proposed collection of information, including the validity of methodology and assumptions used;
(3) Enhance the quality, utility, and clarity of the information to be collected; and,
(4) Minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of this Information Collection:
(1) Type of Information Collection: Reinstatement, with no change, of a previously approved collection for which approval has expired.
(2) Title of Form/Collection: International Terrorism Victim Expense Reimbursement Program (ITVERP) Application.
(3) Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: Form Number: The Office of Management and Budget Number for the certification form is 1121–0309. The Office for Victims of Crime, Office of Justice Programs, United States Department of Justice is sponsoring the collection.
(4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: The form is completed by U.S. nationals and U.S. government employees who become victims of acts of international terrorism that occur outside the United States. Applicants seeking compensation from OVC for expenses associated with their victimization will be required to submit said form. The form will be used to collect necessary information on expenses incurred by the applicant, as well as other pertinent information, and will be used by OVC to make an award determination.
(5) An estimate of the total number of respondents and the amount of time estimated for an average to respond: The total hour burden to complete the forms is 1,500 annual burden hours.
(6) An estimate of the total public burden (in hours) associated with the collection: There are approximately 1,500 hours annual burden associated with this information collection.

If additional information is required contact: Lynn Bryant, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, United States Department of Justice, 601 D Street NW., Patrick Henry Building, Suite 1600, NW., Washington, DC 20530.

Dated: March 18, 2010.

Lynn Bryant, Department Clearance Officer, United States Department of Justice.

DEPARTMENT OF JUSTICE
Office of Justice Programs
[OMB Number 1121–0220]
Agency Information Collection Activities: Extension of a Currently Approved Collection: Comments Requested


The Department of Justice (DOJ), Office of Justice Programs (OJP) will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed information collection is published to obtain comments from the public and affected agencies. The proposed information collection was previously published in the Federal Register Volume 75, Number 8, page 1811 on January 13, 2010, allowing for a 60 day comment period.
The purpose of this notice is to allow for an additional 30 days for public comment until April 23, 2010. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to the Office of Management and Budget, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395–5806. Comments may also be submitted to M. Berry, Bureau of Justice Assistance, Office of Justice Programs, U. S. Department of Justice, 810 7th Street, NW., Washington, DC 20531 via facsimile to (202) 305–1367 or by e-mail at M.A.Berry@ojp.usdoj.gov.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

—Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility;
—Evaluate the accuracy of the agency’s estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;
—Enhance the quality, utility, and clarity of the information to be collected; and,
—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection:
(1) Type of Information Collection: Extension of a currently approved collection.
(2) Title of the Form/Collection: Public Safety Officers’ Educational Assistance.
(3) Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection: None.
(4) Affected public who will be asked or required to respond, as well as a brief abstract: Primary: Dependent spouses and/or children of public safety officers who were killed or permanently and totally disabled in the line of duty.

Abstract: BJA’s Public Safety Officers’ Benefits (PSOB) Office will use the PSOA application information to confirm the eligibility of applicants to receive PSOA benefits. Eligibility is dependent on several factors, including the applicant having received or being eligible to receive a portion of the PSOB death benefit, or having a family member who received the PSOB disability benefit. Also considered are the applicant’s age and the schools being attended. In addition, information to help BJA identify an individual is collected, such as Social Security Number and contact numbers and e-mail addresses. The changes to the application form have been made in an effort to streamline the application process and eliminate requests for information that is either irrelevant or already being collected by other means.

Others: None.
(5) An estimate of the total number of respondents and the amount of time needed for an average respondent to
respond as follows: It is estimated that no more than 100 new respondents will apply a year. Each application takes approximately 20 minutes to complete.

[6] An estimate of the total public burden (in hours) associated with the collection is 33 hours. Total Annual Reporting Burden: 100 x 20 minutes per application = 2000 minutes/by 60 minutes per hour = 33 hours.

If additional information is required, please contact, Lynn Bryant, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Patrick Henry Building, Suite 1600, 601 D Street, NW., Washington, DC 20530.

Dated: March 18, 2010.

Lynn Bryant,
Department Clearance Officer, PRA, United States Department of Justice.

[FR Doc. 2010–6441 Filed 3–23–10; 8:45 am]
BILLING CODE 4410–18–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application

Pursuant to 21 U.S.C. 958(f), 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 regulation under 21 U.S.C. 952(a)(2) authorizing the importation of such a substance, provide manufacturers holding registrations for such basic classes of controlled substances an opportunity for a hearing.

Therefore, in accordance with Title 21 Code of Federal Regulations (CFR), 1301.34(a), this is notice that on January 21, 2010, Roche Diagnostics Operations Inc., Attn: Regulatory Compliance, 9115 Hague Road, Indianapolis, Indiana 46250, 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 schedules I and II:

<table>
<thead>
<tr>
<th>Drug</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lysergic acid diethylamide (7315)</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols (7370)</td>
<td>I</td>
</tr>
<tr>
<td>Alphamethadol (9605)</td>
<td>I</td>
</tr>
<tr>
<td>Cocaine (9041)</td>
<td>II</td>
</tr>
<tr>
<td>Ecgonine (9180)</td>
<td>II</td>
</tr>
<tr>
<td>Methadone (9250)</td>
<td>II</td>
</tr>
<tr>
<td>Morphine (9300)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances for the manufacture of diagnostic products for distribution to its customers.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances 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 April 23, 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.


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

[FR Doc. 2010–6441 Filed 3–23–10; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By a Notice dated June 24, 2009, and published in the Federal Register on July 9, 2009 (74 FR 32954) and by a second Notice (Correction) dated August 21, 2009, and published in the Federal Register on September 8, 2009 (74 FR 46229), Rhodes Technologies, 498 Washington Street, Coventry, Rhode Island 02816, 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>Raw Opium (9600)</td>
<td>II</td>
</tr>
<tr>
<td>Concentrate of Poppy Straw (9670)</td>
<td>II</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances to manufacture bulk active pharmaceutical ingredients.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and §952(a), and determined that the registration of Rhodes Technologies 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, at this time. DEA has investigated Rhodes Technologies to ensure that the company’s registration 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 the basic classes of controlled substances listed.


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

[FR Doc. 2010–6441 Filed 3–23–10; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated November 23, 2009, and published in the Federal Register on December 3, 2009 (74 FR 63411), Noramco, Inc., Division of Ortho-McNeil, Inc., 500 Swedes Landing Road, Wilmington, Delaware 19801, 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>Opium, raw (9600)</td>
<td>II</td>
</tr>
<tr>
<td>Poppy Straw Concentrate (9670)</td>
<td>III</td>
</tr>
</tbody>
</table>