

1 **HOUSE OF REPRESENTATIVES - FLOOR VERSION**

2 STATE OF OKLAHOMA

3 2nd Session of the 59th Legislature (2024)

4 COMMITTEE SUBSTITUTE
5 FOR
6 HOUSE BILL NO. 3361

By: Marti

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8 COMMITTEE SUBSTITUTE

9 An Act relating to medical marijuana; amending 63
10 O.S. 2021, Section 427.18, as amended by Section 18,
11 Chapter 251, O.S.L. 2022 (63 O.S. Supp. 2023, Section
12 427.18), which relates to the Oklahoma Medical
13 Marijuana and Patient Protection Act; modifying
14 certain packaging requirements; requiring business
15 name logos to be designed in a certain manner;
16 providing administrative fines for violations;
17 directing the deposit of administrative fines in
18 specific revolving funds; directing licensed medical
19 marijuana processors and licensed medical marijuana
20 commercial growers to sell certain medical marijuana
21 products in pre-packaged form; providing requirements
22 for packaging; allowing for the display and smelling
23 of marijuana; directing the Oklahoma Medical
24 Marijuana Authority to promulgate certain rules;
 providing for codification; and providing an
 effective date.

BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

SECTION 1. AMENDATORY 63 O.S. 2021, Section 427.18, as
amended by Section 18, Chapter 251, O.S.L. 2022 (63 O.S. Supp. 2023,
Section 427.18), is amended to read as follows:

1 Section 427.18 A. A medical marijuana business shall not sell,
2 transfer or otherwise distribute medical marijuana or medical
3 marijuana product that has not been packaged and labeled in
4 accordance with this section and rules promulgated by the Executive
5 Director of the Oklahoma Medical Marijuana Authority.

6 B. A medical marijuana dispensary shall return medical
7 marijuana and medical marijuana product that does not meet packaging
8 or labeling requirements in this section or rules promulgated
9 pursuant thereto to the entity who transferred it to the dispensary.
10 The medical marijuana dispensary shall document to whom the item was
11 returned, what was returned and the date of the return or dispose of
12 any usable marijuana that does not meet these requirements in
13 accordance with the Oklahoma Medical Marijuana and Patient
14 Protection Act.

15 C. 1. Medical marijuana packaging shall be packaged to
16 minimize its appeal to children and shall not depict images other
17 than the business name logo of the medical marijuana producer ~~and~~
18 ~~image of the product.~~ The business name logo of the medical
19 marijuana producer shall also be designed in a manner that is not
20 appealing to children.

21 2. A medical marijuana business shall not place any content on
22 a container in a manner that reasonably appears to target
23 individuals under the age of twenty-one (21) including, but not
24 limited to, cartoon characters or similar images.

1 3. Labels on a container shall not include any false or
2 misleading statements.

3 4. No container shall be intentionally or knowingly labeled so
4 as to cause a reasonable patient confusion as to whether the medical
5 marijuana, medical marijuana concentrate or medical marijuana
6 product is a trademarked product or labeled in a manner that
7 violates any federal trademark law or regulation.

8 5. The label on the container shall not make any claims
9 regarding health or physical benefits to the patient.

10 6. All medical marijuana, medical marijuana concentrate and
11 medical marijuana products shall be in a child-resistant container
12 at the point of transfer to the patient or caregiver.

13 D. The Executive Director shall develop minimum standards for
14 packaging and labeling of medical marijuana and medical marijuana
15 products. Such standards shall include, but not be limited to, the
16 required contents of labels to be affixed to all medical marijuana
17 and medical marijuana products prior to transfer to a licensed
18 patient or caregiver, which shall include, at a minimum:

19 1. THC and other cannabinoid potency, and terpenoid potency;

20 2. A statement indicating that the product has been tested for
21 contaminants;

22 3. One or more product warnings to be determined by the
23 Executive Director; and
24

1 4. Any other information the Executive Director deems
2 necessary.

3 E. Any licensed medical marijuana dispensary that violates the
4 provisions of subsection B of this section shall be subject to an
5 administrative fine of Five Hundred Dollars (\$500.00) for each
6 separate violation. Administrative fines collected pursuant to the
7 provisions of this subsection shall be collected and deposited to
8 the revolving fund of the law enforcement agency responsible for the
9 investigation, enforcement, and prosecution of medical marijuana
10 dispensary licensees who violate the provisions of subsection B of
11 this section.

12 SECTION 2. NEW LAW A new section of law to be codified
13 in the Oklahoma Statutes as Section 431.1 of Title 63, unless there
14 is created a duplication in numbering, reads as follows:

15 A. Upon the effective date of this act, all medical marijuana
16 flower, trim, shake, kief, medical marijuana product, or other
17 flower-based product not defined as a concentrate, shall be sold by
18 licensed medical marijuana processors and licensed medical marijuana
19 commercial growers to licensed medical marijuana dispensaries only
20 in pre-packaged form in package sizes weighing not less than one-
21 half (1/2) of one (1) gram to not more than three (3) ounces.

22 B. Nonopaque materials may be used when packaging medical
23 marijuana flower provided all other packaging and labeling
24 requirements for medical marijuana products sold in this state are

1 met and it is placed in an opaque container before leaving a
2 licensed medical marijuana dispensary.

3 C. The display and smelling of medical marijuana shall be
4 allowed pursuant to Section 421 of Title 63 of the Oklahoma
5 Statutes.

6 D. The Oklahoma Medical Marijuana Authority shall promulgate
7 rules necessary to allow for pre-packaged products to be returned to
8 the licensed medical marijuana dispensary when found defective or
9 hazardous to the health of the patient. The Authority shall further
10 promulgate rules necessary to allow for the return of medical
11 marijuana products from a licensed medical marijuana dispensary to a
12 licensed medical marijuana processor or licensed medical marijuana
13 commercial grower, from a licensed medical marijuana processor to a
14 licensed medical marijuana commercial grower, or from any other
15 licensed entity that transferred medical marijuana products to
16 another licensed entity.

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18 COMMITTEE REPORT BY: COMMITTEE ON ALCOHOL, TOBACCO AND CONTROLLED
19 SUBSTANCES, dated 02/29/2024 - DO PASS, As Amended.
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