Drug	Schedule
3,4-Methylenedioxyamphetamine (7400).	I
3,4-Methylenedioxy-N-ethylamphetamine (7404).	I
3,4-Methylenedioxymethamphetamine (7405).	I
4-Methoxyamphetamine (7411)	I
Dimethyltryptamine (7435)	I
Psilocybin (7437)	I
Psilocyn (7438)	I
N-Benzylpiperazine (7493)	I
Acetyldihydrocodeine (9051)	I
Dihydromorphine (9145)	I
Heroin (9200)	I
Normorphine (9313)	I
Pholcodine (9314)	I
Tilidine (9750)	I
3-Methylfentanyl (9813)	I
Amphetamine (1100)	II
Methamphetamine (1105)	II
Methylphenidate (1724)	II
Amobarbital (2125)	II
Pentobarbital (2270)	II
Secobarbital (2315)	II
Phencyclidine (7471)	II
Phenylacetone (8501)	II
Cocaine (9041)	II
Codeine (9050)	II
Dihydrocodeine (9120)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Benzoylecgonine (9180)	II
Ethylmorphine (9190)	II
Hydrocodone (9193)	II
Levorphanol (9220)	II
Meperidine (9230)	II
Methadone (9250)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II
Morphine (9300)	II
Thebaine (9333)	II
Oxymorphone (9652)	II
Alfentanil (9737)	II
Sufentanil (9740)	II
Fentanyl (9801)	II

The company plans to import analytical reference standards for distribution to its customers for research and analytical purposes.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic classes of controlled substances listed in schedule I or II, which fall under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may file comments or objections to the issuance of the proposed registration and may, at the same time, file a written request for a hearing on such application pursuant to 21 CFR 1301.43 and in such form as prescribed by 21 CFR 1316.47.

Any such comments or objections being sent via regular or express mail should be addressed, in quintuplicate, to the Drug Enforcement Administration, Attention: DEA Federal Register Representative/ODL, 8701 Morrissette Drive, Springfield, VA 22152; and must be filed no later than May 11, 2009.

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 published in the **Federal Register** on September 23, 1975, (40 FR 43745), all applicants for registration to import a basic class of any controlled substances 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 1301.34(b), (c), (d), (e) and (f) are satisfied.

Dated: April 1, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–8088 Filed 4–8–09; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances; Notice of Registration

By Notice dated November 26, 2008 and published in the **Federal Register** on December 5, 2008, (73 FR 74095), GE Healthcare, 3350 North Ridge Avenue, Arlington Heights, Illinois 60004–1412, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to manufacture a radioactive product used in diagnostic imaging in the diagnosis of Parkinson's Disease and for manufacture in bulk for investigational new drug (IND) submission and clinical trials.

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 GE Healthcare to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated GE Healthcare 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 class of controlled substance listed.

Dated: April 1, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–8089 Filed 4–8–09; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances Notice of Registration

By Notice dated November 26, 2008 and published in the **Federal Register** on December 5, 2008, (73 FR 74196), ISP Freetown Fine Chemicals, 238 South Main Street, Assonet, Massachusetts 02702, 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
2,5-Dimethoxyamphetamine (7396).	I
Amphetamine (1100)	II
Methylphenidate (1724)	II
Phenylacetone (8501)	II
Dextropropoxyphene, bulk (non-dosage forms) (9273).	II

The company plans to manufacture Phenylacetone to be used in the manufacture of Amphetamine for distribution to its customers. The bulk 2,5-Dimethoxyamphetamine will be used for conversion into non-controlled substances.

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 ISP Freetown Fine Chemicals to manufacture the listed basic classes of controlled substances is consistent with the public interest at this time. DEA has investigated ISP Freetown Fine Chemicals 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: April 1, 2009.

Joseph T. Rannazzisi,

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

[FR Doc. E9–8090 Filed 4–8–09; 8:45 am]

BILLING CODE 4410–09–P

DEPARTMENT OF LABOR

Employment And Training Administration

Announcement Regarding States Triggering ''On'' to the Second-Tier of Emergency Unemployment Compensation 2008 (EUC08)

AGENCY: Employment and Training Administration, Labor.

ACTION: Notice.

SUMMARY: Announcement regarding Colorado, Maryland, and Texas triggering ''on'' to the Second-Tier of Emergency Unemployment Compensation (EUC08).

Public law 110–449 created a Second-Tier of benefits for qualified unemployed workers claiming benefits in high unemployment states. The Department of Labor produces a trigger notice indicating which States qualify for the Second-Tier of EUC08 benefits and provides the beginning and ending dates of the Second-Tier period for each qualifying state. The trigger notice covering State eligibility for the Second-Tier of the EUC08 program can be found at: *http://ows.doleta.gov/unemploy/claims_arch.asp.* A new trigger notice is posted at this location each week that the program is in effect.

Beginning April 12, 2009, Colorado, Maryland, and Texas are in a high unemployment period, resulting in their triggering ''on'' to the Second-Tier of the EUC08 program.

Information for Claimants

The duration of benefits payable in the EUC program, and the terms and conditions under which they are payable, are governed by Public Laws 110–252 and 110–449 and the operating instructions issued to the states by the U.S. Department of Labor. The State Workforce Agency in states beginning a high unemployment period will furnish a written notice of potential entitlement to each individual who is potentially eligible for Second-Tier EUC08 benefits.

Persons who believe they may be entitled to additional benefits under the EUC08 program, or who wish to inquire about their rights under the program, should contact their State Workforce Agency.

FOR FURTHER INFORMATION CONTACT: Scott Gibbons, U.S. Department of Labor, Employment and Training Administration, Office of Workforce Security, 200 Constitution Avenue, NW., Frances Perkins Building, Room S–4231, Washington, DC 20210, telephone number (202) 693–3008 (this is not a toll-free number) or by *e-mail: gibbons.scott@dol.gov*.

Signed in Washington, DC, this 2nd day of April 2009.

Douglas F. Small,

Deputy Assistant Secretary, Employment and Training Administration.

[FR Doc. E9–8079 Filed 4–8–09; 8:45 am]

BILLING CODE 4510–FW–P

NUCLEAR REGULATORY COMMISSION

[NRC–2008–0481]

Agency Information Collection Activities: Submission for the Office of Management and Budget (OMB) Review; Comment Request

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

ACTION: Notice of the OMB review of information collection and solicitation of public comment.

SUMMARY: The NRC has recently submitted to OMB for review the following proposal for the collection of information under the provisions of the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). The NRC hereby informs potential respondents that an agency may not conduct or sponsor, and that a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. The NRC published a **Federal Register** Notice with a 60-day comment period on this information collection on January 5, 2009.

1. *Type of submission, new, revision, or extension:* Extension/Revision.

2. *The title of the information collection:* 10 CFR part 51—''Environmental Protection Regulations for Domestic Licensing and Related Regulatory Functions.''

3. *Current OMB approval number:* 3150–0021.

4. *The form number if applicable:* Not applicable.

5. *How often the collection is required:* Upon submittal of an application for a construction permit, operating license, operating license renewal, early site review, design certification review, decommissioning or termination review, or manufacturing license, or upon submittal of a petition for rulemaking.

6. *Who will be required or asked to report:* Licensees and applicants requesting approvals for actions proposed in accordance with the provisions of 10 CFR parts 30, 32, 33, 34, 35, 36, 39, 40, 50, 52, 54, 60, 61, 70, and 72.

7. *An estimate of the number of annual responses:* 23.

8. *The estimated number of annual respondents:* 23.

9. *An estimate of the total number of hours needed annually to complete the requirement or request:* 92,281.

10. *Abstract:* 10 CFR part 51 specifies information to be provided by applicants and licensees so that the NRC can make determinations necessary to adhere to the policies, regulations, and public laws of the United States, which are to be interpreted and administered in accordance with the policies set forth in the National Environmental Policy Act of 1969, as amended.

A copy of the final supporting statement may be viewed free of charge at the NRC Public Document Room, One White Flint North, 11555 Rockville Pike, Room O–1 F21, Rockville, MD 20852. OMB clearance requests are available at the NRC worldwide Web site: *http://www.nrc.gov/public-involve/doc-comment/omb/index.html.* The document will be available on the NRC home page site for 60 days after the signature date of this notice.

Comments and questions should be directed to the OMB reviewer listed below by May 11, 2009. Comments received after this date will be considered if it is practical to do so, but assurance of consideration cannot be given to comments received after this date.

NRC Desk Officer, Office of Information and Regulatory Affairs (3150–0021), NEOB–10202, Office of Management and Budget, Washington, DC 20503.

The NRC Clearance Officer is Gregory Trussell, (301) 415–6445.

Dated at Rockville, Maryland, this 1st day of April 2009.

For the Nuclear Regulatory Commission.

Gregory Trussell,

NRC Clearance Officer, Office of Information Services.

[FR Doc. E9–8063 Filed 4–8–09; 8:45 am]

BILLING CODE 7590–01–P