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

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 1301.34(a), this is notice that on July 25, 2022, Curia CTS, LLC, 10245 Hickman Mills Drive, Kansas City, Missouri 64137–1418, applied to be registered as an importer of the following basic class(es) of controlled substance(s):

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gamma Hydroxybutyric Acid.</td>
<td>2010</td>
<td>I</td>
</tr>
<tr>
<td>Marihuana Extract.</td>
<td>7350</td>
<td>I</td>
</tr>
<tr>
<td>Marihuana ......</td>
<td>7360</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols.</td>
<td>7370</td>
<td>I</td>
</tr>
</tbody>
</table>

The company plans to import the listed controlled substances as dosage unit products for clinical trial studies. In reference to drug code 7370 (Tetrahydrocannabinols), the company plans to import a synthetic tetrahydrocannabinol. No other activities for these drug codes are 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.

Kristi O’Malley, Assistant Administrator.

[FR Doc. 2022–19394 Filed 9–7–22; 8:45 am]

BILLING CODE P

DEPARTMENT OF JUSTICE
Drug Enforcement Administration

[Docket No. DEA–1076]

Bulk Manufacturer of Controlled Substances Application: Curia Wisconsin, Inc.

AGENCY: Drug Enforcement Administration, Justice.

ACTION: Notice of application.

SUMMARY: Curia Wisconsin, Inc., 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 submit electronic comments on or objections to the issuance of the proposed registration on or before November 7, 2022. Such persons may also file a written request for a hearing on the application on or before November 7, 2022.

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 July 25, 2022, Curia Wisconsin, Inc., 870 Badger Circle, Grafton, Wisconsin 53024–0000, applied to be registered as a bulk manufacturer of the following basic class(es) of controlled substance(s):

<table>
<thead>
<tr>
<th>Controlled substance</th>
<th>Drug code</th>
<th>Schedule</th>
</tr>
</thead>
<tbody>
<tr>
<td>Lysergic acid diethylamide</td>
<td>7315</td>
<td>I</td>
</tr>
<tr>
<td>Tetrahydrocannabinols</td>
<td>7370</td>
<td>I</td>
</tr>
<tr>
<td>4-Bromo-2,5-dimethoxymethylamphetamine</td>
<td>7392</td>
<td>I</td>
</tr>
<tr>
<td>3,4-Methylenedioxymethylamphetamine</td>
<td>7400</td>
<td>I</td>
</tr>
<tr>
<td>3,4-Methylenedioxymethylamphetamine</td>
<td>7405</td>
<td>I</td>
</tr>
<tr>
<td>5-Methoxy-N,N-dimethyltryptamine</td>
<td>7431</td>
<td>I</td>
</tr>
<tr>
<td>Dimethyltryptamine</td>
<td>7435</td>
<td>I</td>
</tr>
<tr>
<td>Psilocybin</td>
<td>7437</td>
<td>I</td>
</tr>
<tr>
<td>Psilocyn</td>
<td>7438</td>
<td>I</td>
</tr>
<tr>
<td>Methylenedioxylate</td>
<td>1724</td>
<td>II</td>
</tr>
<tr>
<td>Nabamine</td>
<td>7379</td>
<td>II</td>
</tr>
<tr>
<td>ANPP (4-Anilino-N-phenethyl-4-piperidine)</td>
<td>8333</td>
<td>II</td>
</tr>
<tr>
<td>Fentanyl</td>
<td>9801</td>
<td>II</td>
</tr>
</tbody>
</table>
The company plans to bulk manufacture the listed controlled substances for the purpose of analytical reference standards or for sale to its customers. In reference to the drug code 7370 (Tetrahydrocannabinols), the company plans to bulk manufacture as synthetic. No other activity for this drug code is authorized for this registration.

Kristi O’Malley, 
Assistant Administrator.

DEPARTMENT OF LABOR

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Application for Prevailing Wage Determination

ACTION: Notice of availability; request for comments.

SUMMARY: The Department of Labor (DOL) is submitting this Employment and Training Administration (ETA)-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 the agency receives on or before October 11, 2022.

AFFECTED PUBLIC: The Department of Labor (DOL) is submitting this Employment and Training Administration (ETA)-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.

DEPARTMENT OF LABOR

Mine Safety and Health Administration

Petition for Modification of Application of Existing Mandatory Safety Standards

AGENCY: Mine Safety and Health Administration, Labor.

ACTION: Notice.

SUMMARY: This notice is a summary of a petition for modification submitted to the Mine Safety and Health Administration (MSHA) by the party listed below.

DATES: All comments on the petition must be received by MSHA’s Office of Standards, Regulations, and Variances on or before October 11, 2022.

ADDRESS: You may submit comments identified by Docket No. MSHA—2022–046 by any of the following methods:

2. Fax: 202–693–9441.
3. Email: petitioncomments@dol.gov.

Attention: S. Aronie Noe, Director, Office of Standards, Regulations, and Variances. Persons delivering documents are required to check in at the receptionist’s desk in Suite 4E401. Individuals may inspect copies of the petition and comments during normal business hours at the address listed above. Before visiting MSHA in person, call 202–693–9455 to make an appointment, in keeping with the Department of Labor’s COVID–19 policy. Special health precautions may be required.

FOR FURTHER INFORMATION CONTACT: S. Aronie Noe, Office of Standards, Regulations, and Variances. Persons delivering documents are required to check in at the receptionist’s desk in Suite 4E401. Individuals may inspect copies of the petition and comments during normal business hours at the address listed above. Before visiting MSHA in person, call 202–693–9455 to make an appointment, in keeping with the Department of Labor’s COVID–19 policy. Special health precautions may be required.

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SUPPLEMENTARY INFORMATION: Section 101(c) of the Federal Mine Safety and

Total Estimated Annual Time Burden: 148,629 hours.
Total Estimated Annual Other Costs Burden: $213,953.

Authority: 44 U.S.C. 3507(a)(1)(D)
Dated: September 1, 2022.

Mara Blumenthal, 
Senior PRA Analyst.

[FR Doc. 2022–19359 Filed 9–7–22; 8:45 am]
BILLING CODE 4510–FP–P