

1 STATE OF OKLAHOMA

2 2nd Session of the 57th Legislature (2020)

3 HOUSE BILL 4120

By: Roberts (Sean)

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6 AS INTRODUCED

7 An Act relating to medical marijuana; amending
8 Section 18, Chapter 11, O.S.L. 2019 (63 O.S. Supp.
9 2019, Section 427.18), which relates to the Oklahoma
10 Medical Marijuana and Patient Protection Act;
11 requiring inclusion of certain warnings on labels of
12 medical marijuana products; and providing an
13 effective date.

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

15 SECTION 1. AMENDATORY Section 18, Chapter 11, O.S.L.
16 2019 (63 O.S. Supp. 2019, Section 427.18), is amended to read as
17 follows:

18 Section 427.18 A. An Oklahoma medical marijuana business shall
19 not sell, transfer or otherwise distribute medical marijuana or
20 medical marijuana product that has not been packaged and labeled in
21 accordance with this section and rules promulgated by the State
22 Commissioner of Health.

23 B. A medical marijuana dispensary shall return medical
24 marijuana and medical marijuana product that does not meet packaging
or labeling requirements in this section or rules promulgated

1 pursuant thereto to the entity who transferred it to the dispensary.
2 The medical marijuana dispensary shall document to whom the item was
3 returned, what was returned and the date of the return or dispose of
4 any usable marijuana that does not meet these requirements in
5 accordance with this act.

6 C. 1. Medical marijuana packaging shall be packaged to
7 minimize its appeal to children and shall not depict images other
8 than the business name logo of the medical marijuana producer and
9 image of the product.

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

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

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

21 5. The label on the container shall not make any claims
22 regarding health or physical benefits to the patient.
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1 6. All medical marijuana, medical marijuana concentrate and
2 medical marijuana products shall be in a child-resistant container
3 at the point of transfer to the patient or caregiver.

4 D. The State Department of Health shall develop minimum
5 standards for packaging and labeling of medical marijuana and
6 medical marijuana products. Such standards shall include, but not
7 be limited to, the required contents of labels to be affixed to all
8 medical marijuana and medical marijuana products prior to transfer
9 to a licensed patient or caregiver, which shall include, at a
10 minimum:

11 1. A universal symbol indicating that the product contains
12 tetrahydrocannabinol (THC);

13 2. THC and other cannabinoid potency, and terpenoid potency;

14 3. A statement indicating that the product has been tested for
15 contaminants;

16 4. One or more product warnings to be determined by the
17 Department; ~~and~~

18 5. A warning that states "Use of any product containing
19 tetrahydrocannabinol (THC) may result in adverse drug interactions,
20 side effects or other complications that could significantly
21 jeopardize the health or safety of the patient";

22 6. A warning that states "Do not use while driving or operating
23 heavy machinery"; and

24 7. Any other information the Department deems necessary.

1 SECTION 2. This act shall become effective November 1, 2020.

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