

DEPARTMENT OF JUSTICE**Bureau of Alcohol, Tobacco, Firearms and Explosives**

[OMB 1140–0067]

Agency Information Collection Activities; Proposed eCollection of eComments Requested; Revision of a Currently Approved Collection; Licensed Firearms Manufacturers Records of Production, Disposition, and Supporting Data

AGENCY: Bureau of Alcohol, Tobacco, Firearms and Explosives, Department of Justice.

ACTION: 60-Day notice.

SUMMARY: The Bureau of Alcohol, Tobacco, Firearms and Explosives (ATF), Department of Justice (DOJ), will submit the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The proposed collection OMB 1140–0067 (Licensed Firearms Manufacturers Records of Production, Disposition, and Supporting Data) is being revised due to an increase in respondents, although the total responses and burden hours have reduced since the last renewal in 2018. This information collection is also being published to obtain comments from the public and affected agencies.

DATES: Comments are encouraged and will be accepted for 60 days until September 27, 2021.

FOR FURTHER INFORMATION CONTACT: If you have additional comments regarding the estimated public burden or associated response time, suggestions, or need a copy of the proposed information collection instrument with instructions, or additional information, please contact: Dawn Smith, ATF Firearms Industry Programs Branch, either by mail at 244 Needy Road, Martinsburg, WV 25405, by email at fipb-informationcollection@atf.gov, or by telephone at 202–648–0890.

SUPPLEMENTARY INFORMATION: 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:

—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 practical utility;
 —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;
 —Evaluate whether and if so how the quality, utility, and clarity of the information to be collected can be enhanced; 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 (check justification or form 83):* Revision of a currently approved collection.

2. *The Title of the Form/Collection:* Licensed Firearms Manufacturers Records of Production, Disposition, and Supporting Data.

3. *The agency form number, if any, and the applicable component of the Department sponsoring the collection:*
Form number (if applicable): None.
Component: Bureau of Alcohol, Tobacco, Firearms and Explosives, U.S. Department of Justice.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:*

Primary: Business or other for-profit.
Other (if applicable): None.

Abstract: Firearms manufacturers must create and maintain permanent records of all firearms manufactured and disposed of. These records support the Bureau of Alcohol, Tobacco, Firearms and Explosives' mission to inquire into the disposition of any firearm, during the course of during the course a criminal investigation.

5. *An estimate of the total number of respondents and the amount of time estimated for an average respondent to respond:* An estimated 10,513 respondents will respond 677.12822 times per year to this information collection, and it will take each respondent approximately 1.06 minutes to complete a response.

6. *An estimate of the total public burden (in hours) associated with the collection:* The estimated annual public burden associated with this collection is 123,801 hours, which is equal to 10,513

(# of respondents) * 677.12822 (# of responses per respondent) * .0176728 (1.06 minutes).

7. *An Explanation of the Change in Estimates:* The increase in total respondents by 1,457, is due to more firearms manufacturers responding to this collection. However, the total responses and burdens hours decreased by 4,378,792 and 75,4040 hours respectively, because less firearms were produced since the last renewal of this collection in 2018.

If additional information is required contact: Melody Braswell, Department Clearance Officer, United States Department of Justice, Justice Management Division, Policy and Planning Staff, Two Constitution Square, 145 N Street NE, Mail Stop 3E.405A, Washington, DC 20530.

Dated: July 22, 2021.

Melody Braswell,

Department Clearance Officer for PRA, U.S. Department of Justice.

[FR Doc. 2021–15946 Filed 7–26–21; 8:45 am]

BILLING CODE 4410–FY–P

DEPARTMENT OF JUSTICE**Antitrust Division****Granting of Requests for Early Termination of the Waiting Period Under the Premerger Notification Rules**

Section 7A of the Clayton Act, 15 U.S.C. 18a, as added by Title II of the Hart-Scott-Rodino Antitrust Improvements Act of 1976, requires persons contemplating certain mergers or acquisitions to give the Federal Trade Commission and the Assistant Attorney General advance notice and to wait designated periods before consummation of such plans. Section 7A(b)(2) of the Act permits the agencies, in individual cases, to terminate this waiting period prior to its expiration and requires that notice of this action be published in the **Federal Register**. The following transaction was granted early termination—on the date indicated—of the waiting period provided by law and the premerger notification rules. The listing includes the transaction number and the parties to the transaction. The Federal Trade Commission and the Assistant Attorney General for the Antitrust Division of the Department of Justice made the grants. Neither agency intends to take any action with respect to these proposed acquisitions during the applicable waiting period.

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