

for registration as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in Schedule II.

The firm plans to manufacture a Schedule II cocaine derivative as a final intermediate for the production of dopascan injection.

No comments or objections have been received. DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Guilford Pharmaceuticals, Inc. to manufacture the listed controlled substance is consistent with the public interest at this time. DEA has investigated Guilford Pharmaceuticals, Inc. to ensure that the company's registration is consistent with the public interest. This 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic class of controlled substance listed is granted.

Dated: May 26, 2004.

**William J. Walker,**

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

[FR Doc. 04-13537 Filed 6-15-04; 8:45 am]

**BILLING CODE 4410-09-M**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

**Manufacturer of Controlled Substances; Notice of Registration**

By notice dated February 4, 2004, and published in the **Federal Register** on February 18, 2004, (69 FR 7656), Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, 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
Difenoxin (9168) .....	I
Propiram (9649) .....	I
Amphetamine (1100) .....	II
Methylphenidate (1724) .....	II
Codeine (9050) .....	II
Oxycodone (9143) .....	II

Drug	Schedule
Hydrocodone (9193) .....	II
Meperidine (9230) .....	II
Morphine (9300) .....	II
Thebaine (9333) .....	II
Alfentanil (9737) .....	II
Sufentanil (9740) .....	II
Fentanyl (9801) .....	II

The firm plans to manufacture the listed controlled substances in bulk to supply to its customers.

No comments or objections have been received. DEA has considered the factors in title 21, United States Code, section 823(a) and determined that the registration of Johnson Matthey, Inc. to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated Johnson Matthey, Inc. to ensure that the company's registration is consistent with the public interest. This 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 28 CFR 0.100 and 0.104, the Deputy Assistant Administrator, Office of Diversion Control, hereby orders that the application submitted by the above firm for registration as a bulk manufacturer of the basic classes of controlled substances listed is granted.

Dated: May 26, 2004.

**William J. Walker,**

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

[FR Doc. 04-13536 Filed 6-15-04; 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 October 20, 2003, Siegfried (USA), Inc., Industrial Park Road, Pennsville, New Jersey 08070, made application by renewal to the Drug Enforcement Administration (DEA) for registration as a bulk manufacturer of Methadone Intermediate (9254), a basic class of controlled substance in Schedule II. The code was inadvertently dropped from the subsequent Notices of Application and Renewal. On 5/14/2004, DEA received a telephonic communication

requesting that the code be added back onto the firm's registration.

The firm plans to manufacture the listed controlled substance for distribution as a bulk product to its customers.

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: **Federal Register** Representative, Office of Chief Counsel (CCD) and must be filed no later than August 16, 2004.

Dated: June 1, 2004.

**William J. Walker,**

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

[FR Doc. 04-13532 Filed 6-15-04; 8:45 am]

**BILLING CODE 4410-09-M**

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

[Docket No. 03-48]

**Deborah Y. Strauss, D.V.M.,  
Revocation of Registration**

On August 1, 2003, the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration (DEA), issued an Order to Show Cause to Deborah Y. Strauss, D.V.M. (Respondent) notifying her of an opportunity to show cause as to why DEA should not revoke her Certificate of Registration, BS6351821, and deny any pending applications for renewal or modification of such registration pursuant to 21 U.S.C. 824(a)(3) and (a)(4). Specifically, the Order to Show Cause alleged that the Respondent's State controlled substances registration has been suspended and her continued registration would be inconsistent with the public interest based on matters concerning her purported issuance of prescriptions for Demerol, a Schedule II controlled substance, for no legitimate medical purpose. The Order to Show Cause further alleged that as a result of an accountability audit, the Respondent was unable to account for over 10,000 mg. of injectible Demerol, and her records involving the Schedule IV controlled substance diazepam, were not complete or accurate in violation of DEA regulations.