

Dated: July 28, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8-17973 Filed 8-4-08; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 28, 2008, and published in the **Federal Register** on April 4, 2008 (73 FR 18570), Siegfried (USA), Inc., Industrial Park Road, Pennsville, New Jersey 08070, 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 schedules I and II:

Drug	Schedule
Dihydromorphine (9145)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Glutethimide (2550)	II
Codeine (9050)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Hydrocodone (9193)	II
Methadone (9250)	II
Methadone intermediate (9254) ...	II
Dextropropoxyphene, bulk (non- dosage forms) (9273).	II
Morphine (9300)	II
Oxymorphone (9652)	II

The company plans to manufacture the listed controlled substances in bulk 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 determined that the registration of Siegfried (USA), Inc., to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Siegfried (USA), 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of

the basic classes of controlled substances listed.

Dated: July 28, 2008.

Joseph T. Rannazzisi,

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

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 10, 2008 and published in the **Federal Register** on March 19, 2008 (73 FR 14840), 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 schedules I and II:

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

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

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Mallinckrodt, Inc. to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Mallinckrodt, 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: July 30, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8-18039 Filed 8-5-08; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated March 11, 2008 and published in the **Federal Register** on March 19, 2008 (73 FR 14841), 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-Piperidinocyclohexane- carbonitrile (8603).	II
Benzoylcegonine (9180)	II

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

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Varian, Inc. to manufacture the listed

basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Varian, 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: July 30, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8-18042 Filed 8-5-08; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated April 9, 2008, and published in the **Federal Register** on April 16, 2008, (73 FR 20718), Research Triangle Institute, Kenneth H. Davis Jr., Hermann Building, East Institute Drive, P.O. Box 12194, Research Triangle, North Carolina 27709, 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 schedules I and II:

Drug	Schedule
Marihuana (7360)	I
Tetrahydrocannabinols (7370)	I
Cocaine (9041)	II

The Institute will manufacture small quantities of cocaine and marihuana derivatives for use by their customers in analytical kits, reagents, and reference standards as directed by the National Institute on Drug Abuse.

No comments or objections have been received. DEA has considered the factors in 21 U.S.C. 823(a) and determined that the registration of Research Triangle Institute to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated Research Triangle Institute 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. 823, and in accordance with 21 CFR 1301.33, the above named company is granted registration as a bulk manufacturer of the basic classes of controlled substances listed.

Dated: July 30, 2008.

Joseph T. Rannazzisi,

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

[FR Doc. E8-18067 Filed 8-5-08; 8:45 am]

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DEPARTMENT OF LABOR

Employment Standards Administration

Proposed Extension of the Approval of Information Collection Requirements

ACTION: Notice.

SUMMARY: The Department of Labor, as part of its continuing effort to reduce paperwork and respondent burden, conducts a preclearance consultation program to provide the general public and Federal agencies with an opportunity to comment on proposed and/or continuing collections of information in accordance with the Paperwork Reduction Act of 1995 (PRA95) [44 U.S.C. 3506(c)(2)(A)]. This program helps to ensure that requested data can be provided in the desired format, reporting burden (time and financial resources) is minimized, collection instruments are clearly understood, and the impact of collection requirements on respondents can be properly assessed. Currently, the Employment Standards Administration is soliciting comments concerning the proposal to extend OMB approval of the information collection: Work Experience and Career Exploration (WECEP) Regulations, 29 CFR 570.35a. A copy of the proposed information collection request can be obtained by contacting the office listed below in the addresses section of this Notice.

DATES: Written comments must be submitted to the office listed in the addresses section below on or before October 6, 2008.

ADDRESSES: Ms. Hazel M. Bell, U.S. Department of Labor, 200 Constitution Ave., NW., Room S-3201, Washington, DC 20210, telephone (202) 693-0418, fax (202) 693-1451, E-mail bell.hazel@dol.gov. Please use only one

method of transmission for comments (mail, fax, or E-mail).

SUPPLEMENTARY INFORMATION

I. Background: The Fair Labor Standards Act (FLSA) section 3(l), 29 U.S.C. 203(l), establishes a minimum age of 16 years for most nonagricultural employment but allows the employment of 14- and 15-year olds in occupations other than manufacturing and mining or deemed hazardous, if the Secretary of Labor determines such employment is confined to (1) periods that will not interfere with the minor's schooling and (2) conditions that will not interfere with the minor's health and well-being. FLSA section 11(c), 29 U.S.C. 211(c), requires all employers covered by the FLSA to make, keep and preserve records of their employees' wages, hours and other conditions and practices of employment. Regulations issued by the Secretary of Labor prescribe the recordkeeping and reporting requirements for these records. Subpart C of Regulations, 29 CFR Part 570, Child Labor Regulations, Orders, and Statements of Interpretation, sets forth the employment standards for 14- and 15-year olds (CL Reg. 3). Regulations 29 CFR 570.35a contains the requirements and criteria for the use of 14- and 15-year olds and the occupations permitted for them, and the conditions of employment that allow for the employment of 14- and 15-year olds, pursuant to a school-supervised and school-administered WECEP—under the conditions CL Reg. 3 otherwise prohibits. In order to utilize the CL Reg. 3 WECEP provisions, Regulations 29 CFR 570.35a(b)(2) requires a state educational agency to file an application for approval of a state WECEP program as one not interfering with schooling or with the health and well-being of the minors involved. Regulations 29 CFR 570.35a(b)(3)(vi) requires the preparation of a written training agreement for each student participating in a WECEP and that such agreement be signed by the teacher-coordinator, employer, and student. The regulation also requires the student's parent or guardian to sign or otherwise consent to the agreement in order for it to be valid. Regulations 29 CFR 570.35a(b)(4)(ii) requires state education agencies to keep a record of the names and addresses of each school enrolling WECEP students and the number of enrollees in each unit. The state or local educational agency office must keep a copy of the written training agreement for each student participating in the WECEP. The records and copies must be maintained for three (3) years from the date of each student's enrollment in the