

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 November 17, 2006, Mallinckrodt Inc., 3600 North Second Street, St. Louis, Missouri 63147, 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 I and II:

Drug	Schedule
Codeine-N-oxide (9053)	I
Difenoxin (9168)	I
Dihydromorphine (9145)	I
Morphine-N-oxide (9307)	I
Norlevorphanol (9634)	I
Normorphine (9313)	I
Tetrahydrocannabinols (7370)	I
Nabilone (7379)	II
Alfentanil (9737)	II
Amphetamine (1100)	II
Ecgonine (9180)	II
Codeine (9050)	II
Dextropropoxyphene, bulk (9273)	II
Dihydrocodeine (9120)	II
Diphenoxylate (9170)	II
Diprenorphine (9058)	II
Etorphine HCL (9059)	II
Fentanyl (9801)	II
Hydrocodone (9193)	II
Hydromorphone (9150)	II
Levo-alphaacetylmethadol (9648) ..	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Metopon (9260)	II
Morphine (9300)	II
Opium extracts (9610)	II
Opium fluid extract (9620)	II
Opium tincture (9630)	II
Opium, granulated (9640)	II
Opium, powdered (9639)	II
Oxycodone (9143)	II
Oxymorphone (9652)	II
Noroxymorphone (9668)	II
Phenazocine (9715)	II
Alfentanil (9737)	II
Remifentanil (9739)	II
Sufentanil (9740)	II
Thebaine (9333)	II

The firm plans to manufacture the listed controlled substances for internal use and for sale to other companies.

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 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 April 2, 2007.

Dated: January 23, 2007.

Joseph T. Rannazzisi,

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

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DEPARTMENT OF LABOR

Office of the Secretary

Submission for OMB Review: Comment Request

January 24, 2007.

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 from RegInfo.gov at <http://www.reginfo.gov/public/do/PRAMain> or by contacting Darrin King on 202-693-4129 (this is not a toll-free number) / e-mail: king.darrin@dol.gov.

Comments should be sent to Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for the Employee Benefits Security Administration (EBSA), Office of Management and Budget, Room 10235, Washington, DC 20503, Telephone: 202-395-7316 / Fax: 202-395-6974 (these are not toll-free numbers), 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: Employee Benefits Security Administration.

Type of Review: Extension without change of currently approved collection.

Title: Summary Plan Description Requirements Under ERISA.

OMB Number: 1210-0039.

Type of Response: Third party disclosure.

Affected Public: Private Sector: Business or other for-profit and Not-for-profit institutions.

Estimated Number of Respondents: 3,200,000.

Estimated Number of Annual Responses: 93,457,000.

Estimated Total Burden Hours: 262,000.

Estimated Total Annualized capital/startup costs: \$0.

Estimated Total Annual Costs (operating/maintaining systems or purchasing services): \$257,914,000.

Description: Section 104(b)(1) of the Employee Retirement Security Act of 1974 (ERISA) requires the administrator of an employee benefit plan to furnish each plan participant and each beneficiary receiving benefits under the plan a copy of the plan's summary plan description (SPD) within 90 days after an individual becomes a participant and (in the case of a beneficiary) within 90 days after an individual first receives benefits, or, if later, within 120 days after the plan first becomes subject to Part 2 of Title I of ERISA. Section 104(b)(1) further specifies that an updated SPD must be furnished subsequently every fifth year, integrating all plan amendments made within such five-year period. The information required to be contained in the SPD is set forth in section 102(b) of ERISA.

If a plan is amended to make a material modification in its terms or to change the information required to be contained in the SPD (other than a material reduction in covered services or benefits under a group health plan), section 104(b)(1) requires the plan administrator to furnish participants and beneficiaries receiving benefits a summary of the material modifications