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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than November 17, 1999.

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 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 1311.42(a), (b), (c), (d), (e), and (f) are satisfied.

Dated: October 8, 1999.

John H. King,

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

[FR Doc. 99–27102 Filed 10–15–99; 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 22, 1999, Lifepoint, Inc., 10410 Trademark Street, Rancho Cucamonga, California 91730, 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
Tetrahydrocannabinols (7370)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Phencyclidine (7471)	II
Benzoylecgonine (9180)	II
Morphine (9300)	II

The firm plans to use gram quantities of the listed controlled substances to manufacture drug abuse test kits.

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 November 17, 1999.

Dated: October 8, 1999.

John H. King,

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

[FR Doc. 99–27103 Filed 10–15–99; 8:45 am]

BILLING CODE 4410–09–M

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 August 19, 1999, Nycomed, Inc., 33 Riverside Avenue, Rensselaer, New York 12144, 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
Methylphenidate (1724)	II
Meperidine (9230)	II

The firm plans to manufacture meperidine as bulk product for distribution to its customers and to manufacture methylphenidate for qualification and distribution to a customer.

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 17, 1999.

Dated: October 8, 1999.

John H. King,

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

[FR Doc. 99–27104 Filed 10–15–99; 8:45 am]

BILLING CODE 4410–01–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 August 3, 1999, Pharmacia & Upjohn Company, 7000 Portage Road, 2000–41–109, Kalamazoo, Michigan 49001, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of 2,5-Dimethoxyamphetamine (7396), a basic class of controlled substance listed in Schedule I.

The firm plans to manufacture the controlled substance for distribution as bulk product to a customer.

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, D.C. 20537, Attention: DEA Federal Register Representative (CCR), and must be filed no later than December 17, 1999.

Dated: October 8, 1999.

John H. King,

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

[FR Doc. 99–27105 Filed 10–15–99; 8:45 am]

BILLING CODE 4410–01–M

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Registration

By Notice dated March 3, 1999, and published in the **Federal Register** on April 1, 1999, (64 FR 15810), Roxane Laboratories, Inc., 1809 Wilson Road, P.O. Box 16532, Columbus, Ohio 43216–6532, made application by renewal to the Drug Enforcement Administration to be registered as an

importer of cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to import cocaine to manufacture topical solutions for distribution to customers.

A registered bulk manufacturer of cocaine filed written comments and an objection in response to the Notice of Application. The objector argues that it would not be in the public interest to register Roxane because it would violate United States policy against the use of seized material and that competition is adequate. Both of these issues have already been considered and addressed in Roxane's Notice of Registration published in the **Federal Register** on October 19, 1998 (63 FR 5589).

DEA has considered the factors in Title 21, Untied States Code, Section 823(a) and 952(a) and determined that the registration of Roxane Laboratories, Inc. to import cocaine 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 Roxane Laboratories, Inc. on a regular basis to ensure that the company's continued registration is consistent with the public interest. These investigations have included inspection and testing of the company's physical security systems, audits of the company's records, verification of the company's compliance with state and local laws, and a review of the company's background and history. Therefore, pursuant to Section 1008(a) of the Controlled Substances Import and Export Act and in accordance with Title 21, Code of Federal Regulations, Section 1301.34, the above firm is granted registration as an importer of the basic class of controlled substance listed above.

Dated: October 8, 1999.

John H. King,

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

[FR Doc. 99–27096 Filed 10–15–99; 8:45 am]

BILLING CODE 4410–09–M

DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review; Comment Request

September 14, 1999.

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, Departmental Clearance Officer, Ira Mills (202–219–5096 ext. 143) or E–Mail to Mills-IRA@dol.gov.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for BLS, DM, ESA, ETa, MSHA, OSHA, PWBA, or VETS, 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.

Agency: Employment Standards Administration.

Title: Labor Organization and Auxiliary Reports.

OMB Number: 1215–0188.

Affected Public: Not-for-Profit Institutions; Individuals or Households; Business and other for-profit.

Frequency: Semiannually, Annually.

Form	Responses	Hours per respondent	Reporting burden hours	Minutes per respondent	Recordkeeping hours	Total
LM–1	358	0.83	297	5	30	327
LM–2	6,005	14.75	88,574	30	3,003	91,577
LM–3	14,234	6.50	92,521	15	3,559	96,080
LM–4	9,285	0.83	7,707	2	310	8,017
LM–10	211	0.50	106	5	18	124
LM–15	389	1.50	584	20	128	712
LM–15A	81	0.33	27	2	3	30
LM–16	82	0.33	27	1	2	29
LM–20	254	0.33	84	2	8	92
LM–21	64	0.50	32	5	5	37
LM–30	64	0.50	32	5	5	37
S–1	89	0.50	45	5	7	52
SARF*	2,536	0.17	431	2	85	516
Total	33,652	.....	190,467	.....	7,163	197,630

*Simplified Annual Report Format.

Total Annualized capital/startup costs:

Total annual costs (operating/maintaining systems or purchasing services):

Description: The Labor-Management Reporting and Disclosure Act (LMRDA) requires unions to file annual financial reports, and copies of their constitution and bylaws with DOL. Under certain circumstances, reports are required of union officers and employees, employers, labor relations consultants, and surety companies. All reports are available for public disclosure. Filers are required to retain supporting records