Section 404(d) directs the Commission to conduct an annual review of the program for the purpose of evaluating the effectiveness of the program and making recommendations for improvements. The Commission is required to submit its reports containing the results of its reviews to the House Committee on Ways and Means and the Senate Committee on Finance. The Commission submitted its report on its first annual review (USITC Publication 4175) on July 28, 2010, its report on its second annual review (USITC Publication 4246) on July 22, 2011, and its report on the third annual review (USITC Publication 4175) on July 26, 2012: The Commission expects to submit its report on its fourth annual review by July 26, 2013.

The Commission instituted this investigation pursuant to section 332(g) of the Tariff Act of 1930 to facilitate docketing of submissions and also to facilitate public access to Commission records through the Commission's EDIS electronic records system.

Submissions: Interested parties are invited to file written submissions concerning this fourth annual review. All written submissions should be addressed to the Secretary and must conform to the provisions of section 201.8 of the Commission's Rules of Practice and Procedure (19 CFR 201.8). All written submissions must conform with the provisions of section 201.8 of the Commission's Rules of Practice and Procedure (19 CFR 201.8). Section 201.8 and the Commission's Handbook on Filing Procedures require that interested parties file documents electronically on or before the filing deadline and submit eight (8) true paper copies by 12:00 p.m. eastern time on the next business day. In the event that confidential treatment of a document is requested, interested parties must file, at the same time as the eight paper copies, at least four (4) additional true paper copies in which the confidential information must be deleted (see the following paragraph for further information regarding confidential business information). Persons with questions regarding electronic filing should contact the Secretary (202-205-2000).

Any submissions that contain confidential business information must also conform to the requirements of section 201.6 of the Commission's *Rules of Practice and Procedure* (19 CFR 201.6). Section 201.6 of the rules requires that the cover of the document and the individual pages be clearly marked as to whether they are the "confidential" or "non-confidential" version, and that the confidential business information is clearly

identified by means of brackets. All written submissions, except for confidential business information, will be made available for inspection by interested parties.

The Commission intends to publish only a public report in this review. Consequently, the report that the Commission sends to the committees will not contain any confidential business information. Any confidential business information received by the Commission in this investigation and used in preparing its report will not be published in a manner that would reveal the operations of the firm supplying the information.

Issued: March 8, 2013.

By order of the Commission.

Lisa R. Barton,

Acting Secretary to the Commission. [FR Doc. 2013–05830 Filed 3–13–13; 8:45 am]

BILLING CODE 7020-02-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Manufacturer of Controlled Substances, Notice of Registration; Johnson Matthey, Inc.

By Notice dated November 5, 2012, and published in the **Federal Register** on November 13, 2012, 77 FR 67676, Johnson Matthey, Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066–1742, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as a bulk manufacturer of the following basic classes of controlled substances:

Drug	Schedule
Gamma Hydroxybutyric Acid (2010).	I
Tetrahydrocannabinols (7370)	1
Dihydromorphine (9145)	!
Difenoxin (9168) Propiram (9649)	1
Amphetamine (1100)	
Methamphetamine (1105)	ii
Lisdexamfetamine (1205)	II
Methylphenidate (1724)	II
Nabilone (7379)	II
Cocaine (9041)	II
Codeine (9050)	II II
Dihydrocodeine (9120) Oxycodone (9143)	
Hydromorphone (9150)	
Diphenoxylate (9170)	ii
Ecgonine (9180)	lii
Hydrocodone (9193)	II
Meperidine (9230)	П
Methadone (9250)	II
Methadone intermediate (9254)	II
Morphine (9300)	Ш
Thebaine (9333)	II

Drug	Schedule
Oxymorphone (9652) Noroxymorphone (9668) Alfentanil (9737) Remifentanil (9739) Sufentanil (9740) Fentanyl (9801)	

The company plans to manufacture the listed controlled substances in bulk for sale 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 Johnson Matthey, Inc., to manufacture the listed basic classes of 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. 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(a), 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: February 27, 2013.

Joseph T. Rannazzisi,

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

[FR Doc. 2013-05803 Filed 3-13-13; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF LABOR

Employment and Training Administration

Comment Request for Information Collection: Confidentiality and Disclosure of State Unemployment Compensation Information Final Rule and State Income and Eligibility Verification Provisions of the Deficit Reduction Act of 1984, Extension Without Change

AGENCY: Employment and Training Administration (ETA), Labor.

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