

Louisiana Administrative Code

Title 46 – Professional and Occupational Standards

Part LIII: Pharmacists

Chapter 24. Limited Service Providers

Subchapter E. Marijuana Pharmacy

§2443. Marijuana Products

A. – D.1.e.v. ...

2. Labeling.

- a. Each product shall be labeled by the producer prior to its sale to the marijuana pharmacy.
- b. Each label shall be securely affixed to the package and shall ~~include, at a minimum:~~ comply with labeling standards for marijuana products promulgated by LDAF.
 - i. ~~The batch or lot number assigned by the producer to the marijuana plant(s) from which the marijuana used in the product was harvested;~~
 - ii. ~~A complete list of solvents, chemicals, and pesticides used in the creation of any marijuana concentrate;~~
 - iii. ~~A complete list of all ingredients used to manufacture the product, which may include a list of any potential allergens contained within, or used in the manufacture of, a product;~~
 - iv. ~~The potency of the THC and CBD in the product, expressed in milligrams for each cannabinoid;~~
 - v. ~~The net weight, using a standard of measure compatible with the LMMTS, of the product prior to its placement in the shipping container;~~
 - vi. ~~A product expiration date, upon which the product will no longer be fit for use. Once a label with an expiration date has been affixed to a product, the producer shall not alter that date or affix a new label with a later date; and~~
 - vii. ~~A statement the product has been tested for contaminants, that there were no adverse findings, and the date of such testing.~~
 - viii. ~~A product identification code registered with the board.~~
- b. The labeling text on any marijuana product shall not make any false or misleading statements regarding health or physical benefits to the consumer. Further, each label shall include all of the following statements:
 - i. ~~“Contains Marijuana. For Medical Use Only. KEEP OUT OF THE REACH OF CHILDREN.”~~
 - ii. ~~“Marijuana can impair concentration, coordination, and judgment. Do not operate a vehicle or machinery under the influence of this drug.”~~

- 40 iii. ~~“There may be additional health risks associated with the consumption of this product~~
41 ~~— for women who are pregnant, breastfeeding, or planning to become pregnant.”~~
42 iv. ~~A statement that it is illegal for any person to possess or consume the contents of the~~
43 ~~— package other than the patient for whom it was recommended.~~
44 e. ~~The labeling text required by this Section shall be no smaller than 1/16 of an inch, shall be~~
45 ~~printed in English, and must be unobstructed and conspicuous.~~
46 c. The label for each product shall bear a product identification code registered with the board.
47 d. The producer may utilize a two-dimensional quick response (QR) code or a package insert
48 which is enclosed or attached to the product container to provide the information required in
49 this Section. If the producer elects to use such supplementary labeling, the label affixed to the
50 outer surface of the product container shall contain the following information, at a minimum
51 i. the batch or lot number ~~referenced at Clause D.2.a.i of this Section;~~
52 ii. the potency of ~~the any~~ THC and ~~or~~ CBD ~~referenced at Clause D.2.a.iv of this Section~~
53 contained therein;
54 iii. the net weight ~~referenced at Clause D.2.a.v of this Section;~~
55 iv. the expiration date ~~referenced at Clause D.2.a.vi of this Section;~~ and
56 v. ~~the any~~ caution statements ~~referenced at Clause D.2.b.i of this Section.~~

57 E. – E.4.f. ...

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59 AUTHORITY NOTE: Promulgated in accordance with R.S. 40:1046.

60 HISTORICAL NOTE: Promulgated by the Department of Health, Board of Pharmacy, LR 43:1540 (August 2017),
61 amended LR 45:1473 (October 2019), amended LR 46:568 (April 2020), amended LR
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