EARLY TERMINATION GRANTED
[07/21/2021]

20211736	G	Green Dot Corporation; Republic Bancorp, Inc.; Republic Bank & Trust Company.

Suzanne Morris,
Chief, Premerger and Division Statistics, Antitrust Division, Department of Justice.
[FR Doc. 2021–15904 Filed 7–26–21; 8:45 am]
BILLING CODE 4410–11–P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–866]

Importer of Controlled Substances Application: Catalent Pharma Solutions, LLC.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Catalent Pharma Solutions, LLC. has applied to be registered as an importer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before August 26, 2021. Such persons may also file a written request for a hearing on the application on or before August 26, 2021.

ADDRESSES: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152. All requests for a hearing must be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrissette Drive, Springfield, Virginia 22152. All request for a hearing should also be sent to: (1) Drug Enforcement Administration, Attn: Hearing Clerk/OALJ, 8701 Morrissette Drive, Springfield, Virginia 22152; and (2) Drug Enforcement Administration, Attn: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on June 2, 2021, Catalent Pharma Solutions, LLC., 3031 Red Lion Road, Philadelphia, Pennsylvania 19114, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Gamma Hydroxybutyric Acid	2010	I
Lysergic Acid Diethylamide	7315	I

The company plans to import the above controlled substances as finished dosage unit products for clinical trials, research, and analytical activities. No other activity for these drug codes is authorized for this registration.

Approval of permit applications will occur only when the registrant's business activity is consistent with what is authorized under 21 U.S.C. 952(a)(2). Authorization will not extend to the import of Food and Drug Administration-approved or non-approved finished dosage forms for commercial sale.

Brian S. Besser,
Acting Assistant Administrator.
[FR Doc. 2021–15919 Filed 7–26–21; 8:45 am]
BILLING CODE P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA–867]

Bulk Manufacturer of Controlled Substances Application: Novitium Pharma, LLC

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Novitium Pharma, LLC has applied to be registered as a bulk manufacturer of basic class(es) of controlled substance(s). Refer to **SUPPLEMENTARY INFORMATION** listed below for further drug information.

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may file written comments on or objections to the issuance of the proposed registration on or before September 27, 2021. Such persons may also file a written request for a hearing on the application on or before September 27, 2021.

ADDRESS: Written comments should be sent to: Drug Enforcement Administration, Attention: DEA Federal Register Representative/DPW, 8701 Morrissette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on June 17, 2021, Novitium Pharma, LLC., 70 Lake Drive, East Windsor, New Jersey 08520, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Psilocybin	7347	I
Psilocyn	7348	I

The company plans to bulk manufacture the above controlled substances to produce Active Pharmaceutical Ingredient (API) and finished dosage forms for clinical trial purposes. No other activities for these drug codes are authorized for this registration.

Brian S. Besser,
Acting Assistant Administrator.
[FR Doc. 2021–15920 Filed 7–26–21; 8:45 am]
BILLING CODE 4410–09–P

DEPARTMENT OF JUSTICE

Notice of Lodging of Proposed Consent Decree Under the Clean Water Act

On July 16, 2021, the U.S. Department of Justice (DOJ) lodged a proposed Amendment to Consent Decree with the United States District Court for the Northern District of Indiana in *United States and State of Indiana* v. *City of Elkhart, Indiana,* Civil Action No. 2:11CV328. The lodging of the proposed Amendment to Consent Decree, by the United States on behalf of the U.S. Environmental Protection Agency, with the concurrence of the State of Indiana on behalf of the Indiana Department of Environmental Management, modifies the Consent Decree in this action that was entered by the Court on November 30, 2011.

The 2011 Consent Decree resolved claims for civil penalties as well as injunctive relief in the form of a Long Term Control Plan (LTCP) for violations of the Clean Water Act and related State law claims in connection with the City of Elkhart's operation of its municipal wastewater and sewer system. The proposed Amendment to Consent Decree modifies the LTCP by allowing the City to change the technology

currently used to treat the wastewater originating from its combined sewer system, allowing one additional year to implement such change.

The publication of this notice opens a period for public comment on the proposed Consent Decree. Comments should be addressed to the Acting Assistant Attorney General, Environment and Natural Resources Division, and should refer to *United States and State of Indiana* v. *City of Elkhart, Indiana,* D.J. Ref. No. 90–5–1–1–08202. All comments must be submitted no later than thirty (30) days after the publication date of this notice. Comments may be submitted either by email or by mail:

To submit comments:	*Send them to:*
By email	*pubcomment–ees.enrd@usdoj.gov.*
By mail	Acting Assistant Attorney General, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611.

During the public comment period, the proposed Consent Decree may be examined and downloaded at this Justice Department website: *http://www.justice.gov/enrd/consent-decrees.*

We will provide a paper copy of the proposed Consent Decree upon written request and payment of reproduction costs. Please mail your request and payment to: Consent Decree Library, U.S. DOJ—ENRD, P.O. Box 7611, Washington, DC 20044–7611.

Please enclose a check or money order for $18.25 (25 cents per page reproduction cost), payable to the United States Treasury.

Patricia McKenna,

Assistant Section Chief, Environmental Enforcement Section, Environment and Natural Resources Division.

[FR Doc. 2021–15934 Filed 7–26–21; 8:45 am]

BILLING CODE 4410–15–P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Cotton Dust Standard

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety and Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that agency receives on or before August 26, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

Comments are invited on: (1) Whether the collection of information is necessary for the proper performance of the functions of the Department, including whether the information will have practical utility; (2) if the information will be processed and used in a timely manner; (3) the accuracy of the agency's estimates of the burden and cost of the collection of information, including the validity of the methodology and assumptions used; (4) ways to enhance the quality, utility and clarity of the information collection; and (5) ways to minimize the burden of the collection of information on those who are to respond, including the use of automated collection techniques or other forms of information technology.

FOR FURTHER INFORMATION CONTACT: Crystal Rennie by telephone at 202–693–0456 or by email at *DOL_PRA_PUBLIC@dol.gov.*

SUPPLEMENTARY INFORMATION: The purpose of the cotton dust standard and its information collection requirements is to provide protection for employees from the adverse health effects associated with occupational exposure to cotton dust. Employers must monitor employee exposure, reduce employee exposure to within permissible exposure limits, provide employees with medical examinations and training, and establish and maintain employee exposure monitoring and medical records. For additional substantive information about this ICR, see the related notice published in the **Federal Register** on April 27, 2021 (86 FR 22277).

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 the OMB approves it and displays a currently valid OMB Control Number. In addition, notwithstanding any other provisions of law, no person shall generally be subject to penalty for failing to comply with a collection of information that does not display a valid OMB Control Number. *See* 5 CFR 1320.5(a) and 1320.6.

DOL seeks PRA authorization for this information collection for three (3) years. OMB authorization for an ICR cannot be for more than three (3) years without renewal. The DOL notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Agency: DOL–OSHA.

Title of Collection: Cotton Dust Standard.

OMB Control Number: 1218–0061.

Affected Public: Private Sector: Business or other for-profits.

Total Estimated Number of Respondents: 4,543.

Total Estimated Number of Responses: 17,217.

Total Estimated Annual Time Burden: 6,379 hours.

Total Estimated Annual Other Costs Burden: $845,662.

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

Crystal Rennie,

Senior PRA Analyst.

[FR Doc. 2021–15931 Filed 7–26–21; 8:45 am]

BILLING CODE 4510–26–P

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Bloodborne Pathogens Standard

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Occupational Safety and Health Administration (OSHA)-sponsored information collection request (ICR) to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). Public comments on the ICR are invited.

DATES: The OMB will consider all written comments that agency receives on or before August 26, 2021.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

Comments are invited on: (1) Whether the collection of information is