CFR 1301.34(b), (c), (d), (e), and (f). As noted in a previous notice published in the Federal Register on September 23, 1975, (40 FR 43745–46), all applicants for registration to import a basic class of any controlled substance in schedule I or II are, and will continue to be, required to demonstrate to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, that the requirements for such registration pursuant to 21 U.S.C. 958(a); 21 U.S.C. 823(a); and 21 CFR 1301.34(b), (c), (d), (e), and (f) are satisfied.

Dated: December 17, 2009.

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

BILLING CODE 4410–09–P

OFFICE OF NATIONAL DRUG CONTROL POLICY

Paperwork Reduction Act; 30-Day Notice

AGENCY: Office of National Drug Control Policy.

The Office of National Drug Control Policy (ONDCP) proposes the collection of information concerning arrestee drug use. ONDCP invites interested persons to submit comments to the Office of Management and Budget (OMB) regarding any aspect of this proposed effort.

Type of Information Collection: New collection.

Title: Arrestee Drug Abuse Monitoring (ADAM II) Program Questionnaire.

Use: The information will support statistical trend analysis.

Frequency: Ten sites will each conduct two cycles of surveys from 250 arrestees per cycle.

Annual Number of Respondents: 5000.

Total Annual Responses: 5000.

Average Burden per Response: 25 minutes.

Total Annual Hours: 2,083.

Send comments to John Kraemer, OMB Desk Officer for ONDCP, New Executive Office Building, Room 10235, Washington, DC 20503. Comments must be received within 30 days. Request additional information by facsimile transmission to (202) 395–6562, attention: Robert Cohen, ONDCP, Office of Research and Data Analysis.


Daniel R. Petersen,
Deputy General Counsel.

BILLING CODE 3180–02–P

NUCLEAR REGULATORY COMMISSION

[FR Doc. E9–31165 Filed 12–31–09; 8:45 am]

In the Matter of: Certain Licensees Requesting Unescorted Access to Radioactive Material; Order Imposing Trustworthiness and Reliability Requirements for Unescorted Access to Certain Radioactive Material (Effective Immediately)

I

The Licensees identified in Attachment 1 to this Order hold licenses issued in accordance with the Atomic Energy Act (AEA) of 1954, as amended, by the U.S. Nuclear Regulatory Commission (NRC or Commission) or an Agreement State, authorizing them to perform services on devices containing certain radioactive material for customers licensed by the NRC or an Agreement State to possess and use certain quantities of the radioactive materials listed in Attachment 2 to this Order. Commission regulations at 10 CFR 20.1801 or equivalent Agreement State regulations require Licensees to secure, from unauthorized removal or access, licensed materials that are stored in controlled or unrestricted areas. Commission regulations at 10 CFR 20.1802 or equivalent Agreement State regulations require Licensees to control and maintain constant surveillance of licensed material that is in a controlled or unrestricted area and that is not in storage.

II

Subsequent to the terrorist events of September 11, 2001, the NRC issued immediately effective security Orders to NRC and Agreement State Licensees under the Commission’s authority to protect the common defense and security of the nation. The Orders required certain manufacturing and distribution (M&D) Licensees to implement Additional Security Measures (ASMs) for the radioactive materials listed in Attachment 2 to this Order (the radionuclides of concern), to supplement the existing regulatory requirements. The ASMs included requirements for determining the trustworthiness and reliability of individuals that require unescorted access to the radionuclides of concern. Section 652 of the Energy Policy Act of 2005, which became law on August 8, 2005, amended Section 149 of the AEA to require fingerprinting and a Federal Bureau of Investigation (FBI)