

Alexandria, Virginia 22301; and must be filed no later than November 17, 2006.

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: October 11, 2006.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
[FR Doc. E6–17293 Filed 10–17–06; 8:45 am]
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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated June 7, 2006, and published in the **Federal Register** on June 13, 2006 (71 FR 34162), 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 class of controlled substances listed in Schedule I and II:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Cocaine (9041)	II
Ecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II
Alphamethadol (9605)	II

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

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 Roche Diagnostics Operations, Inc. to import the basic class 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 Roche Diagnostics Operations, Inc. 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 class of controlled substances listed.

Dated: October 11, 2006.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
[FR Doc. E6–17289 Filed 10–17–06; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to Section 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 27, 2006, Varian, Inc., Lake Forest, 25200 Commercentre Drive, Lake Forest, California 92630–8810, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the basic classes of controlled substances listed in Schedule II:

Drug	Schedule
Phencyclidine (7471)	II
1-Piperidinocyclohexanecarbonitrile (8603).	II
Benzoylcegonine (9180)	II

The company plans to manufacture small quantities of the listed controlled substances for use in diagnostic products.

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

Any such written comments or objections being sent via regular mail should be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration,

Washington, DC 20537, *Attention: DEA Federal Register Representative/ODL*; or any being sent via express mail should be sent to DEA Headquarters, *Attention: DEA Federal Register Representative/ODL*, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than December 18, 2006.

Dated: October 11, 2006.

Joseph T. Rannazzisi,
Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration.
[FR Doc. E6–17277 Filed 10–17–06; 8:45 am]
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DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

October 13, 2006.

The Department of Labor (DOL) has submitted the following public information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chapter 35). A copy of this ICR, with applicable supporting documentation, may be obtained by contacting Ira Mills at the Department of Labor on 202–693–4122 (this is not a toll-free number) or E-Mail: Mills.Ira@dol.gov. This ICR can also be accessed online at <http://www.doleta.gov/OMBControlNumber.cfm>. ICRs are filed according to the date the 60-day Notice for Public Comment was published in the **Federal Register** Notice; therefore, on the left hand side of this site, look under July 25, 2006 to access 1205–0441.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ETA, Office of Management and Budget, Room 10235, Washington, DC 20503, 202–395–7316 (this is not a toll free number), within 30 days from the date of this publication in the **Federal Register**.

The OMB is particularly interested in comments which:

- 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.

Agency: Employment and Training Administration (ETA).

Type of Review: Extension of a currently approved collection.

Title: Evaluation of the Individual Training Account Experiment.

OMB Number: 1205-0441.

Frequency: One time, follow-up, as needed.

Affected Public: Individuals or households.

Type of Response: Reporting.

Number of Respondents: 3,840.

Annual Responses: 3,840.

Average Response time: 30 minutes.

Total Annual Burden Hours: 1,920.

Total Annualized Capital/Startup Costs: 0.

Total Annual Costs (operating/maintaining systems or purchasing services): 0.

Description: This clearance package seeks approval for the extension of the currently approved follow-up survey for the Individual Training Account (ITA) Experiment. The experiment is designed to test three different approaches to providing ITAs. Data from the follow-up survey of ITA customers will be used by Mathematica Policy Research, Inc. to describe experiences inside the workforce system and labor market outcomes for ITA customers. Measures of these experiences and outcomes would be used to further evaluate the three approaches. Based on information from the survey and other data sources, the U.S. Department of Labor can provide information to local workforce boards on how to administer their ITA programs.

Ira L. Mills,

Departmental Clearance Officer, Team Leader.

[FR Doc. E6-17356 Filed 10-17-06; 8:45 am]

BILLING CODE 4510-30-P

FEDERAL COUNCIL ON THE ARTS AND THE HUMANITIES

Arts And Artifacts Indemnity Panel Advisory Committee; Notice of Meeting

Pursuant to the provisions of the Federal Advisory Committee Act (Pub. L. 92-463 as amended) notice is hereby given that a meeting of the Arts and Artifacts Indemnity Panel of the Federal Council on the Arts and the Humanities will be held at 1100 Pennsylvania Avenue, NW., Washington, DC 20506, in Room 716, from 9 a.m. to 5 p.m., on Monday, November 6, 2006.

The purpose of the meeting is to review applications for Certificates of Indemnity submitted to the Federal Council on the Arts and the Humanities for exhibitions beginning after January 1, 2007.

Because the proposed meeting will consider financial and commercial data and because it is important to keep values of objects, methods of transportation and security measures confidential, pursuant to the authority granted me by the Chairman's Delegation of Authority to Close Advisory Committee Meetings, dated July 19, 1993, I have determined that the meeting would fall within exemption (4) of 5 U.S.C. 552(b) and that it is essential to close the meeting.

It is suggested that those desiring more specific information contact Acting Advisory Committee Management Officer, Heather Gottry, 1100 Pennsylvania Avenue, NW., Washington, DC 20506, or call 202-606-8322.

Heather Gottry,

Acting Advisory Committee Management Officer.

[FR Doc. E6-17362 Filed 10-17-06; 8:45 am]

BILLING CODE 7536-01-P

NUCLEAR REGULATORY COMMISSION

[Docket No. 50-482]

Wolf Creek Nuclear Operating Corporation; Notice of Receipt and Availability of Application for Renewal of Wolf Creek Generating Station, Unit 1 Facility Operating License No. NPF-42 for an Additional 20-Year Period

The U.S. Nuclear Regulatory Commission (NRC or Commission) has received an application, dated September 27, 2006, from Wolf Creek Nuclear Operating Corporation, filed pursuant to Section 103 of the Atomic Energy Act of 1954, as amended, and Title 10 of the *Code of Federal*

Regulations Part 54 (10 CFR Part 54), to renew the operating license for the Wolf Creek Generating Station (WCGS), Unit 1. Renewal of the license would authorize the applicant to operate the facility for an additional 20-year period beyond the period specified in the current operating license. The current operating license for WCGS, Unit 1, (NPF-42), expires on March 11, 2025. WCGS, Unit 1, is a pressurized water reactor designed by Westinghouse Electric Corporation that is located near Burlington, KS. The acceptability of the tendered application for docketing, and other matters including an opportunity to request a hearing, will be the subject of subsequent **Federal Register** notices.

Copies of the application are available to the public at the Commission's Public Document Room (PDR), located at One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852 or through the internet from the NRC's Agencywide Documents Access and Management System (ADAMS) Public Electronic Reading Room under Accession Number ML062770300. The ADAMS Public Electronic Reading Room is accessible from the NRC Web site at <http://www.nrc.gov/reading-rm/adams.html>. In addition, the application is available at <http://www.nrc.gov/reactors/operating/licensing/renewal/applications.html>. Persons who do not have access to the internet or who encounter problems in accessing the documents located in ADAMS should contact the NRC's PDR reference staff at 1-800-397-4209, extension 4737, or by e-mail to pdr@nrc.gov.

A copy of the license renewal application for the WCGS, Unit 1, is also available to local residents near the site at the Burlington Library, 410 Juniatta, Burlington, KS 66839.

Dated at Rockville, Maryland, this 12th day of October, 2006.

For the Nuclear Regulatory Commission.

Pao-Tsin Kuo,

Deputy Director, Division of License Renewal, Office of Nuclear Reactor Regulation.

[FR Doc. E6-17323 Filed 10-17-06; 8:45 am]

BILLING CODE 7590-01-P

POSTAL SERVICE

Privacy Act of 1974, Computer Matching Program—Postal Service and State Agencies

AGENCY: Postal Service.

ACTION: Notice of Computer Matching Program—Postal Service and states maintaining public sex offender registries.