Dated: June 24, 2002.

Laura M. Nagel,

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

[FR Doc. 02–17213 Filed 7–9–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on March 5, 2002, Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370)	I
Alphamethadol (9605)	I
Phencyclidine (7471)	II
Benzoylecgonine (9180)	II
Methadone (9250)	II
Morphine (9300)	II

Roche Diagnostics Corporation plans to manufacture small quantities of the above listed controlled substances for incorporation in drug of abuse detection kits.

Any other such applicant and any person who is presently registered with DEA to manufacture such substance may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than September 9, 2002.

Dated: June 18, 2002.

Laura M. Nagel,

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

[FR Doc. 02–17207 Filed 7–10–02; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importation of Controlled Substances; Notice of Application

Pursuant to section 1008 of the Controlled Substances Import and Export Act (21 U.S.C. 958(i)), 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 section 1002(a) 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 § 1301.34 of Title 21, Code of Federal Regulations (CFR), notice is hereby given that on March 5, 2002, Roche Diagnostics Corporation, 9115 Hague Road, Indianapolis, Indiana 46250, made application by renewal to the Drug Enforcement Administration to be registered as an importer of the basic classes of controlled substances listed below:

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

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

Any manufacturer holding, or applying for, registration as a bulk manufacturer of this basic class of controlled substance may file written comments on or objections to the application described above and may, at the same time, file a written request for a hearing on such application in accordance with 21 CFR 1301.43 in such form as prescribed by 21 CFR 1316.47.

Any such comments, objections or requests for a hearing may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (30 days from publication).

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 at 40 FR 43745–46 (September 23, 1975), 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(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: June 24, 2002.

Laura M. Nagel,

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

[FR Doc. 02–17210 Filed 7–9–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Application

Pursuant to § 1301.33(a) of Title 21 of the Code of Federal Regulations (CFR), this is notice that on February 11, 2002, Stepan Company, Natural Products Department, 100 W. Hunter Avenue, Maywood, New Jersey 07607, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of the basic classes of controlled substances listed below:

Drug	Schedule
Cocaine (9041)	II
Benzoylecgonine (9180)	II

The firm plans to manufacture bulk controlled substances for distribution to its customers.

Any other such applicant and any person who is presently registered with DEA to manufacture such substances may file comments or objections to the issuance of the proposed registration.

Any such comments or objections may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, United States Department of Justice, Washington, DC 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than (60 days from publication).

Dated: June 24, 2002.

Laura M. Nagel,

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

[FR Doc. 02–17209 Filed 7–9–02; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review; Comment Request

July 1, 2002.

The Department of Labor (DOL) has submitted the following public information collection requests (ICRs) 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 each individual ICR, with applicable supporting documentation, may be obtained by calling the Department of Labor. To obtain documentation contact Marlene Howze at ((202) 693–4158 or Email *Howze-Marlene@dol.gov.*

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for ESA, Office of Management and Budget, Room 10235, Washington, DC 20503 ((202) 395–7316), 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.

Type of Review: Revision of a currently approved collection.

Agency: Employment Standards Administration (ESA).

Title: FECA Medical Report Forms, Claim for Compensation.

OMB Number: 1215–0103.

Affected Public: Business or other for-profit; Individuals or households; and Federal Government.

Frequency: As Needed.

Number of Respondents: 286,010.

Number of Annual Responses: 286,010.

Estimated Response Times and Total Burden Hours:

Form No.	Number of respondents	Average minutes per response	Total burden hours
CA–7	400	13	87
CA–16B	130,000	5	10,833
CA–17B	60,000	5	5,000
CA–20	65,000	5	5,417
CA–1090	200	10	33
CA–1303	2,000	20	6667
CA–1305	10	20	3
CA–1331	200	5	17
CA–1332	200	30	100
QCM Letters	1,000	5	83
OWCP–5A	7,000	15	1,750
OWCP–5B	5,000	15	1,250
OWCP–5C	15,000	15	3,750
Burden Totals	286,010	163	28,990

Total Annualized Capital/Startup Costs: $0.

Total Annual Costs (operating/maintaining systems or purchasing services): $105,824.00.

Description: The Office of Workers' Compensation Programs administers the Federal Employees' Compensation Act (5 U.S.C. 8101, *et seq*). The statute provides for continuation of benefits for wage loss and/or for permanent impairment to a scheduled member, arising out of a work related injury or disease. The Act outlines the elements of pay which are to be included in an individual's pay rate, and sets forth various other criteria for determining eligibility to and the amount of benefits, including augmentation of basic compensation for individuals with qualifying dependents; a requirement to report any earnings during a period that compensation is claimed; a prohibition against concurrent receipt of FECA benefits and benefits from OPM or certain VA benefits; and a mandate that money collected from a liable third party found responsible for the injury for which compensation has been paid be applied to benefits paid or payable. The CA 7 is used to claim compensation and the other forms in this clearance collect medical information necessary to determine entitlement to benefits under the FECA. Without the requested information, an eligible beneficiary could be denied benefits, or benefits could be authorized at an incorrect rate, resulting in an underpayment or overpayment of compensation.

Ira L. Mills,

Departmental Clearance Officer.

[FR Doc. 02–17320 Filed 7–9–02; 8:45 am]

BILLING CODE 4510–CH–M

DEPARTMENT OF LABOR

Employment and Training Administration

Labor Surplus Area Classifications Under Executive Orders 12073 and 10582

ACTION: Notice of additions to the labor surplus area list.

DATE: June 3, 2002.