

Dated: April 2, 1999.

**John H. King,**

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

[FR Doc. 99-9089 Filed 4-12-99; 8:45 am]

BILLING CODE 4410-09-M

**DEPARTMENT OF JUSTICE**

**Drug Enforcement Administration**

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

By Notice dated December 2, 1998, and published in the **Federal Register** on December 23, 1998, (63 FR 71156), High Standard Products, 1100 W. Florence Avenue, #B, Inglewood, California 90301, 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 below:

Drug	Schedule
Methaqualone (2565) .....	I
Lysergic acid diethylamide (7315)	I
Tetrahydrocannabinols (7370) .....	I
3,4-Methylenedioxyamphetamine (7400) .....	I
3,4-Methylenedioxy-N-ethylamphetamine (7404) .....	I
3,4-Methylenedioxymethamphetamine (7405) .....	I
4-Methoxyamphetamine (7411) .....	I
Heroin (9200) .....	I
3-Methylfentanyl (9813) .....	I
Amphetamine (1100) .....	II
Methamphetamine (1105) .....	II
Secobarbital (2315) .....	II
Phencyclidine (7471) .....	II
Cocaine (9041) .....	II
Codeine (9050) .....	II
Hydromorphone (9150) .....	II
Diphenoxylate (9170) .....	II
Hydrocodone (9193) .....	II
Methadone (9250) .....	II
Morphine (9300) .....	II
Fentanyl (9801) .....	II

The firms plans to manufacture analytical reference standards.

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 High Standard Products to manufacture the listed controlled substances is consistent with the public interest at this time. DEA has investigated High Standard Products 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 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 above is granted.

Dated: April 2, 1999.

**John H. King,**

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

[FR Doc. 99-9091 Filed 4-12-99; 8:45 am]

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**NATIONAL TRANSPORTATION SAFETY BOARD**

**Sunshine Act Meeting**

**AGENDA—NATIONAL TRANSPORTATION SAFETY BOARD**

**TIME AND DATE:** 9:30 a.m., Tuesday, April 20, 1999.

**PLACE:** NTSB Board Room, 5th Floor, 490 L'Enfant Plaza, S.W., Washington, D.C. 20594.

**STATUS:** Open.

**MATTERS TO BE CONSIDERED:**

7144—Brief of Accident: Gates Learjet 25B, N627WSx, at Houston, Texas, on January 13, 1998, and Safety Recommendation to the Federal Aviation Administration concerning adherence to standard operating procedures and enhanced ground proximity warning systems.

7141—Accident Summary Report and Recommendation: To the Federal Highway Administration and Dion Oil Company concerning procedures and training for loading and unloading cargo tanks, Key West, Florida on June 29, 1998.

**NEWS MEDIA CONTACT:** Telephone: (202) 314-6100.

**FOR MORE INFORMATION CONTACT:** Rhonda Underwood, (202) 314-6065.

Dated: April 9, 1999.

**Rhonda Underwood,**

*Federal Register Liaison Officer.*

[FR Doc. 99-9338 Filed 4-9-99; 3:26 pm]

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**NUCLEAR REGULATORY COMMISSION**

**Agency Information Collection Activities: Proposed Collection; Comment request**

**AGENCY:** U.S. Nuclear Regulatory Commission (NRC).

**ACTION:** Notice of pending NRC action to submit an information collection request to OMB and solicitation of public comment.

**SUMMARY:** The NRC is preparing a submittal to OMB for review of continued approval of information collections under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35).

Information pertaining to the requirement to be submitted:

1. The title of the information collection: 10 CFR part 26, "Fitness for Duty Program".

2. Current OMB approval number: 3150-0146.

3. How often the collection is required: On occasion.

4. Who is required or asked to report: All licensees authorized to construct or operate a nuclear power reactor and all licensees authorized to possess, use, or transport unirradiated Category 1 nuclear material.

5. The number of annual respondents: 72.

6. The number of hours needed annually to complete the requirement or request: 59,800 (5,786 hours of reporting burden and 54,074 hours of recordkeeping burden).

7. Abstract: 10 CFR part 26, "Fitness for Duty Program," requires licensees of nuclear power plants and licensees authorized to possess, use, or transport unirradiated Category 1 nuclear material to implement fitness-for-duty programs to assure that personnel are not under the influence of any substance or mentally or physically impaired, to retain certain records associated with the management of these programs, and to provide reports concerning significant events and program performance. Compliance with these program requirements is mandatory for licensees subject to 10 CFR part 26.

Submit, by June 14, 1999, comments that address the following questions:

1. Is the proposed collection of information necessary for the NRC to properly perform its functions? Does the information have practical utility?

2. Is the burden estimate accurate?

3. Is there a way to enhance the quality, utility, and clarity of the information to be collected?

4. How can the burden of the information collection be minimized,