

that Applicant's pending application be denied.

Order

Pursuant to the authority vested in me by 21 U.S.C. 823(f), as well as 28 CFR 0.100(b) and 28 CFR 0.104, I order that the application of Wheatland Pharmacy, for a DEA Certificate of Registration as a retail pharmacy, be, and it hereby is, denied. This order is effective immediately.

Dated: November 8, 2013.

Thomas M. Harrigan,
Deputy Administrator.

[FR Doc. 2013-27700 Filed 11-18-13; 8:45 am]

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; Mylan Pharmaceuticals, Inc.

Pursuant to Title 21 Code of Federal Regulations 1301.34(a), this is notice that on October 7, 2013, Mylan Pharmaceuticals, Inc., 781 Chestnut Ridge Road, Morgantown, West Virginia 26505, made application by renewal to the Drug Enforcement Administration (DEA) to be registered as an importer of the following basic classes of controlled substances:

Drug	Schedule
Amphetamine (1100)	II
Methylphenidate (1724)	II
Oxycodone (9143)	II
Hydromorphone (9150)	II
Methadone (9250)	II
Morphine (9300)	II
Fentanyl (9801)	II

The company plans to import the listed controlled substances in finished dosage form (FDF) from foreign sources for analytical testing and clinical trials in which the foreign FDF will be compared to the company's own domestically-manufactured FDF. This analysis is required to allow the company to export domestically-manufactured FDF to foreign markets.

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 II, which falls under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), 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 written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than December 19, 2013.

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 43745-46, all applicants for registration to import basic classes of any controlled substances in schedules 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: November 12, 2013.

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

[FR Doc. 2013-27660 Filed 11-18-13; 8:45 am]

BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

Importer of Controlled Substances; Notice of Application; GE Healthcare

Pursuant to Title 21, Code of Federal Regulations 1301.34(a), this is notice that on September 18, 2013, 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 an importer of Cocaine (9041), a basic class of controlled substance listed in schedule II.

The company plans to import small quantities of ioflupane, in the form of three separate analogues of Cocaine that will be used for the support and manufacture of DaTSCAN (ioflupane 1-123) injection for distribution as a radioactive diagnostic imaging agent utilized in the diagnosis of Parkinson's disease.

Any bulk manufacturer who is presently, or is applying to be, registered with DEA to manufacture such basic class of controlled substance

listed in schedules I and II, which falls under the authority of section 1002(a)(2)(B) of the Act (21 U.S.C. 952(a)(2)(B)) may, in the circumstances set forth in 21 U.S.C. 958(i), 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 written comments or objections should be addressed, in quintuplicate, to the Drug Enforcement Administration, Office of Diversion Control, Federal Register Representative (ODW), 8701 Morrisette Drive, Springfield, Virginia 22152; and must be filed no later than December 19, 2013.

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 43745-46, all applicants for registration to import a basic class of any controlled substance in schedules 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: November 5, 2013.

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

[FR Doc. 2013-27661 Filed 11-18-13; 8:45 am]

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DEPARTMENT OF JUSTICE

Federal Bureau of Investigation

[OMB Number 1110-0046]

Agency Information Collection Activities; Existing Collection, Comments Requested: Friction Ridge Cards: Arrest and Institution; Applicant; Personal Identification; FBI Standard Palm Print; Supplemental Finger and Palm Print

ACTION: 30-day Notice of Information Collection for Reinstatement.

The Department of Justice (DOJ), Federal Bureau of Investigation (FBI), Criminal Justice Information Services (CJIS) Division will be submitting the following information collection renewal to the Office of Management and Budget (OMB) for review in

accordance with established review procedures of the Paperwork Reduction Act of 1995. The information collection is published to obtain comments from the public and affected agencies. The information collection was previously published in the **Federal Register** Volume 78, Number 179, Page 56940, on September 16, 2013, allowing for a 60-day comment period.

The purpose of this notice is to allow for an additional 30 days for public comment until December 19, 2013. This process is conducted in accordance with 5 CFR 1320.10.

Written comments and/or suggestions regarding the items contained in this notice, especially the estimated public burden and associated response time, should be directed to the OMB, Office of Information and Regulatory Affairs, Attention Department of Justice Desk Officer, Washington, DC 20503. Additionally, comments may be submitted to OMB via facsimile to (202) 395-5806. Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address one or more of the following four points:

(1) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have a practical utility;

(2) Evaluate the accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used;

(3) Enhance the quality, utility, and clarity of the information to be collected; and

(4) 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 of other forms of information technology, e.g., permitting electronic submission of responses.

Overview of this information collection:

(1) *Type of information collection:* Reinstatement, with change, of a previously approved collection for which approval has expired. Reference: OMB control number of 1110-0046.

(2) *The title of the form/collection:* Friction Ridge Cards: Arrest and Institution; Applicant; Personal Identification; FBI Standard Palm Print; Supplemental Finger and Palm Print.

(3) *The agency form number, if any, and the applicable component of the*

department sponsoring the collection: Forms FD-249 (Arrest and Institution), FD-258 (Applicant), and FD-353 (Personal Identification); FD-884 (FBI Standard Palm Print); FD-884a (Supplemental Finger and Palm Print) encompassed under OMB 1110-0046; CJIS Division, FBI, DOJ.

(4) *Affected public who will be asked or required to respond, as well as a brief abstract:* Primary: City, county, state, federal and tribal law enforcement agencies; civil entities requesting security clearance and background checks. This collection is needed to collect information on individuals requesting background checks, security clearance, or those individuals who have been arrested for or accused of criminal activities. Acceptable data is stored as part of the Integrated Automated Fingerprint Identification System (IAFIS) of the FBI.

(5) *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* There are approximately 74,793 agencies as respondents at 10 minutes per fingerprint card completed.

(6) *An estimate of the total public burden (in hours) associated with this collection:* There are approximately 10.1 million annual burden hours associated with this collection.

If additional information is required contact: Jerri Murray, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, U.S. Department of Justice, Two Constitution Square, 145 N Street NE., Washington, DC 20530.

Dated: November 14, 2013.

Jerri Murray,

*Department Clearance Officer for PRA,
United States Department of Justice.*

[FR Doc. 2013-27650 Filed 11-18-13; 8:45 am]

BILLING CODE 4410-02-P

DEPARTMENT OF LABOR

Office of the Secretary

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Crawler, Locomotive, and Truck Cranes Standard

ACTION: Notice.

SUMMARY: The Department of Labor (DOL) is submitting the Occupational Safety and Health Administration (OSHA) sponsored information collection request (ICR) titled, "Crawler, Locomotive, and Truck Cranes Standard," to the Office of Management and Budget (OMB) for review and

approval for continued use, without change, in accordance with the Paperwork Reduction Act of 1995 (PRA), 44 U.S.C. 3501 et seq.

DATES: Submit comments on or before December 19, 2013.

ADDRESSES: A copy of this ICR with applicable supporting documentation; including a description of the likely respondents, proposed frequency of response, and estimated total burden may be obtained free of charge from the RegInfo.gov Web site at http://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=201307-1218-004 (this link will only become active on the day following publication of this notice) or by contacting Michel Smyth by telephone at 202-693-4129 (this is not a toll-free number) or sending an email to DOL_PRA_PUBLIC@dol.gov.

Submit comments about this request by mail or courier to the Office of Information and Regulatory Affairs, Attn: OMB Desk Officer for DOL-OSHA, Office of Management and Budget, Room 10235, 725 17th Street NW., Washington, DC 20503; by Fax: 202-395-6881 (this is not a toll-free number); or by email: OIRA_submission@omb.eop.gov. Commenters are encouraged, but not required, to send a courtesy copy of any comments by mail or courier to the U.S. Department of Labor-OASAM, Office of the Chief Information Officer, Attn: Information Policy and Assessment Program, Room N1301, 200 Constitution Avenue NW., Washington, DC 20210; or by email: DOL_PRA_PUBLIC@dol.gov.

FOR FURTHER INFORMATION CONTACT: Contact Michel Smyth by telephone at 202-693-4129 (this is not a toll-free number) or by email at DOL_PRA_PUBLIC@dol.gov.

Authority: 44 U.S.C. 3507(a)(1)(D).

SUPPLEMENTARY INFORMATION: This ICR seeks continued PRA authorization for the Crawler, Locomotive, and Truck Cranes Standard information collection requirements. The Standard requires performance of a monthly inspection on cranes and running ropes and preparation of a certification record for each inspection. A rope that has been idle for a month or more must undergo a thorough inspection and a certification record must be generated. Occupational Safety and Health Act sections 6(b)(7), 29 U.S.C. 655(b)(7), and 8(c), 29 U.S.C. 657(c), authorize this information collection.

This information collection is subject to the PRA. A Federal agency generally cannot conduct or sponsor a collection of information, and the public is generally not required to respond to an information collection, unless it is