

DATES: Registered bulk manufacturers of the affected basic class(es), and applicants therefore, may submit electronic comments on or objections to the issuance of the proposed registration on or before January 29, 2024. Such persons may also file a written request for a hearing on the application on or before January 29, 2024.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.33(a), this is notice that on October 23, 2023, Pharmaron Manufacturing Services (US) LLC, 498 Washington Street, Coventry, Rhode Island 02816, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I

The company plans to bulk manufacture the listed controlled substance for the purpose of producing material for clinical trials. No other activities for this drug code are authorized for this registration.

Claude Redd,
Acting Deputy Assistant Administrator.
[FR Doc. 2023-26343 Filed 11-29-23; 8:45 am]
BILLING CODE 4410-09-P

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

[Docket No. DEA-1298]

**Importer of Controlled Substances
Application: Pharmaron Manufacturing
Services (US) LLC**

AGENCY: Drug Enforcement Administration, Justice.
ACTION: Notice of application.

SUMMARY: Pharmaron Manufacturing Services (US) 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 submit electronic comments on or objections to the issuance of the proposed registration on or before January 2, 2024. Such persons may also file a written request for a hearing on the application on or before January 2, 2024.

ADDRESSES: The Drug Enforcement Administration requires that all comments be submitted electronically through the Federal eRulemaking Portal, which provides the ability to type short comments directly into the comment field on the web page or attach a file for lengthier comments. Please go to <https://www.regulations.gov> and follow the online instructions at that site for submitting comments. Upon submission of your comment, you will receive a Comment Tracking Number. Please be aware that submitted comments are not instantaneously available for public view on <https://www.regulations.gov>. If you have received a Comment Tracking Number, your comment has been successfully submitted and there is no need to resubmit the same comment. All requests for a hearing must 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. All requests for a hearing should also be sent to: Drug Enforcement Administration, Attn: Administrator, 8701 Morrisette Drive, Springfield, Virginia 22152.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on October 30, 2023, Pharmaron Manufacturing Services (US) LLC, 498 Washington Street, Coventry, Rhode Island 02816, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

Controlled substance	Drug code	Schedule
Dimethyltryptamine	7435	I

The company purpose of importing Dimethyltryptamine (7435) is to conduct process and analytical technology transfer, further process, and

analytical development as needed and subsequently manufacture/produce an Active Pharmaceutical Ingredient under Good Manufacturing Practices at the US Pharmaron site (Pharmaron Manufacturing Services (US) LLC in Coventry, Rhode Island. No other activity for this drug code 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.

Claude Redd,
Acting Deputy Assistant Administrator.
[FR Doc. 2023-26342 Filed 11-29-23; 8:45 am]
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DEPARTMENT OF JUSTICE

[OMB Number 1110-0057]

**Agency Information Collection
Activities; Proposed eCollection
eComments Requested; Uniform Crime
Reporting (UCR) Instrument Pretesting
and Burden Estimation Generic
Clearance**

AGENCY: Federal Bureau of Investigation, Department of Justice.
ACTION: 30-Day notice.

SUMMARY: The Federal Bureau of Investigation (FBI), Department of Justice (DOJ), will be submitting 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 information collection was previously published in the **Federal Register** on September 15, 2023, allowing a 60-day comment period.

DATES: Comments are encouraged and will be accepted for 30 days until January 2, 2024.

FOR FURTHER INFORMATION CONTACT: If you have comments especially on 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: Edward L. Abraham, Crime and Law Enforcement Statistics Unit Chief, FBI, CJIS Division, Module D-1, 1000 Custer Hollow Road, Clarksburg, West Virginia 26306; telephone number: 304-625-4830, email: elabraham@fbi.gov.
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;
- Enhance the quality, utility, and clarity of the information to be collected; and/or
- 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.

Written comments and recommendations for this information collection should be submitted within 30 days of the publication of this notice on the following website 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 and entering either the title of the information collection or the OMB Control Number 1110–0057. This information collection request may be viewed at www.reginfo.gov. Follow the instructions to view Department of Justice, information collections currently under review by OMB.

DOJ 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 DOJ notes that information collection requirements submitted to the OMB for existing ICRs receive a month-to-month extension while they undergo review.

Overview of This Information Collection

1. *Type of Information Collection:* Extension of a previously approved collection.
2. *Title of the Form/Collection:* UCR Instrument Pretesting and Burden Estimation Generic Clearance.
3. *Agency form number, if any, and the applicable component of the Department of Justice sponsoring the collection:* The form number is 1110–

0057. The applicable component within DOJ is the CJIS Division, FBI.

4. *Affected public who will be asked or required to respond, as well as a brief abstract:* Abstract: This clearance provides the FBI's UCR Program the ability to conduct pretests which evaluate the validity and reliability of information collection instruments and determine the level of burden state and local agencies have in reporting crime data to the FBI. The PRA only allows for nine or fewer respondents in the collection of information, such as pretesting activities. This clearance request expands the pretesting sample to 350 people for each of the information collections administered by the UCR Program. Further, the clearance will allow for a brief five-minute cost and burden assessment for the 18,000 law enforcement agencies participating in the UCR Program.

5. *Obligation to Respond:* the obligation to respond is voluntary.

6. *Total Estimated Number of Respondents:* 18,000 law enforcement respondents.

7. *Estimated Time per Respondent:* five minutes per submission.

8. *Frequency:* 1/annually.

9. *Total Estimated Annual Time Burden:* 1,850 hours annual burden.

10. *Total Estimated Annual Other Costs Burden:* \$0.

If additional information is required, contact: Darwin Arceo, Department Clearance Officer, Policy and Planning Staff, Justice Management Division, United States Department of Justice, Two Constitution Square, 145 N Street NE, 4W–218 Washington, DC 20530.

Dated: November 9, 2023.

Darwin Arceo,

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

[FR Doc. 2023–26372 Filed 11–29–23; 8:45 am]

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DEPARTMENT OF JUSTICE

National Institute of Justice

[OJP (NIJ) Docket No. 1819]

Publication of NIJ Standard 0101.07, Ballistic Resistance of Body Armor, and NIJ Standard 0123.00, Specification for NIJ Ballistic Protection Levels and Associated Test Threats, and Information About the NIJ Compliance Testing Program

AGENCY: National Institute of Justice, Office of Justice Programs, U.S. Department of Justice.

ACTION: Notice.

SUMMARY: The National Institute of Justice (NIJ) announces publication of NIJ Standard 0101.07, *Ballistic Resistance of Body Armor* and NIJ Standard 0123.00, *Specification for NIJ Ballistic Protection Levels and Associated Test Threats*. The NIJ Compliance Testing Program (CTP) will be transitioning program operations to focus on implementing NIJ Standard 0101.07 as described below.

FOR FURTHER INFORMATION CONTACT: Mark Greene, Office Director, Office of Technology and Standards, National Institute of Justice, 810 7th Street NW, Washington, DC 20531, by telephone at (202) 598–9481 [Note: this is not a toll-free telephone number], or by email at mark.greene2@usdoj.gov.

SUPPLEMENTARY INFORMATION: The National Institute of Justice (NIJ) announces publication of NIJ Standard 0101.07, *Ballistic Resistance of Body Armor*. The document can be found here: <https://www.ojp.gov/pdffiles1/nij/307346.pdf>. It specifies minimum performance requirements and test methods for the ballistic resistance of body armor used by U.S. law enforcement that is intended to protect the torso against handgun and rifle ammunition. This revised standard supersedes NIJ Standard 0101.06, *Ballistic Resistance of Body Armor*, effective immediately. The primary purpose of this standard will be for use by the NIJ Compliance Testing Program (CTP) for testing, evaluation, and certification of ballistic-resistant body armor. It will also be used by NIJ-approved ballistic testing laboratories and body armor suppliers participating in the NIJ CTP. More information on this standard can be found here: <https://nij.ojp.gov/standard-0101-07>.

NIJ also announces publication of NIJ Standard 0123.00, *Specification for NIJ Ballistic Protection Levels and Associated Test Threats*. The document can be found here: <https://www.ojp.gov/pdffiles1/nij/307347.pdf>. It specifies the NIJ ballistic protection levels and associated test threats identified by U.S. law enforcement as representative of current prevalent threats in the United States. This standard should be used in conjunction with other standards to test and evaluate specific ballistic-resistant equipment, such as ballistic-resistant body armor, against contemporary ballistic threats that pose a life-threatening safety hazard to U.S. law enforcement officers. The NIJ CTP will use this standard for testing, evaluation, and certification of ballistic-resistant body armor using NIJ Standard 0101.07 and other types of ballistic-resistant equipment that may be added to the