## <DateSubmitted>

# HOUSE OF REPRESENTATIVES CONFERENCE COMMITTEE REPORT

Mr. President: Mr. Speaker:

The Conference Committee, to which was referred

### **HB4056**

- Marti of the House and Paxton of the Senate By:
- Title: Medical marijuana; Laboratory recommendations, standards and operating procedures; Oklahoma Medical Marijuana Authority; rules; emergency.

Together with Engrossed Senate Amendments thereto, beg leave to report that we have had the same under consideration and herewith return the same with the following recommendations:

- 1. That the Senate recede from its amendment; and
- 2. That the attached Conference Committee Substitute be adopted.

Respectfully submitted,

# SENATE CONFEREES

Paxton	 
Leewright	
Taylor	
Rosino	
Dossett (J.J.)	

1	STATE OF OKLAHOMA		
2	2nd Session of the 58th Legislature (2022)		
3	CONFERENCE COMMITTEE SUBSTITUTE		
4	FOR ENGROSSED HOUSE BILL NO. 4056 By: Marti, Davis, Talley, and		
5	McDugle of the House		
6	and		
7	Paxton of the Senate		
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11	CONFERENCE COMMITTEE SUBSTITUTE		
12	An Act relating to medical marijuana; amending 63 O.S. 2021, Section 427.17, which relates to the		
13	Oklahoma Medical Marijuana and Patient Protection Act; directing certain entities to provide recommendations, standards and operating procedures to the Oklahoma Medical Marijuana Authority; directing the Authority to promulgate rules; requiring licensed medical marijuana testing laboratories to comply with rules; providing for license revocation; and declaring an emergency.		
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17	Ticense revocation, and declaring an emergency.		
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19	BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:		
20	SECTION 1. AMENDATORY 63 O.S. 2021, Section 427.17, is		
21	amended to read as follows:		
22	Section 427.17 A. There is hereby created a medical marijuana		
23	testing laboratory license as a category of the medical marijuana		
24	business license. The Oklahoma Medical Marijuana Authority is		

hereby enabled to monitor, inspect and audit a licensed testing
 laboratory under the Oklahoma Medical Marijuana and Patient
 Protection Act.

4 1. The Authority is hereby authorized to contract with a Β. 5 private laboratory for the purpose of conducting compliance testing of medical marijuana testing laboratories licensed in this state. 6 Any such laboratory under contract for compliance testing shall be 7 prohibited from conducting any other commercial medical marijuana 8 9 testing in this state. The laboratory the Authority contracts with 10 for compliance testing shall not employ, or be owned by, the 11 following:

12 <del>1. Any</del>

any individual that has a direct or indirect interest
 in a licensed medical marijuana business;

15 <del>2. Any</del>

16 any individual or his or her spouse, parent, child, b. 17 spouse of a child, sibling or spouse of a sibling that 18 has an application for a medical marijuana business 19 license pending before the Department or is a member 20 of the board of directors of a medical marijuana 21 business, or is an individual financially interested 22 in any licensee or medical marijuana business located 23 within this state.

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1	2. The private laboratory under contract with the Authority for
2	compliance testing and a board or committee comprised of licensed
3	Oklahoma medical marijuana laboratories currently accredited by the
4	International Organization for Standardization (ISO) shall provide
5	to the Authority its recommendations for all equipment and standards
6	to be utilized by licensed medical marijuana testing laboratories
7	when testing samples of medical marijuana, medical marijuana
8	concentrate, and medical marijuana products as well as standard
9	operating procedures when extracting and testing medical marijuana,
10	medical marijuana concentrate, and medical marijuana products. The
11	recommendations shall be submitted to the Authority no later than
12	June 1, 2023. The Authority shall have ninety (90) days from the
13	date it receives the recommendations to promulgate new rules or
14	modify its current rules for laboratory standards and testing.
15	Beginning June 1, 2024, medical marijuana testing laboratories
16	renewing their medical marijuana business license shall be subject
17	to and comply with any new or modified rules relating to the testing
18	of medical marijuana, medical marijuana concentrate, and medical
19	marijuana products. The refusal or failure of a medical marijuana
20	testing laboratory licensee to comply with new or modified rules
21	relating to laboratory standards and testing procedures promulgated
22	under the provisions of this paragraph shall result in the permanent
23	revocation of the medical marijuana testing laboratory license.

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C. The Authority shall develop acceptable testing practices
 including, but not limited to, testing, standards, quality control
 analysis, equipment certification and calibration, and chemical
 identification and substances used.

D. A person who is a direct beneficial owner of a medical
marijuana dispensary, medical marijuana commercial grower or medical
marijuana processor shall not be an owner of a laboratory.

8 E. A laboratory and a laboratory applicant shall comply with
9 all applicable local ordinances including, but not limited to,
10 zoning, occupancy, licensing and building codes.

F. A separate license shall be required for each specificlaboratory.

13 G. A medical marijuana testing laboratory license may be issued 14 to a person who performs testing on medical marijuana and medical 15 marijuana products for medical marijuana businesses, medical 16 marijuana research facilities, medical marijuana education 17 facilities, and testing on marijuana and marijuana products grown or 18 produced by a patient or caregiver on behalf of a patient, upon 19 verification of registration. A medical marijuana testing 20 laboratory may also conduct research related to the development and 21 improvement of its testing practices and procedures. No state-22 approved medical marijuana testing facility shall operate unless a 23 medical laboratory director is on site during operational hours.

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H. Laboratory applicants and licensees 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 6 medical marijuana, medical marijuana concentrate or medical 7 marijuana product from a medical marijuana business, medical 8 9 marijuana research facility or medical marijuana education facility 10 for testing purposes only, which purposes may include the provision 11 of testing services for samples submitted by a medical marijuana 12 business for product development. The Department may require a 13 medical marijuana business to submit a sample of medical marijuana, 14 medical marijuana concentrate or medical marijuana product to a 15 medical marijuana testing or quality assurance laboratory upon 16 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:

21 1. The individual person is a patient or caregiver pursuant to 22 the Oklahoma Medical Marijuana and Patient Protection Act or is a 23 participant in an approved clinical or observational study conducted 24 by a research facility; and

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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.

9 L. A medical marijuana testing laboratory may utilize a 10 licensed medical marijuana transporter to transport samples of 11 medical marijuana, medical marijuana concentrate and medical 12 marijuana product for testing, in accordance with the Oklahoma 13 Medical Marijuana and Patient Protection Act and the rules adopted 14 pursuant thereto, between the originating medical marijuana business 15 requesting testing services and the destination laboratory 16 performing testing services.

17 Μ. The medical marijuana testing laboratory shall establish 18 policies to prevent the existence of or appearance of undue 19 commercial, financial or other influences that may diminish the 20 competency, impartiality and integrity of the testing processes or 21 results of the laboratory, or that may diminish public confidence in 22 the competency, impartiality and integrity of the testing processes 23 or results of the laboratory. At a minimum, employees, owners or 24 agents of a medical marijuana testing laboratory who participate in

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1 any aspect of the analysis and results of a sample are prohibited from improperly influencing the testing process, improperly 2 manipulating data or improperly benefiting from any ongoing 3 4 financial, employment, personal or business relationship with the 5 medical marijuana business that provided the sample. A medical marijuana testing laboratory shall not test samples for any medical 6 7 marijuana business in which an owner, employee or agent of the medical marijuana testing laboratory has any form of ownership or 8 9 financial interest in the medical marijuana business.

10 N. The Department, pursuant to rules promulgated by the State 11 Commissioner of Health, shall develop standards, policies and 12 procedures as necessary for:

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;

17 2. Testing procedures, testing standards for cannabinoid and 18 terpenoid potency and safe levels of contaminants, and remediation 19 procedures;

20 3. Controlled access areas for storage of medical marijuana and 21 medical marijuana product test samples, waste and reference 22 standards;

4. Records to be retained and computer systems to be utilizedby the laboratory;

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5. The possession, storage and use by the laboratory of
 reagents, solutions and reference standards;

3 6. A certificate of analysis (COA) for each lot of reference4 standard;

5 7. The transport and disposal of unused marijuana, marijuana6 products and waste;

7 8. The mandatory use by a laboratory of an inventory tracking system to ensure all harvest and production batches or samples 8 9 containing medical marijuana, medical marijuana concentrate or 10 medical marijuana products are identified and tracked from the point 11 they are transferred from a medical marijuana business, a patient or 12 a caregiver through the point of transfer, destruction or disposal. 13 The inventory tracking system reporting shall include the results of 14 any tests that are conducted on medical marijuana, medical marijuana 15 concentrate or medical marijuana product;

16 9. Standards of performance;

17 10. The employment of laboratory personnel;

18 11. A written standard operating procedure manual to be 19 maintained and updated by the laboratory;

20 12. The successful participation in a Department-approved 21 proficiency testing program for each testing category listed in this 22 section, in order to obtain and maintain certification;

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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;

4 14. The immediate recall of medical marijuana or medical
5 marijuana products that test above allowable thresholds or are
6 otherwise determined to be unsafe;

7 15. The establishment by the laboratory of a system to document 8 the complete chain of custody for samples from receipt through 9 disposal;

10 16. The establishment by the laboratory of a system to retain 11 and maintain all required records, including business records, and 12 processes to ensure results are reported in a timely and accurate 13 manner; and

14 17. Any other aspect of laboratory testing of medical marijuana15 or medical marijuana product deemed necessary by the Department.

16 A medical marijuana testing laboratory shall promptly Ο. 17 provide the Department or designee of the Department access to a 18 report of a test and any underlying data that is conducted on a 19 sample at the request of a medical marijuana business or qualified 20 patient. A medical marijuana testing laboratory shall also provide 21 access to the Department or designee of the Department to laboratory 22 premises and to any material or information requested by the 23 Department to determine compliance with the requirements of this 24 section.

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P. A medical marijuana testing laboratory shall retain all
 results of laboratory tests conducted on marijuana or products for a
 period of at least seven (7) 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:

10 1. Microbials;

11 2. Mycotoxins;

12 3. Residual solvents;

13 4. Pesticides;

14 5. Tetrahydrocannabinol (THC) and other cannabinoid potency;

15 6. Terpenoid type and concentration; and

16 7. Heavy metals.

17 R. A licensed medical marijuana testing laboratory shall test 18 each individual harvest batch. A grower shall separate each harvest 19 lot of usable marijuana into harvest batches containing no more than 20 fifteen (15) pounds, with the exception of any plant material to be 21 sold to a licensed processor for the purposes of turning the plant 22 material into concentrate which may be separated into harvest 23 batches of no more than fifty (50) pounds. A processor shall 24 separate each medical marijuana production lot into production

1 batches containing no more than four (4) liters of concentrate or nine (9) pounds for nonliquid products, and for final products, the 2 Oklahoma Medical Marijuana Authority shall be authorized to 3 4 promulgate rules on final products as necessary. Provided, however, 5 the Authority shall not require testing of final products less often than every one thousand (1,000) grams of THC. As used in this 6 7 subsection, "final products" shall include, but not be limited to, 8 cookies, brownies, candies, gummies, beverages and chocolates.

9 S. Medical marijuana testing laboratory licensure shall be
10 contingent upon successful on-site inspection, successful
11 participation in proficiency testing and ongoing compliance with the
12 applicable requirements in this section.

T. A medical marijuana testing laboratory shall be inspected prior to initial licensure and up to two (2) times per year thereafter by an inspector approved by the Authority. The Authority may enter the licensed premises of a testing laboratory to conduct investigations and additional inspections when the Authority believes an investigation or additional inspection is necessary due to a possible violation of applicable laws, rules or regulations.

U. Medical marijuana testing laboratories shall obtain accreditation by an accrediting body approved by the Commissioner within one (1) year of the date the initial license is issued. Renewal of any medical marijuana testing laboratory license shall be contingent upon accreditation in accordance with this subsection.

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All medical marijuana testing laboratories shall obtain
 accreditation prior to applying for and receiving a medical
 marijuana testing laboratory license.

V. Unless authorized by the provisions of this section, a 4 5 commercial grower shall not transfer or sell medical marijuana and a processor shall not transfer, sell or process into a concentrate or 6 7 product any medical marijuana, medical marijuana concentrate or medical marijuana product unless samples from each harvest batch or 8 9 production batch from which that medical marijuana, medical 10 marijuana concentrate or medical marijuana product was derived has been tested by a medical marijuana testing laboratory and passed all 11 contaminant tests required by the Oklahoma Medical Marijuana and 12 13 Patient Protection Act and applicable laws, rules and regulations. 14 A licensed commercial grower may transfer medical marijuana that has 15 failed testing to a licensed processor only for the purposes of 16 decontamination or remediation and only in accordance with the 17 provisions of the Oklahoma Medical Marijuana and Patient Protection 18 Act and the rules and regulations of the Department. Remediated and 19 decontaminated medical marijuana may be returned only to the 20 originating licensed commercial grower.

W. Kief shall not be transferred or sold except as authorized
in the rules and regulations of the Department.

23 SECTION 2. It being immediately necessary for the preservation 24 of the public peace, health or safety, an emergency is hereby

1	declared to exist, by reason whereof this act shall take effect and
2	be in full force from and after its passage and approval.
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