HB2660 FULLPCS1 TJ Marti-GRS 2/24/2021 12:19:01 pm

COMMITTEE AMENDMENT

HOUSE OF REPRESENTATIVES
State of Oklahoma

SPEAK	ER:						
CHAIR	:						
I move to	amend <u>HB2</u>	660					'
Page		Section		Line	es	the printed	
					Of th	ne Engrossed	d Bill
inserting	in lieu th	ereof the fo	ollowing la	inguage :	•		
AMEND TITLE	TO CONFORM T	O AMENDMENTS					
Adopted:			Am	endment	submitted b	y: TJ Marti ————	

Reading Clerk

1 STATE OF OKLAHOMA 2 1st Session of the 58th Legislature (2021) 3 PROPOSED COMMITTEE SUBSTITUTE 4 FOR HOUSE BILL NO. 2660 By: Echols 5 6 7 PROPOSED COMMITTEE SUBSTITUTE An Act relating to medical marijuana; amending 8 Section 4, State Question No. 788, Initiative 9 Petition No. 412 (63 O.S. Supp. 2020, Section 423), which relates to medical marijuana processor 10 licensing requirements; providing for the issuance of volatile and nonvolatile processor licenses; updating language; amending Section 17, Chapter 11, O.S.L. 11 2019, as amended by Section 4, Chapter 312, O.S.L. 2019 (63 O.S. Supp. 2020, Section 427.17), which 12 relates to the Oklahoma Medical Marijuana and Patient 1.3 Protection Act; modifying test batch requirements for licensed medical marijuana commercial growers and 14 processors; defining certain term; and providing an effective date. 15 16 17 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 18 Section 4, State Question No. 788, SECTION 1. AMENDATORY 19 Initiative Petition No. 412 (63 O.S. Supp. 2020, Section 423), is 20 amended to read as follows: 2.1 Section 423. A. The Oklahoma State Department of Health shall 22 within thirty (30) days of passage of this initiative, make 23 available, on their its website, in an easy-to-find location, an 24 application for a medical marijuana processing processor license.

1 The Department shall be authorized to issue two types of medical 2 marijuana processor licenses:

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- 1. Nonvolatile, which involves using any solvent in the extraction process that is not a volatile solvent, including carbon dioxide; and
- 2. Volatile, which involves using any solvent that is or produces a flammable gas or vapor that, when present in the air in sufficient quantities, will create explosive or ignitable mixtures and may also include extraction using nonvolatile solvents or no solvents.
- 11 The application fee for a nonvolatile or volatile medical marijuana 12 processor license shall be Two Thousand Five Hundred Dollars 13 (\$2,500.00) and methods. Methods of payment will shall be provided 14 on the website. The Oklahoma State Department of Health shall have 15 two (2) weeks to review the application, approve or reject the 16 application, and mail the approval/rejection letter (if rejected, 17 stating reasons for rejection) to the applicant.
 - The Oklahoma State Department of Health must shall approve all applications which meet the following criteria:
 - Applicant must be age twenty-five (25) or older;
- 21 Any applicant, applying as an individual, must show 22 residency in the State of Oklahoma;
- 23 3. All applying entities must show that all members, managers, and board members are Oklahoma residents;

- 4. An applying entity may show ownership of non-Oklahoma residents, but that percentage ownership may not exceed twenty-five percent (25%);
- 5. All applying individuals or entities must be registered to conduct business in the State of Oklahoma; and
 - 6. All applicants must disclose all ownership.

- 7. Applicant(s) An applicant with only a nonviolent felony conviction(s) in the last two (2) years, or any other felony conviction in five (5) years, inmates, or any person currently incarcerated may not qualify for a medical marijuana processing processor license.
- C. A licensed medical marijuana processor may take marijuana plants and distill or process these plants into concentrates, edibles, and other forms for consumption. As required by subsection D of this section, the Oklahoma State Department of Health will shall, within sixty (60) days of passage of this initiative, make available a set of standards which will be used by licensed medical marijuana processors in the preparation of edible marijuana products. This should be in line with current food preparation guidelines and no excessive or punitive rules may be established by the Oklahoma State Department of Health. Once a year, the Oklahoma State Department of Health. Once a year, the Oklahoma State Department of Health may inspect a processing operation and determine its compliance with the preparation standards. If deficiencies are found, a written report of deficiency will the

1 deficiencies shall be issued to the licensed medical marijuana The licensed medical marijuana processor will shall have processor. one (1) month to correct the deficiency or be subject to a fine of 3 Five Hundred Dollars (\$500.00) for each deficiency. A licensed 4 5 medical marijuana processor may sell marijuana products it creates to a licensed retailer, or any other licensed medical marijuana 6 7 processor. Further, these sales will shall be considered wholesale sales and not subject to taxation. Under no circumstances may a 8 licensed medical marijuana processor sell marijuana, or any 10 marijuana product, directly to a medical marijuana license holder 11 patient licensee or caregiver licensee. However, a licensed medical 12 marijuana processor may process cannabis into a concentrated form, 13 for a medical license holder, marijuana patient licensee for a fee. 14 Processors will Licensed medical marijuana processors shall be 15 required to complete a monthly yield and sales report to the 16 Oklahoma State Department of Health. This report will be due on the 17 15th fifteenth of each month and provide reporting on the previous 18 This report will shall detail the amount of marijuana 19 purchased in pounds, the amount of marijuana cooked or processed in 20 pounds, and the amount of waste in pounds. Additionally, this 21 report will shall show total wholesale sales in dollars. 22 Oklahoma State Department of Health will shall have oversight and 23 auditing responsibilities to ensure that all marijuana being grown 24 is accounted for. A licensed medical marijuana processor will shall

only be subject to a penalty if a gross discrepancy exists and cannot be explained. Penalties for fraudulent reporting occurring within any 2 year two-year time period will shall be an initial fine of Five Thousand Dollars (\$5,000.00) (first) for a first offense and revocation of licensing (second) the medial marijuana processor license for a second offense.

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- The Department shall oversee inspection and compliance of D. licensed medical marijuana processors producing products with marijuana as an additive. The Oklahoma State Department of Health will shall be compelled to, within thirty (30) days of passage of this initiative, appoint a board of twelve (12) Oklahoma residents, who are marijuana industry experts, to create a list of food safety standards for processing and handling medical marijuana in Oklahoma. These standards will shall be adopted by the agency Department and the agency Department can enforce these standards for licensed medical marijuana processors. The agency will Department shall develop a standards review procedure and these standards can may be altered by calling another board of twelve (12) Oklahoma marijuana industry experts. A signed letter of twenty (20) operating licensed medical marijuana processors would constitute a need for a new board and standard standards review.
- E. If it becomes permissible, under federal law, marijuana may be moved across state lines.

- F. Any device used for the consumption of medical marijuana shall be considered legal to be sold, manufactured, distributed, and possessed. No merchant, wholesaler, manufacturer, or individual may unduly be harassed or prosecuted for selling, manufacturing, or possession of medical marijuana paraphernalia.
- SECTION 2. AMENDATORY Section 17, Chapter 11, O.S.L. 2019, as amended by Section 4, Chapter 312, O.S.L. 2019 (63 O.S. Supp. 2020, Section 427.17), is amended to read as follows:

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- Section 427.17 A. There is hereby created a medical marijuana testing laboratory license as a category of the medical marijuana business license. The Authority is hereby enabled to monitor, inspect and audit a licensed testing laboratory under this act.
- B. The Authority is hereby authorized to contract with a private laboratory for the purpose of conducting compliance testing of medical marijuana testing laboratories licensed in this state.

 Any such laboratory under contract for compliance testing shall be prohibited from conducting any other commercial medical marijuana testing in this state.
- C. The Authority shall have the authority to develop acceptable testing and research practices, including but not limited to testing, standards, quality control analysis, equipment certification and calibration, and chemical identification and substances used in bona fide research methods so long as it complies with this act.

- D. A person who is a direct beneficial owner or an indirect beneficial owner of a medical marijuana dispensary, medical marijuana commercial grower, or medical marijuana processor shall not be an owner of a laboratory.
- E. A laboratory and a laboratory applicant shall comply with all applicable local ordinances, including but not limited to zoning, occupancy, licensing and building codes.
- F. A separate license shall be required for each specific laboratory.
- G. A medical marijuana testing laboratory license may be issued to a person who performs testing and research on medical marijuana and medical marijuana products for medical marijuana businesses, medical marijuana research facilities, medical marijuana education facilities, and testing and research on marijuana and marijuana products grown or produced by a patient or caregiver on behalf of a patient, upon verification of registration. No state-approved medical marijuana testing facility shall operate unless a medical laboratory director is on site during operational hours.
- H. A laboratory applicant shall comply with the application requirements of this section and shall submit such other information as required for a medical marijuana business applicant, in addition to any information the Authority may request for initial approval and periodic evaluations during the approval period.

I. A medical marijuana testing laboratory may accept samples of medical marijuana, medical marijuana concentrate or medical marijuana product from a medical marijuana business for testing and research purposes only, which purposes may include the provision of testing services for samples submitted by a medical marijuana business for product development. The Department may require a medical marijuana business to submit a sample of medical marijuana, medical marijuana concentrate or medical marijuana product to a medical marijuana testing laboratory upon demand.

- J. A medical marijuana testing laboratory may accept samples of medical marijuana, medical marijuana concentrate or medical marijuana product from an individual person for testing only under the following conditions:
- 1. The individual person is a patient or caregiver pursuant to this act or is a participant in an approved clinical or observational study conducted by a research facility; and
- 2. The medical marijuana testing laboratory shall require the patient or caregiver to produce a valid patient license and current and valid photo identification.
- K. A medical marijuana testing laboratory may transfer samples to another medical marijuana testing laboratory for testing. All laboratory reports provided to or by a medical marijuana business or to a patient or caregiver shall identify the medical marijuana testing laboratory that actually conducted the test.

L. A medical marijuana testing laboratory may utilize a licensed medical marijuana transporter to transport samples of medical marijuana, medical marijuana concentrate and medical marijuana product for testing, in accordance with this act and the rules adopted pursuant thereto, between the originating medical marijuana business requesting testing services and the destination laboratory performing testing services.

- M. The medical marijuana testing laboratory shall establish policies to prevent the existence of or appearance of undue commercial, financial or other influences that may diminish the competency, impartiality and integrity of the testing processes or results of the laboratory, or that may diminish public confidence in the competency, impartiality and integrity of the testing processes or results of the laboratory. At a minimum, employees, owners or agents of a medical marijuana testing laboratory who participate in any aspect of the analysis and results of a sample are prohibited from improperly influencing the testing process, improperly manipulating data, or improperly benefiting from any ongoing financial, employment, personal or business relationship with the medical marijuana business that provided the sample.
- N. The Department, pursuant to rules promulgated by the State Commissioner of Health, shall develop standards, policies and procedures as necessary for:

1. The cleanliness and orderliness of a laboratory premises and the location of the laboratory in a secure location, and inspection, cleaning and maintenance of any equipment or utensils used for the analysis of test samples;

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- 2. Testing procedures, testing standards for cannabinoid and terpenoid potency and safe levels of contaminants, and remediation procedures;
- 3. Controlled access areas for storage of medical marijuana and medical marijuana product test samples, waste and reference standards:
- 4. Records to be retained and computer systems to be utilized by the laboratory;
 - 5. The possession, storage and use by the laboratory of reagents, solutions and reference standards;
 - 6. A certificate of analysis (COA) for each lot of reference standard;
 - 7. The transport and disposal of unused marijuana, marijuana products and waste;
 - 8. The mandatory use by a laboratory of an inventory tracking system to ensure all test batches or samples containing medical marijuana, medical marijuana concentrate or medical marijuana products are identified and tracked from the point they are transferred from a medical marijuana business, a patient or a caregiver through the point of transfer, destruction or disposal.

The inventory tracking system reporting shall include the results of any tests that are conducted on medical marijuana, medical marijuana concentrate or medical marijuana product;

9. Standards of performance;

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- 10. The employment of laboratory personnel;
- 6 11. A written standard operating procedure manual to be 7 maintained and updated by the laboratory;
 - 12. The successful participation in a Department-approved proficiency testing program for each testing category listed in this section, in order to obtain and maintain certification;
 - 13. The establishment of and adherence to a quality assurance and quality control program to ensure sufficient monitoring of laboratory processes and quality of results reported;
 - 14. The establishment by the laboratory of a system to document the complete chain of custody for samples from receipt through disposal;
 - 15. The establishment by the laboratory of a system to retain and maintain all required records, including business records, and processes to ensure results are reported in a timely and accurate manner; and
 - 16. Any other aspect of laboratory testing of medical marijuana or medical marijuana product deemed necessary by the Department.
 - O. A medical marijuana testing laboratory shall promptly provide the Department or designee of the Department access to a

- report of a test and any underlying data that is conducted on a

 sample at the request of a medical marijuana business or qualified

 patient. A medical marijuana testing laboratory shall also provide

 access to the Department or designee of the Department to laboratory

 premises and to any material or information requested by the

 Department to determine compliance with the requirements of this

 section.
 - P. A medical marijuana testing laboratory shall retain all results of laboratory tests conducted on marijuana or products for a period of at least two (2) years and shall make them available to the Department upon request.
 - Q. A medical marijuana testing laboratory shall test samples from each harvest batch or product batch, as appropriate, of medical marijuana, medical marijuana concentrate and medical marijuana product for each of the following categories of testing, consistent with standards developed by the Commissioner:
- 17 1. Microbials;

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- Mycotoxins;
 - 3. Residual solvents;
- 20 4. Pesticides;
- 5. Tetrahydrocannabinol (THC) and other cannabinoid potency;
- 22 | 6. Terpenoid potency; and
- 7. Heavy metals.

R. A test batch shall not exceed ten (10) pounds of usable marijuana or medical marijuana product, as appropriate. A grower shall separate each harvest lot of usable marijuana into harvest batches containing no more than ten (10) fifty (50) pounds. A processor shall separate each medical marijuana production lot into production batches containing no more than ten (10) two (2) pounds of distillate and for final products, a maximum batch size of fifty thousand (50,000) milligrams of THC. As used in this subsection, "final products" shall include, but not be limited to, cookies, brownies, candies, gummies and chocolates.

- S. Medical marijuana testing laboratory licensure shall be contingent upon successful on-site inspection, successful participation in proficiency testing and ongoing compliance with the applicable requirements in this section.
- T. A medical marijuana testing laboratory shall be inspected prior to initial licensure and annually thereafter by an inspector approved by the Authority.
- U. Beginning on a date determined by the Commissioner, not later than January 1, 2020, medical marijuana testing laboratory licensure shall be contingent upon accreditation by the NELAC Institute (TNI), ANSI/ASQ National Accreditation Board or another accrediting body approved by the Commissioner, and any applicable standards as determined by the Department.

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        V. A commercial grower shall not transfer or sell medical
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    marijuana and a processor shall not transfer, sell or process into a
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    concentrate or product any medical marijuana, medical marijuana
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    concentrate or medical marijuana product unless samples from each
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    harvest batch or production batch from which that medical marijuana,
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    medical marijuana concentrate or medical marijuana product was
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    derived has been tested by a medical marijuana testing facility for
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    contaminants and passed all contaminant tests required by this act.
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        SECTION 3.
                    This act shall become effective November 1, 2021.
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