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State of Minnesota

HOUSE OF REPRESENTATIVES

NINETY-FIRST SESSION

H. F. No. 4179

03/05/2020 Authored by Vang and Koznick
The bill was read for the first time and referred to the Committee on Health and Human Services Policy

1.1 A bill for an act
1.2 relating to health; requiring cannabinoid product labels to contain a bar code or
1.3 QR code; amending Minnesota Statutes 2019 Supplement, section 151.72,
1.4 subdivision 5.

1.5 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:

1.6 Section 1. Minnesota Statutes 2019 Supplement, section 151.72, subdivision 5, is amended
1.7 to read:

1.8 Subd. 5. Labeling requirements. (a) A product regulated under this section must bear
1.9 a label that contains, at a minimum:

1.10 (1) the name, location, contact phone number, and website of the manufacturer of the
1.11 product;

1.12 (2) the name and address of the independent, accredited laboratory used by the
1.13 manufacturer to test the product;

1.14 (3) an accurate statement of the amount or percentage of cannabinoids found in each
1.15 unit of the product meant to be consumed; and

1.16 (4) a scannable bar code or QR code that links to the following information on the
1.17 product:

1.18 (i) batch identification number;

1.19 (ii) product name;

1.20 (iii) batch date;

1.21 (iv) expiration date;

- 2.1 (v) batch size;
- 2.2 (vi) total quantity produced; and
- 2.3 (vii) ingredient used, including the:
- 2.4 (A) ingredient name;
- 2.5 (B) name of company that manufactured the ingredient;
- 2.6 (C) company or product identification number, if applicable; and
- 2.7 (D) ingredient lot number; and
- 2.8 ~~(4)~~ (5) a statement stating that this product does not claim to diagnose, treat, cure, or
- 2.9 prevent any disease and has not been evaluated or approved by the United States Food and
- 2.10 Drug Administration (FDA) unless the product has been so approved.
- 2.11 (b) The information required to be on the label must be prominently and conspicuously
- 2.12 placed and in terms that can be easily read and understood by the consumer.
- 2.13 (c) The label must not contain any claim that the product may be used or is effective for
- 2.14 the prevention, treatment, or cure of a disease or that it may be used to alter the structure
- 2.15 or function of human or animal bodies, unless the claim has been approved by the FDA.