

Minn.Rule 4770.0850. Packaging and Labeling

Subpart 3. Labeling.

A. A medical cannabis manufacturer must ensure that all medical cannabis that is distributed is labeled with the following information:

- (1) the patient's registry identification number, name, and date of birth;
- (2) the name and date of birth of the designated registered caregiver, if applicable;
- (3) the name of the patient's parent or legal guardian, if listed on the registry verification, if applicable;
- (4) the patient's address;
- (5) the name ~~and address~~ of the medical cannabis manufacturer where the medical cannabis was manufactured;
- (6) the medical cannabis's chemical composition;
- (7) ~~the recommended dosage~~dosing instructions recommended for the patient;
- (8) directions for use of the product;
- (9) all ingredients of the product shown with common or usual names, including any colors, artificial flavors, and preservatives, listed in descending order by predominance of weight;
- (10) the date of manufacture and batch number;
- (11) a notice with the statement, including capitalization: "This product has not been analyzed or approved by the United States Food and Drug Administration. There is limited information on the side effects of using this product, and there may be associated health risks. Do not drive or operate heavy machinery when under the influence of this product. KEEP THIS PRODUCT OUT OF REACH OF CHILDREN."; and
- (12) a notice with the statement: "This medical cannabis is for therapeutic use only. Diversion of this product is unlawful and may result in the revocation of the patient's registration."

B. Labeling text must not include any false or misleading statements regarding health or physical benefits to the patient.

C. A package may contain multiple labels if the information required by this part is not obstructed.