## 1 SENATE FLOOR VERSION 2 February 22, 2021 3 SENATE BILL NO. 680 By: Daniels 4 5 6 7 An Act relating to medical marijuana; amending Section 2, Chapter 11, O.S.L. 2019, as last amended by Section 48, Chapter 161, O.S.L. 2020 (63 O.S. 8 Supp. 2020, Section 427.2), which relates to 9 definitions used in the Oklahoma Medical Marijuana and Patient Protection Act; modifying definition; amending Section 17, Chapter 11, O.S.L. 2019, as 10 amended by Section 4, Chapter 312, O.S.L. 2019 (63 11 O.S. Supp. 2020, Section 427.17), which relates to medical marijuana testing laboratory license; 12 requiring testing of medical marijuana waste prior to transfer; requiring separation of medical marijuana waste into waste batches; modifying provisions to 13 include medical marijuana waste; clarifying language; updating statutory references; and providing an 14 effective date. 15 16 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA: 17 Section 2, Chapter 11, O.S.L. 18 SECTION 1. AMENDATORY 2019, as last amended by Section 48, Chapter 161, O.S.L. 2020 (63 19 O.S. Supp. 2020, Section 427.2), is amended to read as follows: 20 Section 427.2. As used in this act the Oklahoma Medical 21 Marijuana and Patient Protection Act: 22 "Advertising" means the act of providing consideration for 23 1. 24 the publication, dissemination, solicitation, or circulation, of

- visual, oral, or written communication to induce directly or indirectly any person to patronize a particular medical marijuana business, or to purchase particular medical marijuana or a medical marijuana product. Advertising includes marketing, but does not
  - 2. "Authority" means the Oklahoma Medical Marijuana Authority;
  - 3. "Batch number" means a unique numeric or alphanumeric identifier assigned prior to testing to allow for inventory tracking and traceability;
  - 4. "Cannabinoid" means any of the chemical compounds that are active principles of marijuana;
  - 5. "Caregiver" means a family member or assistant who regularly looks after a medical marijuana license holder whom a physician attests needs assistance;
    - 6. "Child-resistant" means special packaging that is:
      - a. designed or constructed to be significantly difficult for children under five (5) years of age to open and not difficult for normal adults to use properly as defined by 16 C.F.R. 1700.15 (1995) and 16 C.F.R. 1700.20 (1995),
      - b. opaque so that the outermost packaging does not allow the product to be seen without opening the packaging material, and

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include packaging and labeling;

| 1 | С. | resealable to maintain its child-resistant          |
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| 2 |    | effectiveness for multiple openings for any product |
| 3 |    | intended for more than a single use or containing   |
| 4 |    | multiple servings;                                  |

- 7. "Clone" means a nonflowering plant cut from a mother plant that is capable of developing into a new plant and has shown no signs of flowering;
  - 8. "Commissioner" means the State Commissioner of Health;
- 9. "Complete application" means a document prepared in accordance with the provisions set forth in this act the Oklahoma Medical Marijuana and Patient Protection Act, rules promulgated pursuant thereto, and the forms and instructions provided by the Department, including any supporting documentation required and the applicable license application fee;
  - 10. "Department" means the State Department of Health;
- 11. "Director" means the Executive Director of the Oklahoma Medical Marijuana Authority;
- 12. "Dispense" means the selling of medical marijuana or a medical marijuana product to a qualified patient or the designated caregiver of the patient that is packaged in a suitable container appropriately labeled for subsequent administration to or use by a qualifying patient;
- 13. "Dispensary" means a medical marijuana dispensary, an entity that has been licensed by the Department pursuant to this act

- 1 the Oklahoma Medical Marijuana and Patient Protection Act to 2 purchase medical marijuana or medical marijuana products from a 3 licensed medical marijuana commercial grower or medical marijuana processor, sell medical marijuana or medical marijuana products to 4 5 patients and caregivers as defined under this act Section 427.1 et seq. of this title, or sell or transfer products to another 6 7 dispensary;
- 14. "Edible medical marijuana product" means any medical-8 9 marijuana-infused product for which the intended use is oral 10 consumption including, but not limited to, any type of food, drink
  - 15. "Entity" means an individual, general partnership, limited partnership, limited liability company, trust, estate, association, corporation, cooperative, or any other legal or commercial entity;
    - 16. "Flower" means the reproductive organs of the marijuana or cannabis plant referred to as the bud or parts of the plant that are harvested and used to consume in a variety of medical marijuana products;
    - 17. "Flowering" means the reproductive state of the marijuana or cannabis plant in which there are physical signs of flower or budding out of the nodes of the stem;
- 18. "Food-based medical marijuana concentrate" means a medical 22 marijuana concentrate that was produced by extracting cannabinoids from medical marijuana through the use of propylene glycol,

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or pill;

1 glycerin, butter, olive oil, coconut oil or other typical food-safe cooking fats;

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- 19. "Good cause" for purposes of an initial, renewal or reinstatement license application, or for purposes of discipline of a licensee, means:
  - a. the licensee or applicant has violated, does not meet, or has failed to comply with any of the terms, conditions or provisions of the act, any rules promulgated pursuant thereto, or any supplemental relevant state or local law, rule or regulation,
  - b. the licensee or applicant has failed to comply with any special terms or conditions that were placed upon the license pursuant to an order of the State Department of Health, Oklahoma Medical Marijuana Authority or the municipality, or
  - c. the licensed premises of a medical marijuana business or applicant have been operated in a manner that adversely affects the public health or welfare or the safety of the immediate vicinity in which the establishment is located;
- 20. "Harvest batch" means a specifically identified quantity of medical marijuana that is uniform in strain, cultivated utilizing the same cultivation practices, harvested at the same time from the same location and cured under uniform conditions;

21. "Harvested marijuana" means post-flowering medical marijuana not including trim, concentrate or waste;

- 22. "Heat- or pressure-based medical marijuana concentrate" means a medical marijuana concentrate that was produced by extracting cannabinoids from medical marijuana through the use of heat or pressure;
- 23. "Immature plant" means a nonflowering marijuana plant that has not demonstrated signs of flowering;
- 24. "Inventory tracking system" means the required tracking system that accounts for medical marijuana from either the seed or immature plant stage until the medical marijuana or medical marijuana product is sold to a patient at a medical marijuana dispensary, transferred to a medical marijuana research facility, destroyed by a medical marijuana business or used in a research project by a medical marijuana research facility;
- 25. "Licensed patient" or "patient" means a person who has been issued a medical marijuana patient license by the State Department of Health or Oklahoma Medical Marijuana Authority;
- 26. "Licensed premises" means the premises specified in an application for a medical marijuana business license, medical marijuana research facility license or medical marijuana education facility license pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act that are owned or in possession of the licensee and within which the licensee is authorized to cultivate,

manufacture, distribute, sell, store, transport, test or research
medical marijuana or medical marijuana products in accordance with
the provisions of this act the Oklahoma Medical Marijuana and
Patient Protection Act and rules promulgated pursuant thereto;

- 27. "Manufacture" means the production, propagation, compounding or processing of a medical marijuana product, excluding marijuana plants, either directly or indirectly by extraction from substances of natural or synthetic origin, or independently by means of chemical synthesis, or by a combination of extraction and chemical synthesis;
- 28. "Marijuana" shall have the same meaning as such term is defined in Section 2-101 of Title 63 of the Oklahoma Statutes;
- 29. "Material change" means any change that would require a substantive revision to the standard operating procedures of a licensee for the cultivation or production of medical marijuana, medical marijuana concentrate or medical marijuana products;
- 30. "Mature plant" means a harvestable female marijuana plant that is flowering;
  - 31. "Medical marijuana business (MMB)" means a licensed medical marijuana dispensary, medical marijuana processor, medical marijuana commercial grower, medical marijuana laboratory, medical marijuana business operator, or a medical marijuana transporter;
- 32. "Medical marijuana concentrate" or "concentrate" means a specific subset of medical marijuana that was produced by extracting

- cannabinoids from medical marijuana. Categories of medical
  marijuana concentrate include water-based medical marijuana

  concentrate, food-based medical marijuana concentrate, solvent-based
  medical marijuana concentrate, and heat- or pressure-based medical
  marijuana concentrate;
  - 33. "Medical marijuana commercial grower" or "commercial grower" means an entity licensed to cultivate, prepare and package medical marijuana and transfer or contract for transfer medical marijuana to a medical marijuana dispensary, medical marijuana processor, any other medical marijuana commercial grower, medical marijuana research facility, medical marijuana education facility and pesticide manufacturers. A commercial grower may sell seeds, flower or clones to commercial growers pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act;
  - 34. "Medical marijuana education facility" or "education facility" means a person or entity approved pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act to operate a facility providing training and education to individuals involving the cultivation, growing, harvesting, curing, preparing, packaging or testing of medical marijuana, or the production, manufacture, extraction, processing, packaging or creation of medical-marijuana-infused products or medical marijuana products as described in this act the Oklahoma Medical Marijuana and Patient Protection Act;

- 35. "Medical-marijuana-infused product" means a product infused with medical marijuana including, but not limited to, edible products, ointments and tinctures;
- 36. "Medical marijuana product" or "product" means a product that contains cannabinoids that have been extracted from plant material or the resin therefrom by physical or chemical means and is intended for administration to a qualified patient including, but not limited to, oils, tinctures, edibles, pills, topical forms, gels, creams, vapors, patches, liquids, and forms administered by a nebulizer, excluding live plant forms which are considered medical marijuana;
- 37. "Medical marijuana processor" means a person or entity licensed pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act to operate a business including the production, manufacture, extraction, processing, packaging or creation of concentrate, medical-marijuana-infused products or medical marijuana products as described in this act the Oklahoma Medical Marijuana and Patient Protection Act;
- 19 38. "Medical marijuana research facility" or "research
  20 facility" means a person or entity approved pursuant to this act the
  21 Oklahoma Medical Marijuana and Patient Protection Act to conduct
  22 medical marijuana research. A medical marijuana research facility
  23 is not a medical marijuana business;

- 39. "Medical marijuana testing laboratory" or "laboratory" means a public or private laboratory licensed pursuant to this act, the Oklahoma Medical Marijuana and Patient Protection Act to conduct testing and research on medical marijuana and, medical marijuana products and medical marijuana waste;
- 40. "Medical marijuana transporter" or "transporter" means a person or entity that is licensed pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act. A medical marijuana transporter does not include a medical marijuana business that transports its own medical marijuana, medical marijuana concentrate or medical marijuana products to a property or facility adjacent to or connected to the licensed premises if the property is another licensed premises of the same medical marijuana business;
- 41. "Medical marijuana waste" or "waste" means unused, surplus, returned or out-of-date marijuana, plant debris of the plant of the genus Cannabis, including dead plants and all unused plant parts and roots, except the term shall not include roots, stems, stalks and fan leaves;
- 42. "Medical use" means the acquisition, possession, use, delivery, transfer or transportation of medical marijuana, medical marijuana products, medical marijuana devices or paraphernalia relating to the administration of medical marijuana to treat a licensed patient;

- 43. "Mother plant" means a marijuana plant that is grown or maintained for the purpose of generating clones, and that will not be used to produce plant material for sale to a medical marijuana processor or medical marijuana dispensary;
  - 44. "Oklahoma physician" or "physician" means a physician
    licensed by and in good standing with the State Board of Medical
    Licensure and Supervision, the State Board of Osteopathic Examiners
    or the Board of Podiatric Medical Examiners;
  - 45. "Oklahoma resident" means an individual who can provide proof of residency as required by this act the Oklahoma Medical Marijuana and Patient Protection Act;
- 46. "Owner" means, except where the context otherwise requires, a direct beneficial owner including, but not limited to, all persons or entities as follows:
  - a. all shareholders owning an interest of a corporate entity and all officers of a corporate entity,
  - b. all partners of a general partnership,
  - c. all general partners and all limited partners that own an interest in a limited partnership,
  - d. all members that own an interest in a limited liability company,
  - e. all beneficiaries that hold a beneficial interest in a trust and all trustees of a trust,

1 f. all persons or entities that own interest in a joint venture,

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- all persons or entities that own an interest in an q. association,
- h. the owners of any other type of legal entity, and
- any other person holding an interest or convertible i. note in any entity which owns, operates or manages a licensed facility;
- 47. "Package" or "packaging" means any container or wrapper that may be used by a medical marijuana business to enclose or contain medical marijuana;
- 48. "Person" means a natural person, partnership, association, business trust, company, corporation, estate, limited liability company, trust or any other legal entity or organization, or a manager, agent, owner, director, servant, officer or employee thereof, except that "person" does not include any governmental organization;
- 49. "Pesticide" means any substance or mixture of substances 18 intended for preventing, destroying, repelling or mitigating any 19 pest or any substance or mixture of substances intended for use as a 20 plant regulator, defoliant or desiccant, except that the term 21 "pesticide" shall not include any article that is a "new animal 22 drug" as designated by the United States Food and Drug 23 Administration; 24

50. "Production batch" means:

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- a. any amount of medical marijuana concentrate of the same category and produced using the same extraction methods, standard operating procedures and an identical group of harvest batch of medical marijuana, or
- b. any amount of medical marijuana product of the same exact type, produced using the same ingredients, standard operating procedures and the same production batch of medical marijuana concentrate;
- 51. "Public institution" means any entity established or controlled by the federal government, state government, or a local government or municipality including, but not limited to, institutions of higher education or related research institutions;
- 52. "Public money" means any funds or money obtained by the holder from any governmental entity including, but not limited to, research grants;
- 53. "Recommendation" means a document that is signed or electronically submitted by a physician on behalf of a patient for the use of medical marijuana pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act;
- 54. "Registered to conduct business" means a person that has provided proof that the business applicant is in good standing with the Oklahoma Secretary of State and Oklahoma Tax Commission;

| 55. "Remediation" means the process by which the medical            |
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| marijuana flower or trim, which has failed microbial testing, is    |
| processed into solvent-based medical marijuana concentrate and      |
| retested as required by this act the Oklahoma Medical Marijuana and |
| Patient Protection Act;   |

- 56. "Research project" means a discrete scientific endeavor to answer a research question or a set of research questions related to medical marijuana and is required for a medical marijuana research license. A research project shall include a description of a defined protocol, clearly articulated goals, defined methods and outputs, and a defined start and end date. The description shall demonstrate that the research project will comply with all requirements in this act the Oklahoma Medical Marijuana and Patient Protection Act and rules promulgated pursuant thereto. All research and development conducted by a medical marijuana research facility shall be conducted in furtherance of an approved research project;
- 57. "Revocation" means the final decision by the Department that any license issued pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act is rescinded because the individual or entity does not comply with the applicable requirements set forth in this act the Oklahoma Medical Marijuana and Patient Protection Act or rules promulgated pursuant thereto;
- 58. "School" means a public or private preschool or a public or private elementary or secondary school used for school classes and

- instruction. A homeschool, daycare or child-care facility shall not be considered a "school" as used in this act the Oklahoma Medical
- 3 | Marijuana and Patient Protection Act;
  - 59. "Shipping container" means a hard-sided container with a lid or other enclosure that can be secured in place. A shipping container is used solely for the transport of medical marijuana, medical marijuana concentrate, or medical marijuana products between medical marijuana businesses, a medical marijuana research facility, or a medical marijuana education facility;
  - 60. "Solvent-based medical marijuana concentrate" means a medical marijuana concentrate that was produced by extracting cannabinoids from medical marijuana through the use of a solvent approved by the Department;
  - 61. "State Question" means Oklahoma State Question No. 788, Initiative Petition No. 412, approved by a majority vote of the citizens of Oklahoma on June 26, 2018;
  - 62. "Strain" means the classification of marijuana or cannabis plants in either pure sativa, indica, afghanica, ruderalis or hybrid varieties;
- 20 63. "THC" means tetrahydrocannabinol, which is the primary
  21 psychotropic cannabinoid in marijuana formed by decarboxylation of
  22 naturally tetrahydrocannabinolic acid, which generally occurs by
  23 exposure to heat;

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- 64. "Test batch" means with regard to usable marijuana, a homogenous, identified quantity of usable marijuana by strain, no greater than ten (10) pounds, that is harvested during a seven-day period from a specified cultivation area, and with regard to oils, vapors and waxes derived from usable marijuana, means an identified quantity that is uniform, that is intended to meet specifications for identity, strength and composition, and that is manufactured, packaged and labeled during a specified time period according to a single manufacturing, packaging and labeling protocol;
- 65. "Transporter agent" means a person who transports medical marijuana or medical marijuana products for a licensed transporter and holds a transporter agent license pursuant to this act the Oklahoma Medical Marijuana and Patient Protection Act;
- 66. "Universal symbol" means the image established by the State Department of Health or Oklahoma Medical Marijuana Authority and made available to licensees through its website indicating that the medical marijuana or the medical marijuana product contains THC;
- 67. "Usable marijuana" means the dried leaves, flowers, oils, vapors, waxes and other portions of the marijuana plant and any mixture or preparation thereof, excluding seed, roots, stems, stalks and fan leaves; and
- 68. "Water-based medical marijuana concentrate" means a concentrate that was produced by extracting cannabinoids from medical marijuana through the use of only water, ice, or dry ice.

- 1 SECTION 2. AMENDATORY Section 17, Chapter 11, O.S.L.
- 2 | 2019, as amended by Section 4, Chapter 312, O.S.L. 2019 (63 O.S.
- 3 | Supp. 2020, Section 427.17), is amended to read as follows:
- 4 Section 427.17. A. There is hereby created a medical marijuana
- 5 | testing laboratory license as a category of the medical marijuana
- 6 business license. The Authority is hereby enabled to monitor,
- 7 | inspect and audit a licensed testing laboratory under this act
- 8 | Section 427.1 et seq. of this title.
- 9 B. The Authority is hereby authorized to contract with a
- 10 private laboratory for the purpose of conducting compliance testing
- 11 of medical marijuana testing laboratories licensed in this state.
- 12 | Any such laboratory under contract for compliance testing shall be
- 13 | prohibited from conducting any other commercial medical marijuana
- 14 testing in this state.
- 15 C. The Authority shall have the authority to develop acceptable
- 16 testing and research practices, including but not limited to
- 17 testing, standards, quality control analysis, equipment
- 18 | certification and calibration, and chemical identification and
- 19 substances used in bona fide research methods so long as it complies
- 20 | with this act the Oklahoma Medical Marijuana and Patient Protection
- 21 <u>Act</u>.
- 22 D. A person who is a direct beneficial owner or an indirect
- 23 beneficial owner of a medical marijuana dispensary, medical

1 marijuana commercial grower, or medical marijuana processor shall 2 not be an owner of a laboratory.

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- 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 and medical marijuana waste for medical marijuana businesses, medical marijuana research facilities, medical marijuana education facilities, and testing and research on medical 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.
  - 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 23 medical marijuana, medical marijuana concentrate or, medical

- marijuana product <u>or medical marijuana waste</u> 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 <del>or</del>,
  medical marijuana product <u>or medical marijuana waste</u> 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 the Oklahoma Medical Marijuana and Patient Protection 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 requires 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 and medical marijuana waste for testing, in accordance with this act the Oklahoma Medical Marijuana and Patient Protection 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
  2 the location of the laboratory in a secure location, and inspection,
  3 cleaning and maintenance of any equipment or utensils used for the
  4 analysis of test samples;
  - 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 or medical marijuana waste 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

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- 1 disposal. The inventory tracking system reporting shall include the
- 2 results of any tests that are conducted on medical marijuana,
- 3 | medical marijuana concentrate or, medical marijuana product or
- 4 | medical marijuana waste;

- 9. Standards of performance;
  - 10. The employment of laboratory personnel;
- 7 11. A written standard operating procedure manual to be
- 8 maintained and updated by the laboratory;
- 9 12. The successful participation in a Department-approved
- 10 proficiency testing program for each testing category listed in this
- 11 | section, in order to obtain and maintain certification;
- 12 13. The establishment of and adherence to a quality assurance
- 13 | and quality control program to ensure sufficient monitoring of
- 14 | laboratory processes and quality of results reported;
- 15 14. The establishment by the laboratory of a system to document
- 16 | the complete chain of custody for samples from receipt through
- 17 | disposal;
- 18 15. The establishment by the laboratory of a system to retain
- 19 and maintain all required records, including business records, and
- 20 processes to ensure results are reported in a timely and accurate
- 21 | manner; and
- 22 16. Any other aspect of laboratory testing of medical marijuana
- 23 | or, medical marijuana product or medical marijuana waste deemed
- 24 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 <a href="medical">medical</a> marijuana <a href="medical marijuana">or,</a>
  <a href="medical marijuana">medical marijuana</a> waste 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, production batch, as appropriate, or waste batch of medical marijuana, medical marijuana concentrate and, medical marijuana product or medical marijuana waste for each of the following categories of testing, consistent with standards developed by the Commissioner:
  - 1. Microbials;
  - 2. Mycotoxins;
- Residual solvents;
- 24 4. Pesticides;

- 1 | 5. Tetrahydrocannabinol (THC) and other cannabinoid potency;
  - 6. Terpenoid potency; and
  - 7. Heavy metals.

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- R. A test batch shall not exceed ten (10) pounds of usable medical marijuana er, medical marijuana product, as appropriate or medical marijuana waste. A grower shall separate each harvest lot of usable marijuana into harvest batches containing no more than ten (10) pounds. A processor shall separate each medical marijuana production lot into production batches containing no more than ten (10) pounds. A grower or processor shall separate each medical marijuana waste lot into waste batches containing no more than ten (10) pounds.
  - 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.
- V. 1. A commercial grower shall not transfer or sell medical marijuana and a processor shall not transfer, sell or process into a concentrate or product any medical marijuana, medical marijuana concentrate or medical marijuana product unless samples from each harvest batch or production batch from which that medical marijuana, medical marijuana concentrate or medical marijuana product was derived has have been tested by a medical marijuana testing facility for contaminants and passed all contaminant tests required by this act the Oklahoma Medical Marijuana and Patient Protection Act.
- 2. A processor shall not transfer, sell or process into a concentrate or product any medical marijuana, medical marijuana concentrate or medical marijuana product unless samples from each production batch from which that medical marijuana, medical marijuana concentrate or medical marijuana product was derived have been tested by a medical marijuana testing facility for contaminants and passed all contaminant tests required by this act.
- 3. A commercial grower or processor shall not transfer medical marijuana waste to a medical marijuana waste disposal facility unless samples from each waste batch from which that medical marijuana waste was derived have been tested by a medical marijuana testing facility for contaminants.

| 1   | SECTION 3. This act shall become effective November 1, 2021.                            |  |
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| 2   | COMMITTEE REPORT BY: COMMITTEE ON HEALTH AND HUMAN SERVICES February 22, 2021 - DO PASS |  |
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