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Subd. 5. Labeling requirements. (a) A product regulated under this section m a label that contains, at a minimum: (1) the name, location, contact phone number, and website of the manufacturer product; (2) the name and address of the independent, accredited laboratory used by the manufacturer to test the product; (3) an accurate statement of the amount or percentage of cannabinoids found in unit of the product meant to be consumed; and or (4) a scannable bar code or QR code that links to the manufacturer's website; a (4) (5) a statement stating that this product does not claim to diagnose, treat, or prevent any disease and has not been evaluated or approved by the United States F. Drug Administration (FDA) unless the product has been so approved. (b) The information required to be on the label must be prominently and conspit placed and in terms that can be easily read and understood by the consumer. (c) The label must not contain any claim that the product may be used or is effect the prevention, treatment, or cure of a disease or that it may be used to alter the str or function of human or animal bodies, unless the claim has been approved by the On R106, House language, (H2128-4), Article 3 Page 254, after line 22, insert: "EFFECTIVE DATE. This section is effective the day following final enactm On R109, House language, (H2128-4), Article 3 Page 257, after line 24, insert:	1.1	Senator moves to amend H.F. No. 2128, in conference committee, as
Page 232, after line 9, insert: "Sec. 67. Minnesota Statutes 2020, section 151.72, subdivision 5, is amended to Subd. 5. Labeling requirements. (a) A product regulated under this section m a label that contains, at a minimum: (1) the name, location, contact phone number, and website of the manufacturer product; (2) the name and address of the independent, accredited laboratory used by the manufacturer to test the product; (3) an accurate statement of the amount or percentage of cannabinoids found in unit of the product meant to be consumed; and or (4) a scannable bar code or QR code that links to the manufacturer's website; at the prevent any disease and has not been evaluated or approved by the United States F. Drug Administration (FDA) unless the product has been so approved. (b) The information required to be on the label must be prominently and conspipalized and in terms that can be easily read and understood by the consumer. (c) The label must not contain any claim that the product may be used or is effect the prevention, treatment, or cure of a disease or that it may be used to alter the stroof function of human or animal bodies, unless the claim has been approved by the On R106, House language, (H2128-4), Article 3 Page 254, after line 22, insert: "EFFECTIVE DATE, This section is effective the day following final enacting the product of the product	1.2	follows:
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On R109, House language, (H2128-4), Article 3 Page 257, after line 24, insert: "EFFECTIVE DATE. This section is effective the day following final enactors."	1.24	Page 254, after line 22, insert:
Page 257, after line 24, insert: "EFFECTIVE DATE. This section is effective the day following final enactors."	1.25	"EFFECTIVE DATE. This section is effective the day following final enactment."
1.28 "EFFECTIVE DATE. This section is effective the day following final enactors	1.26	On R109, House language, (H2128-4), Article 3
	1.27	Page 257, after line 24, insert:
On R110, House language, (H2128-4), Article 3	1.28	"EFFECTIVE DATE. This section is effective the day following final enactment."
	1.29	On R110, House language, (H2128-4), Article 3

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2.1	Page 258, after line 13, insert:
2.2	"EFFECTIVE DATE. This section is effective the day following final enactment."
2.3	On R112, House language, (H2128-4), Article 3
2.4	Page 260, after line 25, insert:
2.5	"EFFECTIVE DATE. This section is effective the day following final enactment."
2.6	On R115, House language, (H2128-4), Article 3
2.7	Page 264, line 23, before "Paragraph" insert "Paragraph (c) is effective the day following
2.8	final enactment. "
2.9	Page 264, line 27, delete "this section" and insert "paragraph (e)"
2.10	On R116, House language, (H2128-4), Article 3
2.11	Page 265, after line 22, insert:
2.12	"EFFECTIVE DATE. This section is effective the day following final enactment."
2.13	Page 265, before line 23, insert:
2.14	"Sec. 89. Minnesota Statutes 2020, section 152.27, subdivision 2, is amended to read:
2.15	Subd. 2. Commissioner duties. (a) The commissioner shall:
2.16	(1) give notice of the program to health care practitioners in the state who are eligible
2.17	to serve as health care practitioners and explain the purposes and requirements of the
2.18	program;
2.19	(2) allow each health care practitioner who meets or agrees to meet the program's
2.20	requirements and who requests to participate, to be included in the registry program to
2.21	collect data for the patient registry;
2.22	(3) provide explanatory information and assistance to each health care practitioner in
2.23	understanding the nature of therapeutic use of medical cannabis within program requirements;
2.24	(4) create and provide a certification to be used by a health care practitioner for the
2.25	practitioner to certify whether a patient has been diagnosed with a qualifying medical
2.26	condition and include in the certification an option for the practitioner to certify whether
2.27	the patient, in the health care practitioner's medical opinion, is developmentally or physically
2.28	disabled and, as a result of that disability, the patient requires assistance in administering
2.29	medical cannabis or obtaining medical cannabis from a distribution facility;

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(5) supervise the participation of the health care practitioner in conducting patient treatment and health records reporting in a manner that ensures stringent security and record-keeping requirements and that prevents the unauthorized release of private data on individuals as defined by section 13.02;

- (6) develop safety criteria for patients with a qualifying medical condition as a requirement of the patient's participation in the program, to prevent the patient from undertaking any task under the influence of medical cannabis that would constitute negligence or professional malpractice on the part of the patient; and
- (7) conduct research and studies based on data from health records submitted to the registry program and submit reports on intermediate or final research results to the legislature and major scientific journals. The commissioner may contract with a third party to complete the requirements of this clause. Any reports submitted must comply with section 152.28, subdivision 2.
- (b) The commissioner may add a delivery method under section 152.22, subdivision 6, or add, remove, or modify a qualifying medical condition under section 152.22, subdivision 14, upon a petition from a member of the public or the task force on medical cannabis therapeutic research or as directed by law. The commissioner shall evaluate all petitions to add a qualifying medical condition or to remove or modify an existing qualifying medical condition submitted by the task force on medical cannabis therapeutic research or as directed by law and shall may make the addition, removal, or modification if the commissioner determines the addition, removal, or modification is warranted based on the best available evidence and research. If the commissioner wishes to add a delivery method under section 152.22, subdivision 6, or add or remove a qualifying medical condition under section 152.22, subdivision 14, the commissioner must notify the chairs and ranking minority members of the legislative policy committees having jurisdiction over health and public safety of the addition or removal and the reasons for its addition or removal, including any written comments received by the commissioner from the public and any guidance received from the task force on medical cannabis research, by January 15 of the year in which the commissioner wishes to make the change. The change shall be effective on August 1 of that year, unless the legislature by law provides otherwise."
 - Page 265, after line 30, insert:
- 3.32 **"EFFECTIVE DATE.** This section is effective the day following final enactment."
- Renumber the sections in sequence and correct the internal references
- 3.34 Amend the title accordingly

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Sec. 89.