including the validity of the methodology and assumptions used;

—Enhance the quality, utility, and clarity of the information to be collected; and

—Minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, e.g., permitting electronic submission of responses.

Overview of This Information Collection

(1) *Type of Information Collection:* Extension of a currently approved collection.

(2) *Title of the Form/Collection:* Application For Restoration of Firearms Privileges.

(3) *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* Form Number: ATF F 3210.1, Bureau of Alcohol, Tobacco, Firearms and Explosives.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: Individuals or households. Other: Business or other for profit. Certain categories of persons are prohibited from possessing firearms. ATF F 3210.1, Application For Restoration of Firearms Privileges is the basis for ATF investigating the merits of an applicant to have his/her rights restored.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* It is estimated that 250 respondents will complete a 30 minute form.

(6) *An estimate of the total public burden (in hours) associated with the collection:* There are an estimated 125 annual total burden hours associated with this collection.

FOR FURTHER INFORMATION CONTACT: Robert B. Briggs, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, Department of Justice, Patrick Henry Building, Suite 1600, 601 D Street, NW., Washington, DC 20530.

Dated: May 10, 2006.

Robert B. Briggs,

Department Clearance Officer, Department of Labor.

[FR Doc. 06–4513 Filed 5–12–06; 8:45 am]

BILLING CODE 4410–FY–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 December 14, 2005, and February 14, 2006, Johnson Matthey Inc., Custom Pharmaceuticals Department, 2003 Nolte Drive, West Deptford, New Jersey 08066, made application by letter 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
Dihydrocodeine (9120)	II
Oxymorphone (9652)	II

The company plans to manufacture in bulk, for distribution to its customers, who are final dosage manufacturers.

Any other such applicant and any person who is presently registered with DEA to manufacture such a substance may file comments or objections to the issuance of the proposed registration pursuant to 21 CFR 1301.33(a).

Any such written comments or objections being sent via regular mail may be addressed, in quintuplicate, to the Deputy Assistant Administrator, Office of Diversion Control, Drug Enforcement Administration, Washington, DC 20537, Attention: DEA Federal Register Representative, Liaison and Policy Section (ODL); or any being sent via express mail should be sent to DEA Headquarters, Attention: DEA Federal Register Representative/ODL, 2401 Jefferson-Davis Highway, Alexandria, Virginia 22301; and must be filed no later than July 14, 2006.

Dated: May 9, 2006.

Joseph T. Rannazzisi,

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

[FR Doc. E6–7338 Filed 5–12–06; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–218N]

RIN 1117–AA61

Electronic Prescriptions for Controlled Substances; Notice of Meeting

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Notice of meeting.

SUMMARY: The Drug Enforcement Administration (DEA), in conjunction with the Department of Health and Human Services (HHS), is conducting a public meeting to discuss electronic prescriptions for controlled substances. Specifically, this meeting is intended to allow industry—prescribers, pharmacies, software/hardware vendors, and other interested third parties—to address how electronic prescribing systems can meet DEA's prescription requirements under the Controlled Substances Act, without unduly burdening the parties to electronic prescribing transactions.

DATES: This meeting will be held Tuesday, July 11, 2006, and Wednesday, July 12, 2006, 8:30 a.m. until 5:30 p.m. Registration will begin at 7:30 a.m. This meeting will be held at the Marriott Crystal City at Reagan National Airport, 1999 Jefferson-Davis Highway, Arlington, VA 22202; (703) 413–5500. The meeting will take place in the Crystal Forum amphitheatre, adjacent to the hotel.

Meeting Attendance: To ensure proper handling, please reference ''Docket No. DEA–218N'' on all written and electronic correspondence regarding this meeting. Persons wishing to attend this meeting, space permitting, must provide attendee information to the Liaison and Policy Section, Office of Diversion Control, Drug Enforcement Administration, via e-mail to *dea.diversion.policy@usdoj.gov,* or via facsimile, (202) 353–1079, as specified below. Persons wishing to attend the meeting must provide this information to the Liaison and Policy Section no later than June 26, 2006.

Comments: All written comments will be made available at the Diversion Control Program Web site, *http://www.deadiversion.usdoj.gov* prior to the public meeting. Therefore, as this is a public meeting, confidential business information or other proprietary information SHOULD NOT be presented at this meeting.

Persons wishing to provide written comments must do so no later than June 26, 2006. To ensure proper handling of