

1 ENGROSSED HOUSE AMENDMENT
TO
2 ENGROSSED SENATE BILL NO. 518 By: Alvord of the Senate
3 and
4 West (Kevin) of the House
5
6
7 An Act relating to medical marijuana packaging;
8 amending 63 O.S. 2021, Section 427.18, as last
9 amended by Section 144, Chapter 452, O.S.L. 2024 (63
10 O.S. Supp. 2024, Section 427.18), which relates to
11 packaging and labeling requirements; requiring
12 certain labeling; and providing an effective date.
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14 AMENDMENT NO. 1. Strike the title, enacting clause, and entire bill
15 and insert:
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17 "An Act relating to medical marijuana packaging;
18 amending 63 O.S. 2021, Section 427.18, as last
19 amended by Section 144, Chapter 452, O.S.L. 2024 (63
20 O.S. Supp. 2024, Section 427.18), which relates to
21 packaging and labeling requirements; requiring
22 certain warnings on labels; and providing an
23 effective date.
24
25 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

1 SECTION 1. AMENDATORY 63 O.S. 2021, Section 427.18, as
2 last amended by Section 144, Chapter 452, O.S.L. 2024 (63 O.S. Supp.
3 2024, Section 427.18), is amended to read as follows:

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

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

18 C. 1. Medical marijuana packaging shall be packaged to
19 minimize its appeal to children and shall not depict images other
20 than the business name logo of the medical marijuana producer and
21 image of the product.

22 2. A medical marijuana business shall not place any content on
23 a container in a manner that reasonably appears to target
24

1 individuals under the age of twenty-one (21) including, but not
2 limited to, cartoon characters or similar images.

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

5 4. No container shall be intentionally or knowingly labeled so
6 as to cause a reasonable patient confusion as to whether the medical
7 marijuana, medical marijuana concentrate or medical marijuana
8 product is a trademarked product or labeled in a manner that
9 violates any federal trademark law or regulation. The label on the
10 container shall include a warning that states the following:

11 a. "For use by licensed medical marijuana patients only",

12 ~~and~~

13 b. "Keep out of reach of children",

14 c. "It is illegal to drive a motor vehicle while under
15 the influence of marijuana or marijuana products",

16 d. "Women should not use marijuana or marijuana products
17 during pregnancy because of the risk of birth
18 defects", and

19 e. "This product has been tested for contaminants".

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

22 6. The container itself may be clear in order to allow licensed
23 medical marijuana patients and licensed medical marijuana caregivers
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1 the ability to view the product inside the container but shall be
2 child-resistant, as defined in Section 427.2 of this title.

3 7. At the point of sale and transfer of any medical marijuana,
4 medical marijuana concentrate, or medical marijuana products to a
5 licensed medical marijuana patient or licensed medical marijuana
6 caregiver, the dispensary shall place the medical marijuana, medical
7 marijuana concentrate, or medical marijuana products in an exit
8 package, as such term is defined in Section 427.2 of this title.

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

- 16 1. THC and other cannabinoid potency, and terpenoid potency;
- 17 2. A statement indicating that the product has been tested for
18 contaminants;
- 19 3. One or more product warnings to be determined by the
20 Executive Director; and
- 21 4. Any other information the Executive Director deems
22 necessary.

23 SECTION 2. This act shall become effective November 1, 2025."
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1 Passed the House of Representatives the 29th day of April, 2025.

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4 Presiding Officer of the House of
Representatives
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6 Passed the Senate the ____ day of _____, 2025.

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9 Presiding Officer of the Senate
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2 BILL NO. 518

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4 West (Kevin) of the House

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7 amending 63 O.S. 2021, Section 427.18, as last
8 amended by Section 144, Chapter 452, O.S.L. 2024 (63
9 O.S. Supp. 2024, Section 427.18), which relates to
10 packaging and labeling requirements; requiring
11 certain labeling; and providing an effective date.

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

13 SECTION 3. AMENDATORY 63 O.S. 2021, Section 427.18, as
14 last amended by Section 144, Chapter 452, O.S.L. 2024 (63 O.S. Supp.
15 2024, Section 427.18), is amended to read as follows:

16 Section 427.18. A. A medical marijuana business shall not
17 sell, transfer or otherwise distribute medical marijuana or medical
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20 Director of the Oklahoma Medical Marijuana Authority.

21 B. A medical marijuana dispensary shall return medical
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23 or labeling requirements in this section or rules promulgated
24 pursuant thereto to the entity who transferred it to the dispensary.
The medical marijuana dispensary shall document to whom the item was

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3 accordance with the Oklahoma Medical Marijuana and Patient
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10 a container in a manner that reasonably appears to target
11 individuals under the age of twenty-one (21) including, but not
12 limited to, cartoon characters or similar images.

13 3. Labels on a container shall not include any false or
14 misleading statements.

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16 as to cause a reasonable patient confusion as to whether the medical
17 marijuana, medical marijuana concentrate or medical marijuana
18 product is a trademarked product or labeled in a manner that
19 violates any federal trademark law or regulation. The label on the
20 container shall include a warning that states the following:

- 21 a. "For use by licensed medical marijuana patients only",
22 ~~and~~
23 b. "Keep out of reach of children" ~~and~~ and

1 c. “Marijuana and marijuana products can impair
2 concentration, coordination, and judgment: a person
3 should not operate a motor vehicle while under the
4 influence of marijuana or marijuana products. The
5 ingestion of any amount of marijuana or marijuana
6 products before driving may result in criminal
7 prosecution for driving under the influence.”

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

10 6. The container itself may be clear in order to allow licensed
11 medical marijuana patients and licensed medical marijuana caregivers
12 the ability to view the product inside the container but shall be
13 child-resistant, as defined in Section 427.2 of this title.

14 7. At the point of sale and transfer of any medical marijuana,
15 medical marijuana concentrate, or medical marijuana products to a
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24 be affixed to all medical marijuana, medical marijuana concentrate,

1 and medical marijuana products prior to transfer to a licensed
2 patient or caregiver, which shall include, at a minimum:

- 3 1. THC and other cannabinoid potency, and terpenoid potency;
- 4 2. A statement indicating that the product has been tested for
5 contaminants;
- 6 3. One or more product warnings to be determined by the
7 Executive Director; and
- 8 4. Any other information the Executive Director deems
9 necessary.

10 SECTION 4. This act shall become effective November 1, 2025.

11 Passed the Senate the 27th day of March, 2025.

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13 _____
14 Presiding Officer of the Senate

15 Passed the House of Representatives the ____ day of _____,
16 2025.

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18 _____
19 Presiding Officer of the House
20 of Representatives
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