	MEDICAL CANNABIS REVISIONS
	2021 GENERAL SESSION
	STATE OF UTAH
	Chief Sponsor: Jennifer Dailey-Provost
	Senate Sponsor:
LONG T	ITLE
General	Description:
Tł	nis bill amends provisions related to the processing and dispensing of medical
cannabis.	
Highligh	ted Provisions:
Tł	nis bill:
•	prohibits the inclusion of a certain oil in medical cannabis products;
►	requires the inclusion of synthetic additives on the label of a medical cannabis
product;	
•	addresses allowed delivery of medical cannabis shipments by a medical cannabis
pharmacy	; and
►	makes technical and conforming changes.
Money A	ppropriated in this Bill:
N	one
Other Sp	ecial Clauses:
N	one
Utah Coo	le Sections Affected:
AMEND	5:
4-	41a-602, as last amended by Laws of Utah 2020, Chapter 12
4-	41a-603, as last amended by Laws of Utah 2020, Chapter 12
26	-61a-305 , as last amended by Laws of Utah 2020, Chapter 12

H.B. 435

Be it enacted by the Legislature of the state of Utah:
Section 1. Section 4-41a-602 is amended to read:
4-41a-602. Cannabis product Labeling and child-resistant packaging.
(1) For any cannabis product that a cannabis processing facility processes or produces
and for any raw cannabis that the facility packages, the facility shall:
(a) label the cannabis or cannabis product with a label that:
(i) clearly and unambiguously states that the cannabis product or package contains
cannabis;
(ii) clearly displays the amount of total composite tetrahydrocannabinol and
cannabidiol in the labeled container;
(iii) has a unique identification number that:
(A) is connected to the inventory control system; and
(B) identifies the unique cannabis product manufacturing process the cannabis
processing facility used to manufacture the cannabis product;
(iv) identifies the cannabinoid extraction process that the cannabis processing facility
used to create the cannabis product;
(v) does not display an image, word, or phrase that the facility knows or should know
appeals to children; and
(vi) discloses each:
(A) active or potentially active ingredient, in order of prominence[;];
(B) synthetic additive; and
(C) possible allergen; and
(b) package the raw cannabis or cannabis product in a medicinal dosage form in a
container that:
(i) is tamper evident and tamper resistant;
(ii) does not appeal to children;
(iii) does not mimic a candy container;
(iv) is opaque;
(v) complies with child-resistant effectiveness standards that the United States
Consumer Product Safety Commission establishes; and

59	(vi) includes a warning label that states: "WARNING: Cannabis has intoxicating
60	effects and may be addictive. Do not operate a vehicle or machinery under its influence. KEEP
61	OUT OF REACH OF CHILDREN. This product is for medical use only. Use only as directed
62	by a qualified medical provider."
63	(2) For any cannabis or cannabis product that the cannabis processing facility processes
64	into a gelatinous cube, gelatinous rectangular cuboid, or lozenge in a cube or rectangular
65	cuboid shape, the facility shall:
66	(a) ensure that the label described in Subsection (1)(a) does not contain a photograph or
67	other image of the content of the container; and
68	(b) include on the label described in Subsection (1)(a) a warning about the risks of
69	over-consumption.
70	(3) The department shall make rules in accordance with Title 63G, Chapter 3, Utah
71	Administrative Rulemaking Act to establish:
72	(a) a standard labeling format that:
73	(i) complies with the requirements of this section; and
74	(ii) ensures inclusion of a pharmacy label; and
75	(b) additional requirements on packaging for cannabis and cannabis products to ensure
76	safety and product quality.
77	Section 2. Section 4-41a-603 is amended to read:
78	4-41a-603. Cannabis product Product quality.
79	(1) As used in this section, "MCT oil" means a supplement made from medium-chain
80	triglycerides.
81	[(1)] (2) A cannabis processing facility:
82	(a) may not produce a cannabis product in a physical form that:
83	(i) the facility knows or should know appeals to children;
84	(ii) is designed to mimic or could be mistaken for a candy product; or
85	(iii) for a cannabis product used in vaporization, includes a candy-like flavor or another
86	flavor that the facility knows or should know appeals to children; [and]
87	(b) may not include MCT oil in a medical cannabis product; and
88	[(b)] (c) notwithstanding Subsection $[(1)]$ (2)(a)(iii), may produce a concentrated oil
89	with a flavor that the department approves to facilitate minimizing the taste or odor of

H.B. 435

90	cannabis.
91	[(2)] (3) A cannabis product may vary in the cannabis product's labeled cannabinoid
92	profile by up to 10% of the indicated amount of a given cannabinoid, by weight.
93	[(3)] (4) The department shall adopt by rule, in accordance with Title 63G, Chapter 3,
94	Utah Administrative Rulemaking Act, human safety standards for the manufacturing of
95	cannabis products that are consistent with best practices for the use of cannabis.
96	Section 3. Section 26-61a-305 is amended to read:
97	26-61a-305. Maximum number of licenses Home delivery medical cannabis
98	pharmacies.
99	(1) (a) Except as provided in Subsections (1)(b) or (d), if a sufficient number of
100	applicants apply, the department shall issue 14 medical cannabis pharmacy licenses in
101	accordance with this section.
102	(b) If fewer than 14 qualified applicants apply for a medical cannabis pharmacy
103	license, the department shall issue a medical cannabis pharmacy license to each qualified
104	applicant.
105	(c) The department may issue the licenses described in Subsection (1)(a) in two phases
106	in accordance with this Subsection (1)(c).
107	(i) Using one procurement process, the department may issue eight licenses to an initial
108	group of medical cannabis pharmacies and six licenses to a second group of medical cannabis
109	pharmacies.
110	(ii) If the department issues licenses in two phases in accordance with this Subsection
111	(1)(c), the department shall:
112	(A) divide the state into no less than four geographic regions;
113	(B) issue at least one license in each geographic region during each phase of issuing
114	licenses; and
115	(C) complete the process of issuing medical cannabis pharmacy licenses no later than
116	July 1, 2020.
117	(d) (i) The department may issue licenses to operate a medical cannabis pharmacy in
118	addition to the licenses described in Subsection (1)(a) if the department determines, in
119	consultation with the Department of Agriculture and Food and after an annual or more frequent
120	analysis of the current and anticipated market for medical cannabis, that each additional license

121 is necessary to provide an adequate supply, quality, or variety of medical cannabis to medical 122 cannabis cardholders. 123 (ii) The department shall: 124 (A) in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, 125 make rules to establish criteria and processes for the consultation, analysis, and application for 126 a license described in Subsection (1)(d)(i); 127 (B) before November 30, 2020, report on the rules described in Subsection 128 (1)(d)(ii)(A) to the Executive Appropriations Committee of the Legislature; and 129 (C) report to the Executive Appropriations Committee of the Legislature before each 130 time the department issues an additional license under Subsection (1)(d)(i) regarding the results 131 of the consultation and analysis described in Subsection (1)(d)(i) and the application of the 132 criteria described in Subsection (1)(d)(ii)(A) to the intended licensee. 133 (2) (a) If there are more qualified applicants than there are available licenses for 134 medical cannabis pharmacies, the department shall: 135 (i) evaluate each applicant and award the license to the applicant that best 136 demonstrates: 137 (A) experience with establishing and successfully operating a business that involves 138 complying with a regulatory environment, tracking inventory, and training, evaluating, and 139 monitoring employees; 140 (B) an operating plan that will best ensure the safety and security of patrons and the 141 community; 142 (C) positive connections to the local community; 143 (D) the suitability of the proposed location and the location's accessibility for 144 qualifying patients; (E) the extent to which the applicant can increase efficiency and reduce the cost of 145 146 medical cannabis for patients; and 147 (F) a strategic plan described in Subsection 26-61a-304(7) that has a comparatively 148 high likelihood of success; and 149 (ii) ensure a geographic dispersal among licensees that is sufficient to reasonably 150 maximize access to the largest number of medical cannabis cardholders. 151 (b) In making the evaluation described in Subsection (2)(a), the department may give

H.B. 435

02-25-21 1:26 PM

152	increased consideration to applicants who indicate a willingness to:
153	(i) operate as a home delivery medical cannabis pharmacy that accepts electronic
154	medical cannabis orders that the state central patient portal facilitates; and
155	(ii) accept payments through:
156	(A) a payment provider that the Division of Finance approves, in consultation with the
157	state treasurer, in accordance with Section 26-61a-603; or
158	(B) a financial institution in accordance with Subsection 26-61a-603(4).
159	(3) The department may conduct a face-to-face interview with an applicant for a
160	license that the department evaluates under Subsection (2).
161	(4) (a) The department may designate a medical cannabis pharmacy as a home delivery
162	medical cannabis pharmacy if the department determines that the medical cannabis pharmacy's
163	operating plan demonstrates the functional and technical ability to:
164	(i) safely conduct transactions for medical cannabis shipments;
165	(ii) accept electronic medical cannabis orders that the state central patient portal
166	facilitates; and
167	(iii) accept payments through:
168	(A) a payment provider that the Division of Finance approves, in consultation with the
169	state treasurer, in accordance with Section 26-61a-603; or
170	(B) a financial institution in accordance with Subsection 26-61a-603(4).
171	(b) An applicant seeking a designation as a home delivery medical cannabis pharmacy
172	shall identify in the applicant's operating plan any information relevant to the department's
173	evaluation described in Subsection (4)(a), including:
174	(i) the name and contact information of the payment provider;
175	(ii) the nature of the relationship between the prospective licensee and the payment
176	provider;
177	(iii) the processes of the following to safely and reliably conduct transactions for
178	medical cannabis shipments:
179	(A) the prospective licensee; and
180	(B) the electronic payment provider or the financial institution described in Subsection
181	(4)(a)(iii); and
182	(iv) the ability of the licensee to comply with the department's rules regarding the

- 183 secure transportation and delivery of medical cannabis or medical cannabis product to a
- 184 medical cannabis cardholder.
- 185 (c) Notwithstanding any county or municipal ordinance, a medical cannabis pharmacy
- 186 that the department designates as a home delivery medical cannabis pharmacy may deliver
- 187 medical cannabis shipments in accordance with this chapter.
- 188 (d) A home delivery medical cannabis pharmacy may not deliver a medical cannabis
- 189 shipment within a 35-mile radius of a home delivery medical cannabis pharmacy that is located
- 190 within a county of the third through sixth class.