# Jennifer Dailey-Provost proposes the following substitute bill:

1

## **Cannabis Amendments**

# 2025 GENERAL SESSION STATE OF UTAH

**Chief Sponsor: Jennifer Dailey-Provost** 

Senate Sponsor: Evan J. Vickers

2

5

#### LONG TITLE

## **4** General Description:

This bill amends provisions related to medical cannabis.

# 6 **Highlighted Provisions:**

- 7 This bill:
- 8 defines terms;
- 9 allows for additional medical cannabis pharmacies;
- creates a new medical cannabis pharmacy license for independent medical cannabis
- 11 pharmacies;
- 12 creates ownership restrictions for independent medical cannabis pharmacies;
- 13 adjusts fees for certain medical cannabis pharmacy licenses;
- 14 amends provisions regarding cannabis production and sanitation;
- 15 modifies provisions related to enforcement and appeals;
- → amends provisions related to closed-door medical cannabis pharmacies;
- 17 allows a cannabis processing facility to have a website that includes product information;
- limits the number of licenses that the Department of Agriculture and Food (department)
- may issue for cannabis processing facilities;
- 20 amends provisions regarding when the department may seize products and test products;
- 21 amends provisions related to information a medical cannabis pharmacy must have
- 22 available to a patient purchasing medical cannabis;
- creates a reporting requirement for the department;
- repeals sections related to the state central patient portal;
- creates a medical cannabis ombudsman and duties for the ombudsman;
- 26 authorizes the creation of patient product information inserts;
- ▶ moves the repeal of the Cannabis Research Review Board earlier one year;
- 28 extends the repeal date for the Medical Cannabis Governance Structure Working Group:

29 and 30 makes technical and conforming changes. 31 **Money Appropriated in this Bill:** 32 None 33 **Other Special Clauses:** 34 None **Utah Code Sections Affected:** 35 36 AMENDS: 37 **4-41a-102**, as last amended by Laws of Utah 2024, Chapters 217, 238 and 240 38 **4-41a-110**, as enacted by Laws of Utah 2023, Chapter 273 39 **4-41a-205**, as last amended by Laws of Utah 2020, Chapter 12 40 4-41a-401, as last amended by Laws of Utah 2024, Chapter 217 41 4-41a-403, as last amended by Laws of Utah 2023, Chapter 327 42 **4-41a-501**, as last amended by Laws of Utah 2023, Chapter 313 43 **4-41a-701**, as last amended by Laws of Utah 2023, Chapters 313, 317 44 4-41a-801, as renumbered and amended by Laws of Utah 2018, Third Special Session, 45 Chapter 1 46 **4-41a-802**, as last amended by Laws of Utah 2024, Chapter 217 47 **4-41a-1001**, as last amended by Laws of Utah 2024, Chapters 217, 238 and 240 48 4-41a-1003, as last amended by Laws of Utah 2023, Chapter 435 and renumbered and 49 amended by Laws of Utah 2023, Chapters 273, 307 and last amended by Coordination Clause, 50 Laws of Utah 2023, Chapter 307 51 **4-41a-1005**, as last amended by Laws of Utah 2024, Chapter 217 52 **4-41a-1101**, as last amended by Laws of Utah 2024, Chapter 217 53 **4-41a-1201**, as enacted by Laws of Utah 2023, Chapter 273 54 **4-41a-1202**, as last amended by Laws of Utah 2024, Chapters 217, 240 55 4-41a-1203, as renumbered and amended by Laws of Utah 2023, Chapters 273, 307 and 56 last amended by Coordination Clause, Laws of Utah 2023, Chapter 307 57 **4-41a-1206**, as enacted by Laws of Utah 2024, Chapter 238 58 26B-1-310, as last amended by Laws of Utah 2023, Chapters 273, 281 and renumbered 59 and amended by Laws of Utah 2023, Chapter 305 and last amended by Coordination Clause, 60 Laws of Utah 2023, Chapter 305 61 **26B-1-435**, as last amended by Laws of Utah 2024, Chapters 238, 240 62 **26B-4-201**, as last amended by Laws of Utah 2024, Chapters 217, 240

- 63 **26B-4-202**, as last amended by Laws of Utah 2024, Chapters 217, 240 64 **26B-4-214**, as last amended by Laws of Utah 2024, Chapter 240 65 **26B-4-222**, as last amended by Laws of Utah 2024, Chapter 240 66 **26B-4-243**, as enacted by Laws of Utah 2023, Chapter 281 67 **26B-4-247**, as enacted by Laws of Utah 2023, Chapter 273 68 63I-2-204, as last amended by Laws of Utah 2024, Third Special Session, Chapter 5 69 63I-2-226, as last amended by Laws of Utah 2024, Third Special Session, Chapter 5 70 63I-2-236, as last amended by Laws of Utah 2024, Third Special Session, Chapter 5 71 **ENACTS**: 72 **4-41a-1006**, Utah Code Annotated 1953 73 **26B-4-248**, Utah Code Annotated 1953 74 **REPEALS:** 75 4-41a-801.1, as renumbered and amended by Laws of Utah 2023, Chapters 273, 307 and 76 last amended by Coordination Clause, Laws of Utah 2023, Chapter 307 77 26B-4-236, as last amended by Laws of Utah 2023, Chapters 273, 317 and renumbered 78 and amended by Laws of Utah 2023, Chapter 307 and last amended by Coordination Clause, 79 Laws of Utah 2023, Chapter 307 80 81 *Be it enacted by the Legislature of the state of Utah:* 82 Section 1. Section **4-41a-102** is amended to read: 83 4-41a-102 . Definitions. 84 As used in this chapter: 85 (1) "Adulterant" means any poisonous or deleterious substance in a quantity that may be 86 injurious to health, including: 87 (a) pesticides;
- 88 (b) heavy metals;
- 89 (c) solvents;
- 90 (d) microbial life;
- 91 (e) artificially derived cannabinoid;
- 92 (f) toxins; or
- 93 (g) foreign matter.
- 94 (2) "Advertise" or "advertising" means information provided by a person in any medium:
- 95 (a) to the public; and
- 96 (b) that is not age restricted to an individual who is at least 21 years old.

97	(3) "Advisory board" means the Medical Cannabis Policy Advisory Board created in
98	Section 26B-1-435.
99	(4)(a) "Anticompetitive business practice" means any practice that reduces the amount
100	of competition in the medical cannabis market that would be considered an attempt to
101	monopolize, as defined in Section 76-10-3103.
102	(b) "Anticompetitive business practice" may include:
103	(i) agreements that may be considered unreasonable when competitors interact to the
104	extent that they are:
105	(A) no longer acting independently; or
106	(B) when collaborating are able to wield market power together;
107	(ii) monopolizing or attempting to monopolize trade by:
108	(A) acting to maintain or acquire a dominant position in the market; or
109	(B) preventing new entry into the market; or
110	(iii) other conduct outlined in rule.
111	(5)(a) "Artificially derived cannabinoid" means a chemical substance that is created by a
112	chemical reaction that changes the molecular structure of any chemical substance
113	derived from the cannabis plant.
114	(b) "Artificially derived cannabinoid" does not include:
115	(i) a naturally occurring chemical substance that is separated from the cannabis plant
116	by a chemical or mechanical extraction process; or
117	(ii) a cannabinoid that is produced by decarboxylation from a naturally occurring
118	cannabinoid acid without the use of a chemical catalyst.
119	(6) "Batch" means a quantity of:
120	(a) cannabis extract produced on a particular date and time and produced between
121	completion of equipment and facility sanitation protocols until the next required
122	sanitation cycle during which lots of cannabis are used;
123	(b) cannabis product produced on a particular date and time and produced between
124	completion of equipment and facility sanitation protocols until the next required
125	sanitation cycle during which cannabis extract is used; or
126	(c) cannabis flower packaged on a particular date and time and produced between
127	completion of equipment and facility sanitation protocols until the next required
128	sanitation cycle during which lots of cannabis are being used.
129	[(6)] (7) "Cannabis Research Review Board" means the Cannabis Research Review Board
130	created in Section 26B-1-420.

131	[(7)] (8) "Cannabis" means the same as that term is defined in Section 26B-4-201.
132	[(8)] (9) "Cannabis concentrate" means:
133	(a) the product of any chemical or physical process applied to naturally occurring
134	biomass that concentrates or isolates the cannabinoids contained in the biomass; and
135	(b) any amount of a natural cannabinoid or artificially derived cannabinoid in an
136	artificially derived cannabinoid's purified state.
137	[(9)] (10) "Cannabis cultivation byproduct" means any portion of a cannabis plant that is not
138	intended to be sold as a cannabis plant product.
139	[(10)] (11) "Cannabis cultivation facility" means a person that:
140	(a) possesses cannabis;
141	(b) grows or intends to grow cannabis; and
142	(c) sells or intends to sell cannabis to a cannabis cultivation facility, a cannabis
143	processing facility, or a medical cannabis research licensee.
144	[(11)] (12) "Cannabis cultivation facility agent" means an individual who
145	holds a valid cannabis production establishment agent registration card with a cannabis
146	cultivation facility designation.
147	[(12)] (13) "Cannabis derivative product" means a product made using cannabis concentrate.
148	[(13)] (14) "Cannabis plant product" means any portion of a cannabis plant intended to be
149	sold in a form that is recognizable as a portion of a cannabis plant.
150	[(14)] (15) "Cannabis processing facility" means a person that:
151	(a) acquires or intends to acquire cannabis from a cannabis production establishment;
152	(b) possesses cannabis with the intent to manufacture a cannabis product;
153	(c) manufactures or intends to manufacture a cannabis product from unprocessed
154	cannabis or a cannabis extract; and
155	(d) sells or intends to sell a cannabis product to a medical cannabis pharmacy or a
156	medical cannabis research licensee.
157	[(15)] (16) "Cannabis processing facility agent" means an individual who
158	holds a valid cannabis production establishment agent registration card with a cannabis
159	processing facility designation.
160	[(16)] (17) "Cannabis product" means the same as that term is defined in Section 26B-4-201.
161	[(17)] (18) "Cannabis production establishment" means a cannabis cultivation facility, a
162	cannabis processing facility, or an independent cannabis testing laboratory.
163	[(18)] (19) "Cannabis production establishment agent" means a cannabis cultivation facility
164	agent, a cannabis processing facility agent, or an independent cannabis testing laboratory

165	agent.
166	[(19)] (20) "Cannabis production establishment agent registration card" means a registration
167	card that the department issues that:
168	(a) authorizes an individual to act as a cannabis production establishment agent; and
169	(b) designates the type of cannabis production establishment for which an individual is
170	authorized to act as an agent.
171	[(20)] (21) "Closed-door medical cannabis pharmacy" means a facility operated by a home
172	delivery medical cannabis pharmacy for delivering [eannabis or a medical cannabis
173	product] medical cannabis.
174	[(21)] (22) "Community location" means a public or private elementary or secondary school
175	a church, a public library, a public playground, or a public park.
176	[(22)] (23) "Cultivation space" means, quantified in square feet, the horizontal area in which
177	a cannabis cultivation facility cultivates cannabis, including each level of horizontal area
178	if the cannabis cultivation facility hangs, suspends, stacks, or otherwise positions plants
179	above other plants in multiple levels.
180	[ <del>(23)</del> ] <u>(24)</u> "Delivery address" means:
181	(a) for a medical cannabis cardholder who is not a facility:
182	(i) the medical cannabis cardholder's home address; or
183	(ii) an address designated by the medical cannabis cardholder that:
184	(A) is the medical cannabis cardholder's workplace; and
185	(B) is not a community location; or
186	(b) for a medical cannabis cardholder that is a facility, the facility's address.
187	[(24)] (25) "Department" means the Department of Agriculture and Food.
188	[(25)] (26) "Family member" means a parent, step-parent, spouse, child, sibling,
189	step-sibling, uncle, aunt, nephew, niece, first cousin, mother-in-law, father-in-law,
190	brother-in-law, sister-in-law, son-in-law, daughter-in-law, grandparent, or grandchild.
191	[(26)] (27) "Government issued photo identification" means the same as that term is defined
192	in Section 26B-4-201, including expired identification in accordance with Section
193	26B-4-244.
194	[(27)] (28) "Home delivery medical cannabis pharmacy" means a medical cannabis
195	pharmacy that the department authorizes, as part of the pharmacy's license, to deliver
196	medical cannabis shipments to a delivery address to fulfill electronic orders[-that the
197	state central patient portal facilitates].
198	[(28)] (29)(a) "Independent cannabis testing laboratory" means a person that:

199	(i) conducts a chemical or other analysis of cannabis or a cannabis product; or
200	(ii) acquires, possesses, and transports cannabis or a cannabis product with the intent
201	to conduct a chemical or other analysis of the cannabis or cannabis product.
202	(b) "Independent cannabis testing laboratory" includes a laboratory that the department
203	or a research university operates in accordance with Subsection 4-41a-201(14).
204	[(29)] (30) "Independent cannabis testing laboratory agent" means an individual who
205	holds a valid cannabis production establishment agent registration card with an
206	independent cannabis testing laboratory designation.
207	[(30)] (31) "Inventory control system" means a system described in Section 4-41a-103.
208	[(31)] (32) "Licensing board" or "board" means the Cannabis Production Establishment and
209	Pharmacy Licensing Advisory Board created in Section 4-41a-201.1.
210	[(32)] (33) "Medical cannabis" or "medical cannabis product" means the same as that term is
211	defined in Section 26B-4-201.
212	[(33)] (34) "Medical cannabis card" means the same as that term is defined in Section
213	26B-4-201.
214	[ <del>(34)</del> ] <u>(35)</u> "Medical cannabis courier" means a courier that:
215	(a) the department licenses in accordance with Section 4-41a-1201; and
216	(b) contracts with a home delivery medical cannabis pharmacy to deliver medical
217	cannabis shipments to fulfill electronic orders[-that the state central patient portal
218	facilitates].
219	[(35)] (36) "Medical cannabis courier agent" means an individual who:
220	(a) is an employee of a medical cannabis courier; and
221	(b) who holds a valid medical cannabis courier agent registration card.
222	(37) "Medical cannabis ombudsman" means the ombudsman created in Section 26B-4-248.
223	[(36)] (38) "Medical cannabis pharmacy" means the same as that term is defined in Section
224	26B-4-201.
225	[(37)] (39) "Medical cannabis pharmacy agent" means the same as that term is defined in
226	Section 26B-4-201.
227	[(38)] (40) "Medical cannabis research license" means a license that the department issues to
228	a research university for the purpose of obtaining and possessing medical cannabis for
229	academic research.
230	[(39)] (41) "Medical cannabis research licensee" means a research university that the
231	department licenses to obtain and possess medical cannabis for academic research, in
232	accordance with Section 4-41a-901.

- 233 [(40)] (42) "Medical cannabis shipment" means a shipment of medical cannabis that a home
- 234 delivery medical cannabis pharmacy or a medical cannabis courier delivers to a delivery
- address to fulfill an electronic medical cannabis order[that the state central patient portal
- 236 <u>facilitates</u>].
- 237 [(41)] (43) "Medical cannabis treatment" means the same as that term is defined in Section
- 238 26B-4-201.
- 239 [(42)] (44) "Medicinal dosage form" means the same as that term is defined in Section
- 240 26B-4-201.
- 241 (45) "Patient product information insert" means the same as that term is defined in Section
- 242 <u>26B-4-201.</u>
- [(43)] (46) "Pharmacy ownership limit" means an amount equal to 30% of the total number
- of medical cannabis pharmacy licenses issued by the department rounded down to the
- 245 nearest whole number.
- 246 [(44)] (47) "Pharmacy medical provider" means the same as that term is defined in Section
- 247 26B-4-201.
- 248 [(45)] (48) "Qualified medical provider" means the same as that term is defined in Section
- 249 26B-4-201.
- 250 [(46)] (49) "Qualified Production Enterprise Fund" means the fund created in Section
- 251 4-41a-104.
- 252 [(47)] (50) "Recommending medical provider" means the same as that term is defined in
- 253 Section 26B-4-201.
- 254 [(48)] (51) "Research university" means the same as that term is defined in Section
- 255 53B-7-702 and a private, nonprofit college or university in the state that:
- 256 (a) is accredited by the Northwest Commission on Colleges and Universities;
- (b) grants doctoral degrees; and
- (c) has a laboratory containing or a program researching a schedule I controlled
- substance described in Section 58-37-4.
- 260 [(49)] (52) "State electronic verification system" means the system described in Section
- 261 26B-4-202.
- 262 [(50)] (53) "Targeted marketing" means the promotion of [a cannabis product,] medical
- 263 <u>cannabis</u>, a medical cannabis brand, or a medical cannabis device using any of the
- 264 following methods:
- 265 (a) electronic communication to an individual who is at least 21 years old and has
- requested to receive promotional information;

300

267	(b) an in-person marketing event that is:
268	(i) held inside a medical cannabis pharmacy; and
269	(ii) in an area where only a medical cannabis cardholder may access the event;
270	(c) other marketing material that is physically available or digitally displayed in a
271	medical cannabis pharmacy; or
272	(d) a leaflet a medical cannabis pharmacy places in the opaque package or box that is
273	provided to an individual when obtaining medical cannabis:
274	(i) in the medical cannabis pharmacy;
275	(ii) at the medical cannabis pharmacy's drive-through pick up window; or
276	(iii) in a medical cannabis shipment.
277	[(51)] (54) "Tetrahydrocannabinol" or "THC" means the same as that term is defined in
278	Section 4-41-102.
279	[(52)] (55) "THC analog" means the same as that term is defined in Section 4-41-102.
280	[(53)] (56) "Total composite tetrahydrocannabinol" means all detectable forms of
281	tetrahydrocannabinol.
282	[(54)] (57) "Total tetrahydrocannabinol" or "total THC" means the same as that term is
283	defined in Section 4-41-102.
284	Section 2. Section <b>4-41a-110</b> is amended to read:
285	4-41a-110 . Department coordination.
286	The department shall:
287	(1) provide draft rules made under this chapter to:
288	(a) the advisory board for the advisory board's review; and
289	(b) the medical cannabis ombudsman;
290	(2) consult with the advisory board before issuing an additional:
291	(a) cultivation facility license under Section 4-41a-205; or
292	(b) pharmacy license under Section 4-41a-1005;
293	(3) consult with the advisory board regarding fees set by the department that pertain to the
294	medical cannabis program; and
295	(4) when appropriate, consult with the advisory board regarding issues that arise in the
296	medical cannabis program.
297	Section 3. Section <b>4-41a-205</b> is amended to read:
298	4-41a-205 . Number of licenses Cannabis cultivation facilities Cannabis
299	processing facilities.

(1) Except as provided in Subsection (2)(a), the department shall issue at least five but not

334

301	more than eight licenses to operate a cannabis cultivation facility.
302	(2)(a) The department may issue a number of licenses to operate a cannabis cultivation
303	facility that, in addition to the licenses described in Subsection (1), does not cause the
304	total number of licenses to exceed 15 if the department determines, in consultation
305	with the Department of Health and Human Services and after an annual or more
306	frequent analysis of the current and anticipated market for medical cannabis, that
307	each additional license is necessary to provide an adequate supply, quality, or variety
308	of medical cannabis to medical cannabis cardholders.
309	(b) If the recipient of one of the initial licenses described in Subsection (1) ceases
310	operations for any reason or otherwise abandons the license, the department may but
311	is not required to grant the vacant license to another applicant based on an analysis as
312	described in Subsection (2)(a).
313	(3) If there are more qualified applicants than the number of available licenses for cannabis
314	cultivation facilities under Subsections (1) and (2), the department shall evaluate the
315	applicants and award the limited number of licenses described in Subsections (1) and (2)
316	to the applicants that best demonstrate:
317	(a) experience with establishing and successfully operating a business that involves:
318	(i) complying with a regulatory environment;
319	(ii) tracking inventory; and
320	(iii) training, evaluating, and monitoring employees;
321	(b) an operating plan that will best ensure the safety and security of patrons and the
322	community;
323	(c) positive connections to the local community; and
324	(d) the extent to which the applicant can increase efficiency and reduce the cost to
325	patients of medical cannabis.
326	(4) The department may conduct a face-to-face interview with an applicant for a license that
327	the department evaluates under Subsection (3).
328	(5) The licensing board may not issue more than 18 cannabis processing facility licenses.
329	Section 4. Section <b>4-41a-401</b> is amended to read:
330	4-41a-401 . Cannabis production establishment General operating
331	requirements.
332	(1)(a) A cannabis production establishment shall operate in accordance with the
333	operating plan described in Sections 4-41a-201 and 4-41a-204.

(b) A cannabis production establishment shall notify the department before a change in

335	the cannabis production establishment's operating plan.
336	(c)(i) If a cannabis production establishment changes the cannabis production
337	establishment's operating plan, the establishment shall ensure that the new
338	operating plan complies with this chapter.
339	(ii) The department shall establish by rule, in accordance with Title 63G, Chapter 3,
340	Utah Administrative Rulemaking Act, a process to:
341	(A) review a change notification described in Subsection (1)(b);
342	(B) identify for the cannabis production establishment each point of
343	noncompliance between the new operating plan and this chapter;
344	(C) provide an opportunity for the cannabis production establishment to address
345	each identified point of noncompliance; and
346	(D) suspend or revoke a license if the cannabis production establishment fails to
347	cure the noncompliance.
348	(2) A cannabis production establishment shall operate:
349	(a) except as provided in Subsection (5), in a facility that is accessible only by an
350	individual with a valid cannabis production establishment agent registration card
351	issued under Section 4-41a-301; and
352	(b) at the physical address provided to the department under Section 4-41a-201.
353	(3) A cannabis production establishment may not employ an individual who is younger than
354	21 years old.
355	(4) A cannabis production establishment may not employ an individual who has been
356	convicted, under state or federal law, of:
357	(a) a felony in the preceding 10 years; or
358	(b) after December 3, 2018, a misdemeanor for drug distribution.
359	(5) A cannabis production establishment may authorize an individual who is at least 18
360	years old and is not a cannabis production establishment agent to access the cannabis
361	production establishment if the cannabis production establishment:
362	(a) tracks and monitors the individual at all times while the individual is at the cannabis
363	production establishment; and
364	(b) maintains a record of the individual's access, including arrival and departure.
365	(6) A cannabis production establishment shall operate in a facility that has:
366	(a) a single, secure public entrance;
367	(b) a security system with a backup power source that:
368	(i) detects and records entry into the cannabis production establishment; and

369	(11) provides notice of an unauthorized entry to law enforcement when the cannabis
370	production establishment is closed; and
371	(c) a lock or equivalent restrictive security feature on any area where the cannabis
372	production establishment stores cannabis or a cannabis product.
373	(7) The department shall make rules establishing requirements for cannabis production
374	establishments regarding:
375	(a) master manufacturing plans;
376	(b) batch production records;
377	(c) sanitary operations;
378	(d) sanitary facilities and controls;
379	(e) equipment and utensils;
380	(f) production and process controls;
381	(g) warehousing and distribution; and
382	(h) employee personal hygiene.
383	Section 5. Section <b>4-41a-403</b> is amended to read:
384	4-41a-403 . Advertising.
385	(1) Except as provided in this section and Section 4-41a-604, a cannabis production
386	establishment may not advertise to the general public in any medium.
387	(2) A cannabis production establishment may advertise an employment opportunity at the
388	cannabis production establishment.
389	(3)(a) A cannabis production establishment may maintain a website that:
390	[(a)] (i) contains information about the establishment and employees; and
391	[(b)] (ii) except as provided in Subsection (3)(b), does not advertise any medical
392	cannabis, cannabis products, or medical cannabis devices.
393	(b) A cannabis processing facility may:
394	(i) if the website has age verification mechanisms that effectively prevent access by
395	individuals under 21 years of age, maintain a website that contains:
396	(A) educational information regarding medical cannabis produced by the cannabis
397	processing facility, including the certificate of analysis that is created by an
398	independent cannabis testing facility; and
399	(B) where medical cannabis produced by the cannabis processing facility may be
400	purchased in the state; and
401	(ii) engage in targeted marketing in accordance with Section 4-41a-604 for
402	advertising a particular medical cannabis product, medical cannabis device, or

403	medical cannabis brand.
404	(4)(a) Notwithstanding any municipal or county ordinance prohibiting signage, a
405	cannabis production establishment may use signage on the outside of the cannabis
406	production establishment that:
407	(i) includes only:
408	(A) in accordance with Subsection (4)(b), the cannabis production establishment's
409	name, logo, and hours of operation; and
410	(B) a green cross; and
411	(ii) complies with local ordinances regulating signage.
412	(b) The department shall define standards for a cannabis production establishment's
413	name and logo to ensure a medical rather than recreational disposition.
414	(5)(a) A cannabis production establishment may hold an educational event for the public
415	or medical providers in accordance with this Subsection (5) and the rules described in
416	Subsection (5)(c).
417	(b) A cannabis production establishment may not include in an educational event
418	described in Subsection (5)(a):
419	(i) any topic that conflicts with this chapter or Title 26B, Chapter 4, Part 2,
420	Cannabinoid Research and Medical Cannabis;
421	(ii) any gift items or merchandise other than educational materials, as those terms are
422	defined by the department;
423	(iii) any marketing for a specific product from the cannabis production establishment
424	or any other statement, claim, or information that would violate the federal Food,
425	Drug, and Cosmetic Act, 21 U.S.C. Sec. 301, et seq.; or
426	(iv) a presenter other than the following:
427	(A) a cannabis production establishment agent;
428	(B) a pharmacist licensed under Title 58, Chapter 17b, Pharmacy Practice Act;
429	(C) an advanced practice registered nurse licensed under Title 58, Chapter 31b,
430	Nurse Practice Act;
431	(D) a physician licensed under Title 58, Chapter 67, Utah Medical Practice Act, or
432	Title 58, Chapter 68, Utah Osteopathic Medical Practice Act;
433	(E) a physician assistant licensed under Title 58, Chapter 70a, Utah Physician
434	Assistant Act; or
435	(F) a state employee.
436	(c) The department shall make rules, in accordance with Title 63G, Chapter 3, Utah

437	Administrative Rulemaking Act, to define the elements of and restrictions on the
438	educational event described in Subsection (5)(a), including a minimum age of 21
439	years old for attendees.
440	Section 6. Section <b>4-41a-501</b> is amended to read:
441	4-41a-501. Cannabis cultivation facility Operating requirements.
442	(1) A cannabis cultivation facility shall ensure that any cannabis growing at the cannabis
443	cultivation facility is not visible from the ground level of the cannabis cultivation facility
444	perimeter.
445	(2) A cannabis cultivation facility shall use a unique identifier that is connected to the
446	facility's inventory control system to identify:
447	(a) beginning at the time a cannabis plant is eight inches tall and has a root ball, each
448	cannabis plant;
449	(b) each unique harvest of cannabis plants;
450	(c) each batch of cannabis the facility transfers to a medical cannabis pharmacy, a
451	cannabis processing facility, or an independent cannabis testing laboratory; and
452	(d) any excess, contaminated, or deteriorated cannabis of which the cannabis cultivation
453	facility disposes.
454	(3) A cannabis cultivation facility shall identify cannabis biomass as cannabis byproduct or
455	cannabis plant product before transferring the cannabis biomass from the facility.
456	(4) A cannabis cultivation facility shall either:
457	(a) ensure that a cannabis processing facility chemically or physically processes
458	cannabis cultivation byproduct to produce a cannabis concentrate for incorporation
459	into cannabis derivative products; or
460	(b) destroy cannabis cultivation byproduct in accordance with Section 4-41a-405.
461	(5) A cannabis cultivation facility may utilize radiation-based methods and equipment for
462	quality assurance or remediation purposes.
463	(6) The department shall make rules establishing:
464	(a) the records a cannabis cultivation facility must keep regarding each batch, amount of
465	product treated, and the methods used; and
466	(b) disclosure requirements to a cannabis processor receiving the material subject to the
467	radiation including the methods and equipment used.
468	Section 7. Section <b>4-41a-701</b> is amended to read:
469	4-41a-701. Cannabis and cannabis product testing.
470	(1) In accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, the

471	department may make rules to:
472	(a) determine required adulterant tests for a cannabis plant product, cannabis
473	concentrate, or cannabis product;
474	(b) determine the amount of any adulterant that is safe for human consumption;
475	(c) immediately ban or limit the presence of any ingredient in a medical cannabis
476	product after receiving a recommendation to do so from a public health authority
477	under Section 26B-1-102;
478	(d) establish protocols for a recall of [eannabis or a cannabis product] medical cannabis
479	by a cannabis production establishment; or
480	(e) allow the propagation of testing results forward to derived product if the processing
481	steps the cannabis production establishment uses to produce the product are unlikely
482	to change the results of the test.
483	(2)(a) The department may require testing for a toxin if:
484	[(a)] (i) the department receives information indicating the potential presence of a
485	toxin; or
486	[(b)] (ii) the department's inspector has reason to believe a toxin may be present based
487	on the inspection of a facility.
488	(b) The department may not require a cannabis processor to test a cannabis batch or a
489	cannabis product batch a third time if the cannabis batch or cannabis product has
490	previously met all testing requirements after being tested by:
491	(i) an independent cannabis testing laboratory that is not the department; and
492	(ii) the department.
493	(3)(a) A cannabis production establishment may not:
494	(i) incorporate cannabis concentrate into a cannabis derivative product until an
495	independent cannabis testing laboratory tests the cannabis concentrate in
496	accordance with department rule; or
497	(ii) transfer cannabis or a cannabis product to a medical cannabis pharmacy until an
498	independent cannabis testing laboratory tests a representative sample of the
499	cannabis or cannabis product in accordance with department rule.
500	(b) A medical cannabis pharmacy may not offer any cannabis or cannabis product for
501	sale unless an independent cannabis testing laboratory has tested a representative
502	sample of the cannabis or cannabis product in accordance with department rule.
503	(4) Before the sale of a <u>medical</u> cannabis product, an independent cannabis testing
504	laboratory shall:

505	(a) identify and quantify any cannabinoid known to be present in [a] the medical
506	cannabis product; and
507	(b) test terpene profiles for the following products:
508	(i) raw cannabis; or
509	(ii) a cannabis product:
510	(A) contained in a vaporizer cartridge; or
511	(B) in concentrate form; and
512	(c) record the five highest terpene profiles tested under Subsection (4)(b).
513	(5) The department shall establish by rule, in accordance with Title 63G, Chapter 3, Utah
514	Administrative Rulemaking Act, the standards, methods, practices, and procedures for
515	the testing of cannabis and cannabis products by independent cannabis testing
516	laboratories.
517	(6) The department may require an independent cannabis testing laboratory to participate in
518	a proficiency evaluation that the department conducts or that an organization that the
519	department approves conducts.
520	Section 8. Section <b>4-41a-801</b> is amended to read:
521	4-41a-801 . Enforcement Fine Citation.
522	(1)(a) If a person that is a cannabis production establishment[-or], a cannabis production
523	establishment agent, a medical cannabis pharmacy, a medical cannabis pharmacy
524	agent, or a medical cannabis courier violates this chapter, the department may:
525	[(a)] (i) revoke the person's license or [eannabis production establishment] agent
526	registration card;
527	[(b)] (ii) decline to renew the person's license [or cannabis production establishment]
528	agent registration card;
529	(iii) provide a letter of concern in accordance with Subsection (10); or
530	[(e)] (iv) assess the person an administrative penalty that the department establishes
531	by rule in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking
532	Act.
533	(b) Except for a violation that threatens public health or for the third violation of the
534	same rule or statute in a 24-month period, the department shall issue a letter of
535	concern before taking other administrative action under this section.
536	(2) The department shall deposit an administrative penalty imposed under this section into
537	the General Fund.
538	(3)(a) The department may take an action described in Subsection (3)(b) if the

539		department concludes, upon investigation, that[, for a person that is] a cannabis
540		production establishment[ $-or$ ], a cannabis production establishment agent[ $\div$ ], a
541		medical cannabis pharmacy, a medical cannabis pharmacy agent, or a medical
542		cannabis courier
543		[(i) the person] has violated the provisions of this chapter, a rule made under this
544		chapter, or an order issued under this chapter[; or] .
545		[(ii) the person produced cannabis or a cannabis product batch that contains a
546		substance, other than cannabis, that poses a significant threat to human health.]
547		(b) If the department makes the determination about a person described in Subsection
548		(3)(a), the department shall:
549		(i) issue the person a written administrative citation;
550		(ii) attempt to negotiate a stipulated settlement;
551		[(iii) seize, embargo, or destroy the cannabis or cannabis product batch;]
552		[(iv)] (iii) order the person to cease and desist from the action that creates a violation; [
553		and] <u>or</u>
554		[(v)] (iv) direct the person to appear before an adjudicative proceeding conducted
555		under Title 63G, Chapter 4, Administrative Procedures Act.
556		(c) If the department concludes, upon investigation, that a cannabis production
557		establishment or a cannabis production establishment agent has produced a cannabis
558		batch or a cannabis product batch that contains a substance that poses a significant
559		threat to human health, the department shall seize, embargo, or destroy the cannabis
560		batch or cannabis product batch.
561	(4)	The department may, for a person subject to an uncontested citation, a stipulated
562		settlement, or a finding of a violation in an adjudicative proceeding under this section,
563		for a fine amount not already specified in law, assess the person, who is not an
564		individual, a fine of up to \$5,000 per violation, in accordance with a fine schedule that
565		the department establishes by rule in accordance with Title 63G, Chapter 3, Utah
566		Administrative Rulemaking Act.
567	(5)	The department may not revoke a [eannabis production establishment's-]license without
568		first directing the [eannabis production establishment] <u>licensee</u> to appear before an
569		adjudicative proceeding conducted under Title 63G, Chapter 4, Administrative
570		Procedures Act.
571	(6)	If within $[20]$ 30 calendar days after the day on which a department serves a citation for
572		a violation of this chapter, the person that is the subject of the citation fails to request a

573	hearing to contest the citation, the citation becomes the department's final order.
574	(7) The department may, for a person who fails to comply with a citation under this section:
575	(a) refuse to issue or renew the person's license or cannabis production establishment
576	agent registration card; or
577	(b) suspend, revoke, or place on probation the person's license or cannabis production
578	establishment registration card.
579	(8)(a) Except where a criminal penalty is expressly provided for a specific violation of
580	this chapter, if an individual:
581	(i) violates a provision of this chapter, the individual is:
582	(A) guilty of an infraction; and
583	(B) subject to a \$100 fine; or
584	(ii) intentionally or knowingly violates a provision of this chapter or violates this
585	chapter three or more times, the individual is:
586	(A) guilty of a class B misdemeanor; and
587	(B) subject to a \$1,000 fine.
588	(b) An individual who is guilty of a violation described in Subsection (8)(a) is not guilty
589	of a violation of Title 58, Chapter 37, Utah Controlled Substances Act, for the
590	conduct underlying the violation described in Subsection (8)(a).
591	(9) Nothing in this section prohibits the department from referring potential criminal
592	activity to law enforcement.
593	(10)(a) A letter of concern shall describe:
594	(i) the violation including the statute or rule being violated;
595	(ii) possible options to remedy the issue; and
596	(iii) possible consequences for not remedying the violation.
597	(b) Under a letter of concern, the department shall provide the person at least 30 days to
598	remedy the violation.
599	(c) If the person fails to remedy the violation described in a letter of concern, the
600	department may take other enforcement action as described in this section.
601	(d) If a letter of concern is resolved without an enforcement action being taken under
602	Subsection (10)(c), the department may not report that a letter of concern was issued
603	to the licensing board.
604	(11)(a) An appeal of administrative action taken under this chapter shall be heard by the
605	medical cannabis ombudsman as an informal proceeding in accordance with Title
606	63G, Chapter 4, Administrative Procedures Act.

607	(b) Subsection (11)(a) is only effective when the position of medical cannabis
608	ombudsman is actively occupied by an employed individual.
609	Section 9. Section <b>4-41a-802</b> is amended to read:
610	4-41a-802 . Report.
611	(1) At or before the November interim meeting each year, the department shall report to the
612	Health and Human Services Interim Committee on:
613	(a) the number of applications and renewal applications that the department receives
614	under this chapter;
615	(b) the number of each type of cannabis production facility that the department licenses
616	in each county;
617	(c) the amount of cannabis that licensees grow;
618	(d) the amount of cannabis that licensees manufacture into cannabis products;
619	(e) the number of licenses the department revokes under this chapter;
620	(f) the department's operation of an independent cannabis testing laboratory under
621	Section 4-41a-201, including:
622	(i) the cannabis and cannabis products the department tested; and
623	(ii) the results of the tests the department performed;
624	(g) the expenses incurred and revenues generated under this chapter; and
625	(h) an analysis of product availability in medical cannabis pharmacies in consultation
626	with the Department of Health and Human Services.
627	(2) The department may not include personally identifying information in the report
628	described in this section.
629	(3) The department shall report to the working group described in Section 36-12-8.2 as
630	requested by the working group.
631	(4)(a) Before August 1, of each year, the department shall provide a report to the
632	working group described in Section 36-12-8.2 that provides the following for each
633	fine issued by the department under this chapter:
634	(i) the date of the fine;
635	(ii) the reference to the statute or rule that was violated for each fine issued; and
636	(iii) a short description explaining why the fine was issued.
637	(b) The report described in Subsection (4)(a) may not include identifying information of
638	the person that was subject to the fine.
639	Section 10. Section <b>4-41a-1001</b> is amended to read:
640	4-41a-1001 . Medical cannabis pharmacy License Eligibility.

541	(1) A person may not:
542	(a) operate as a medical cannabis pharmacy without a license that the department issues
543	under this part;
544	(b) obtain a medical cannabis pharmacy license if obtaining the license would cause the
545	person to exceed the pharmacy ownership limit;
546	(c) obtain a partial ownership share of a medical cannabis pharmacy if obtaining the
547	partial ownership share would cause the person to exceed the pharmacy ownership
548	limit; or
549	(d) enter into any contract or agreement that allows the person to directly or indirectly
550	control the operations of a medical cannabis pharmacy if the person's control of the
551	medical cannabis pharmacy would cause the person to effectively exceed the
552	pharmacy ownership limit.
553	(2)(a)(i) Subject to Subsections (4) and (5) and to Section 4-41a-1005, the department
554	shall issue a license to operate a medical cannabis pharmacy through the licensing
555	board created under Section 4-41a-201.1.
656	(ii) The department may not issue a license to operate a medical cannabis pharmacy
557	to an applicant who is not eligible for a license under this section.
558	(b) An applicant is eligible for a license under this section if the applicant submits to the
559	department:
560	(i) subject to Subsection (2)(c), a proposed name and address where the applicant will
561	operate the medical cannabis pharmacy;
562	(ii) the name and address of an individual who:
563	(A) for a publicly traded company, has a financial or voting interest of 10% or
564	greater in the proposed medical cannabis pharmacy;
565	(B) for a privately held company, a financial or voting interest in the proposed
566	medical cannabis pharmacy; or
567	(C) has the power to direct or cause the management or control of a proposed
568	medical cannabis pharmacy;
569	(iii) for each application that the applicant submits to the department, a statement
570	from the applicant that the applicant will obtain and maintain:
571	(A) a performance bond in the amount of \$100,000 issued by a surety authorized
572	to transact surety business in the state; or
573	(B) a liquid cash account in the amount of \$100,000 with a financial institution;
574	(iv) an operating plan that:

675	(A) complies with Section 4-41a-1004;
676	(B) includes operating procedures to comply with the operating requirements for a
677	medical cannabis pharmacy described in this part and with a relevant municipal
678	or county law that is consistent with Section 4-41a-1106; and
679	(C) the department approves;
680	(v) an application fee in an amount that, subject to Subsection 4-41a-104(5), the
681	department sets in accordance with Section 63J-1-504; and
682	(vi) a description of any investigation or adverse action taken by any licensing
683	jurisdiction, government agency, law enforcement agency, or court in any state for
684	any violation or detrimental conduct in relation to any of the applicant's
685	cannabis-related operations or businesses.
686	(c)(i) A person may not locate a medical cannabis pharmacy:
687	(A) within 200 feet of a community location; or
688	(B) in or within 600 feet of a district that the relevant municipality or county has
689	zoned as primarily residential.
690	(ii) The proximity requirements described in Subsection (2)(c)(i) shall be measured
691	from the nearest entrance to the medical cannabis pharmacy establishment by
692	following the shortest route of ordinary pedestrian travel to the property boundary
693	of the community location or residential area.
694	(iii) The department may grant a waiver to reduce the proximity requirements in
695	Subsection (2)(c)(i) by up to 20% if the department determines that it is not
696	reasonably feasible for the applicant to cite the proposed medical cannabis
697	pharmacy without the waiver.
698	(iv) An applicant for a license under this section shall provide evidence of
699	compliance with the proximity requirements described in Subsection (2)(c)(i).
700	(d) The department may not issue a license to an eligible applicant that the department
701	has selected to receive a license until the selected eligible applicant complies with the
702	bond or liquid cash requirement described in Subsection (2)(b)(iii).
703	(e) If the department receives more than one application for a medical cannabis
704	pharmacy within the same city or town, the department shall consult with the local
705	land use authority before approving any of the applications pertaining to that city or
706	town.
707	(f) In considering the issuance of a medical cannabis pharmacy license under this
708	section, the department may consider the extent to which the pharmacy can increase

709	efficiency and reduce cost to patients of medical cannabis.
710	[(3) If the department selects an applicant-]
711	(3)(a) After an entity has been selected for a medical cannabis pharmacy license under
712	this section, the department shall:
713	[(a)] (i) charge the applicant an initial license fee in an amount that, subject to
714	Subsection 4-41a-104(5), the department sets in accordance with Section
715	63J-1-504;
716	[(b)] (ii) notify the Department of Public Safety of the license approval and the names
717	of each individual described in Subsection (2)(b)(ii); and
718	[(e)] (iii) charge the licensee a fee in an amount that, subject to Subsection 4-41a-104
719	(5), the department sets in accordance with Section 63J-1-504, for any change in
720	location, ownership, or company structure.
721	(b) For a fee described in Subsection (3)(a)(i), a license fee for a medical cannabis
722	pharmacy located in a medically underserved area as determined by the federal
723	Health Resources and Services Administration shall be 50% less than what is charged
724	for other medical cannabis pharmacies.
725	(4) The department may not issue a license to operate a medical cannabis pharmacy to an
726	applicant if an individual described in Subsection (2)(b)(ii):
727	(a) has been convicted under state or federal law of:
728	(i) a felony in the preceding 10 years; or
729	(ii) after December 3, 2018, a misdemeanor for drug distribution;
730	(b) is younger than 21 years old; or
731	(c) after September 23, 2019, until January 1, 2023, is actively serving as a legislator.
732	(5)(a) If an applicant for a medical cannabis pharmacy license under this section holds
733	another license under this chapter, the department may not give preference to the
734	applicant based on the applicant's status as a holder of the license.
735	(b) If an applicant for a medical cannabis pharmacy license under this section holds a
736	license to operate a cannabis cultivation facility under this section, the department
737	may give consideration to the applicant's status as a holder of the license if:
738	(i) the applicant demonstrates that a decrease in costs to patients is more likely to
739	result from the applicant's vertical integration than from a more competitive
740	marketplace; and
741	(ii) the department finds multiple other factors, in addition to the existing license, that
742	support granting the new license.

- 743 (6) The licensing board may revoke a license under this part:
- 744 (a) if the medical cannabis pharmacy does not begin operations within one year after the 745 day on which the department issues an announcement of the department's intent to 746 award a license to the medical cannabis pharmacy;
  - (b) after the third the same violation of this chapter in any of the licensee's licensed cannabis production establishments or medical cannabis pharmacies;
  - (c) if an individual described in Subsection (2)(b)(ii) is convicted, while the license is active, under state or federal law of:
  - (i) a felony; or
    - (ii) after December 3, 2018, a misdemeanor for drug distribution;
  - (d) if the licensee fails to provide the information described in Subsection (2)(b)(vi) at the time of application, or fails to supplement the information described in Subsection (2)(b)(vi) with any investigation or adverse action that occurs after the submission of the application within 14 calendar days after the licensee receives notice of the investigation or adverse action;
  - (e) if the medical cannabis pharmacy demonstrates a willful or reckless disregard for the requirements of this chapter or the rules the department makes in accordance with this chapter;
  - (f) if, after a change of ownership described in Subsection (11)(c), the department determines that the medical cannabis pharmacy no longer meets the minimum standards for licensure and operation of the medical cannabis pharmacy described in this chapter; or
  - (g) if through an investigation conducted under Subsection 4-41a-201.1(11) and in accordance with Title 63G, Chapter 4, Administrative Procedures Act, the board finds that the licensee has participated in anticompetitive business practices.
  - (7)(a) A person who receives a medical cannabis pharmacy license under this chapter, if the municipality or county where the licensed medical cannabis pharmacy will be located requires a local land use permit, shall submit to the department a copy of the licensee's approved application for the land use permit within 120 days after the day on which the department issues the license.
    - (b) If a licensee fails to submit to the department a copy the licensee's approved land use permit application in accordance with Subsection (7)(a), the department may revoke the licensee's license.
  - (8) The department shall deposit the proceeds of a fee imposed by this section into the

777	Qualified Production Enterprise Fund.
778	(9) The department shall begin accepting applications under this part on or before March 1,
779	2020.
780	(10)(a) The department's authority to issue a license under this section is plenary and is
781	not subject to review.
782	(b) Notwithstanding Subsection (2), the decision of the department to award a license to
783	an applicant is not subject to:
784	(i) Title 63G, Chapter 6a, Part 16, Protests; or
785	(ii) Title 63G, Chapter 6a, Part 17, Procurement Appeals Board.
786	(11)(a) A medical cannabis pharmacy license is not transferrable or assignable.
787	(b) A medical cannabis pharmacy shall report in writing to the department no later than
788	10 business days before the date of any change of ownership of the medical cannabis
789	pharmacy.
790	(c) If the ownership of a medical cannabis pharmacy changes by 50% or more:
791	(i) concurrent with the report described in Subsection (11)(b), the medical cannabis
792	pharmacy shall submit a new application described in Subsection (2)(b), subject to
793	Subsection (2)(c);
794	(ii) within 30 days of the submission of the application, the department shall:
795	(A) conduct an application review; and
796	(B) award a license to the medical cannabis pharmacy for the remainder of the
797	term of the medical cannabis pharmacy's license before the ownership change
798	if the medical cannabis pharmacy meets the minimum standards for licensure
799	and operation of the medical cannabis pharmacy described in this chapter; and
800	(iii) if the department approves the license application, notwithstanding Subsection
801	(3), the medical cannabis pharmacy shall pay a license fee that the department sets
802	in accordance with Section 63J-1-504 in an amount that covers the department's
803	cost of conducting the application review.
804	Section 11. Section <b>4-41a-1003</b> is amended to read:
805	4-41a-1003 . Renewal - Notice of available license.
806	(1)(a) The department shall renew a license [under Sections 4-41a-1001 through
807	4-41a-1005] issued under this part every year if, at the time of renewal:
808	[(a)] (i) the licensee meets the requirements of Section 4-41a-1001;
809	[(b)] (ii) the licensee pays the department a license renewal fee in an amount that,
810	subject to Subsection 4-41a-1004(5), the department sets in accordance with

811	Section 63J-1-504; and
812	[(e)] (iii) if the medical cannabis pharmacy changes the operating plan described in
813	Section 4-41a-1004 that the department approved under Subsection
814	4-41a-1001(2)(b)(iv), the department approves the new operating plan.
815	(b) A license fee for a medical cannabis pharmacy located in a county of the third,
816	fourth, fifth, or sixth class shall be 50% less than what is charged for other medical
817	cannabis pharmacies.
818	(2)(a) If a licensed medical cannabis pharmacy abandons the medical cannabis
819	pharmacy's license, the department shall publish notice of an available license[-], for
820	the geographic area in which the medical cannabis pharmacy license is available, as a
821	class A notice under Section 63G-30-102, for at least seven days.
822	(b) The department may establish criteria, in collaboration with the Division of
823	Professional Licensing and the Board of Pharmacy and in accordance with Title 63G,
824	Chapter 3, Utah Administrative Rulemaking Act, to identify the medical cannabis
825	pharmacy actions that constitute abandonment of a medical cannabis pharmacy
826	license.
827	(3) If the department has not completed the necessary processes to make a determination on
828	a license renewal under Subsections (1)(a) and (c) before the expiration of a license, the
829	department may issue a conditional medical cannabis pharmacy license to a licensed
830	medical cannabis pharmacy that has applied for license renewal under this section and
831	paid the fee described in Subsection (1)(b).
832	Section 12. Section <b>4-41a-1005</b> is amended to read:
833	4-41a-1005 . Maximum number of licenses.
834	(1)(a) [Except as provided in Subsection (1)(b) or (d), if a sufficient number of
835	applicants apply, the department] The licensing board shall issue up to [15] 17 medical
836	cannabis pharmacy licenses in accordance with this section including the two medical
837	cannabis pharmacy licenses in accordance with Section 4-41a-1006.
838	(b) The medical cannabis ombudsman shall select the entities to receive a license in
839	accordance with this chapter.
840	(c) The medical cannabis ombudsman may choose to select entities as an entity is
841	qualified for a license and in accordance with Subsection (2)(c).
842	[(b) If an insufficient number of qualified applicants apply for the available number of
843	medical cannabis pharmacy licenses, the department shall issue a medical cannabis
844	pharmacy license to each qualified applicant.]

845	[(e) The department may issue the licenses described in Subsection (1)(a) in accordance
846	with this Subsection (1)(c).]
847	[(i) Using one procurement process, the department may issue eight licenses to an
848	initial group of medical cannabis pharmacies and six licenses to a second group of
849	medical cannabis pharmacies.]
850	[(ii) The department shall:]
851	[(A) divide the state into no less than four geographic regions, set by the
852	department in rule;]
853	[(B) issue at least one license in each geographic region during each phase of
854	issuing licenses; and]
855	[(C) complete the process of issuing medical cannabis pharmacy licenses no later
856	than July 1, 2020.]
857	[(iii) In issuing a 15th license under Subsection (1), the department shall ensure that
858	the license recipient will locate the medical cannabis pharmacy within Dagget,
859	Duchesne, Uintah, Carbon, Sevier, Emery, Grand, or San Juan County.]
860	[(d)(i) The department may issue licenses to operate a medical cannabis pharmacy in
861	addition to the licenses described in Subsection (1)(a) if the department
862	determines, in consultation with the Department of Health and Human Services
863	and after an annual or more frequent analysis of the current and anticipated market
864	for medical cannabis, that each additional license is necessary to provide an
865	adequate supply, quality, or variety of medical cannabis to medical cannabis
866	cardholders.]
867	[(ii) The department shall:]
868	[(A) in accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking
869	Act, make rules to establish criteria and processes for the consultation,
870	analysis, and application for a license described in Subsection (1)(d)(i); and]
871	[(B) report to the Executive Appropriations Committee of the Legislature before
872	each time the department issues an additional license under Subsection
873	(1)(d)(i) regarding the results of the consultation and analysis described in
874	Subsection (1)(d)(i) and the application of the criteria described in Subsection
875	$\frac{(1)(d)(ii)(A)}{(ii)(A)}$
876	(2)(a) [If there are more qualified applicants than there are available licenses for medical
877	eannabis pharmacies, the department] The medical cannabis ombudsman shall:
878	(i) evaluate each applicant and award the license to the applicant that best

879	demonstrates:
880	(A) experience with establishing and successfully operating a business that
881	involves complying with a regulatory environment, tracking inventory, and
882	training, evaluating, and monitoring employees;
883	(B) an operating plan that will best ensure the safety and security of patrons and
884	the community;
885	(C) positive connections to the local community;
886	(D) the suitability of the proposed location and the location's accessibility for
887	qualifying patients;
888	(E) the extent to which the applicant can increase efficiency and reduce the cost of
889	medical cannabis for patients; and
890	(F) a strategic plan described in Subsection 4-41a-1004(7) that has a
891	comparatively high likelihood of success; and
892	(ii) ensure a geographic dispersal among licensees that is sufficient to reasonably
893	maximize access to the largest number of medical cannabis cardholders.
894	(b) In making the evaluation described in Subsection (2)(a), the [department] medical
895	cannabis ombudsman may give increased consideration to applicants who indicate a
896	willingness to:
897	(i) site a medical cannabis pharmacy in an area or population center designated as a
898	medically underserved area or population as determined by the federal Health
899	Resources and Services Administration;
900	(ii) operate as a home delivery medical cannabis pharmacy that accepts electronic
901	medical cannabis orders[that the state central patient portal facilitates]; and
902	[(ii)] (iii) accept payments through:
903	(A) a payment provider that the Division of Finance approves, in consultation
904	with the state treasurer, in accordance with Section 4-41a-108; or
905	(B) a financial institution in accordance with Subsection 4-41a-108(4).
906	(c) Except for the licenses described in Section 26B-4-249, before each new license may
907	be issued under this section, the medical cannabis ombudsman shall:
908	(i) consider the number of active patients in the program;
909	(ii) geographic locations of current medical cannabis pharmacies; and
910	(iii) consult with other government agencies, licensees, and other stakeholders to
911	determine the economic impact of an additional license.
912	(3) The [department] medical cannabis ombudsman may conduct a face-to-face interview

913	with an applicant for a license that the [department] medical cannabis ombudsman
914	evaluates under Subsection (2).
915	Section 13. Section <b>4-41a-1006</b> is enacted to read:
916	4-41a-1006 . Licensees selected by medical cannabis ombudsman.
917	(1) Upon receiving a recommendation from the medical cannabis ombudsman under
918	Section 26B-4-248, the licensing board shall issue a license to the entity.
919	(2) An entity selected for a license under Section 26B-4-248 is subject to all of the
920	applicable requirements of this chapter and Title 26B, Chapter 4, Part 2, Cannabinoid
921	Research and Medical Cannabis.
922	(3) The department shall ensure compliance with Subsection 26B-4-248(3)(e).
923	Section 14. Section <b>4-41a-1101</b> is amended to read:
924	4-41a-1101 . Operating requirements General.
925	(1)(a) A medical cannabis pharmacy shall operate:
926	(i) at the physical address provided to the department under Section 4-41a-1001; and
927	(ii) in accordance with the operating plan provided to the department under Section
928	4-41a-1001 and, if applicable, Section 4-41a-1004.
929	(b) A medical cannabis pharmacy shall notify the department before a change in the
930	medical cannabis pharmacy's physical address or operating plan.
931	(2) An individual may not enter a medical cannabis pharmacy unless the individual:
932	(a) is at least 18 years old or is an emancipated minor under Section 80-7-105; and
933	(b) except as provided in Subsection (4):
934	(i) possesses a valid:
935	(A) medical cannabis pharmacy agent registration card;
936	(B) pharmacy medical provider registration card; or
937	(C) medical cannabis card;
938	(ii) is an employee of the department performing an inspection under Section
939	4-41a-1103; or
940	(iii) is another individual as the department provides.
941	(3) A medical cannabis pharmacy may not employ an individual who is younger than 21
942	years old.
943	(4) Notwithstanding Subsection (2)(a), a medical cannabis pharmacy may authorize an
944	individual who is not a medical cannabis pharmacy agent or pharmacy medical provider
945	to access the medical cannabis pharmacy if the medical cannabis pharmacy tracks and
946	monitors the individual at all times while the individual is at the medical cannabis

947	pharmacy and maintains a record of the individual's access.
948	(5) A medical cannabis pharmacy shall operate in a facility that has:
949	(a) a single, secure public entrance;
950	(b) a security system with a backup power source that:
951	(i) detects and records entry into the medical cannabis pharmacy; and
952	(ii) provides notice of an unauthorized entry to law enforcement when the medical
953	cannabis pharmacy is closed; and
954	(c) a lock on each area where the medical cannabis pharmacy stores [eannabis or a
955	eannabis product] medical cannabis.
956	(6) A medical cannabis pharmacy shall post, both clearly and conspicuously in the medical
957	cannabis pharmacy, the limit on the purchase of cannabis described in Subsection
958	4-41a-1102(2).
959	(7) Except for an emergency situation described in Subsection 26B-4-213(3)(c), a medical
960	cannabis pharmacy may not allow any individual to consume cannabis on the property
961	or premises of the medical cannabis pharmacy.
962	(8) A medical cannabis pharmacy may not sell [eannabis or a cannabis product] medical
963	cannabis without first indicating on the [cannabis or cannabis product] medical cannabis
964	label the name of the medical cannabis pharmacy.
965	(9)(a) Each medical cannabis pharmacy shall retain in the pharmacy's records the
966	following information regarding each recommendation underlying a transaction:
967	(i) the recommending medical provider's name, address, and telephone number;
968	(ii) the patient's name and address;
969	(iii) the date of issuance;
970	(iv) directions of use and dosing guidelines or an indication that the recommending
971	medical provider did not recommend specific directions of use or dosing
972	guidelines; and
973	(v) if the patient did not complete the transaction, the name of the medical cannabis
974	cardholder who completed the transaction.
975	(b)(i) Except as provided in Subsection (9)(b)(iii), a medical cannabis pharmacy may
976	not sell medical cannabis unless the medical cannabis has a label securely affixed
977	to the container indicating the following minimum information:
978	(A) the name, address, and telephone number of the medical cannabis pharmacy;
979	(B) the unique identification number that the medical cannabis pharmacy assigns;
980	(C) the date of the sale;

981	(D) the name of the patient;
982	(E) the name of the recommending medical provider who recommended the
983	medical cannabis treatment;
984	(F) directions for use and cautionary statements, if any;
985	(G) the amount dispensed and the cannabinoid content;
986	(H) the suggested use date;
987	(I) for unprocessed cannabis flower, the legal use termination date; and
988	(J) any other requirements that the department determines, in consultation with the
989	Division of Professional Licensing and the Board of Pharmacy.
990	(ii) A medical cannabis pharmacy is exempt from the requirement to provide the
991	following information under Subsection (9)(b)(i) if the information is already
992	provided on the product label that a cannabis production establishment affixes:
993	(A) a unique identification number;
994	(B) directions for use and cautionary statements;
995	(C) amount and cannabinoid content; and
996	(D) a suggested use date.
997	(iii) If the size of a medical cannabis container does not allow sufficient space to
998	include the labeling requirements described in Subsection (9)(b)(i), the medical
999	cannabis pharmacy may provide the following information described in
1000	Subsection (9)(b)(i) on a supplemental label attached to the container or an
1001	informational enclosure that accompanies the container:
1002	(A) the cannabinoid content;
1003	(B) the suggested use date; and
1004	(C) any other requirements that the department determines.
1005	(iv) A medical cannabis pharmacy may sell medical cannabis to another medical
1006	cannabis pharmacy without a label described in Subsection (9)(b)(i).
1007	(10) A pharmacy medical provider or medical cannabis pharmacy agent shall:
1008	(a) upon receipt of an order from a limited medical provider in accordance with
1009	Subsections 26B-4-204(1)(b) through (d):
1010	(i) for a written order or an electronic order under circumstances that the department
1011	determines, contact the limited medical provider or the limited medical provider's
1012	office to verify the validity of the recommendation; and
1013	(ii) for an order that the pharmacy medical provider or medical cannabis pharmacy
1014	agent verifies under Subsection (10)(a)(i) or an electronic order that is not subject

1015	to verification under Subsection (10)(a)(i), enter the limited medical provider's
1016	recommendation or renewal, including any associated directions of use, dosing
1017	guidelines, or caregiver indication, in the state electronic verification system;
1018	(b) in processing an order for a holder of a conditional medical cannabis card described
1019	in Subsection 26B-4-213(1)(b) that appears irregular or suspicious in the judgment of
1020	the pharmacy medical provider or medical cannabis pharmacy agent, contact the
1021	recommending medical provider or the recommending medical provider's office to
1022	verify the validity of the recommendation before processing the cardholder's order;
1023	(c) unless the medical cannabis cardholder has had a consultation under Subsection
1024	26B-4-231(5), verbally offer to a medical cannabis cardholder at the time of a
1025	purchase of [eannabis, a cannabis product,] medical cannabis or a medical cannabis
1026	device, personal counseling with the pharmacy medical provider; and
1027	(d) provide a telephone number or website by which the cardholder may contact a
1028	pharmacy medical provider for counseling.
1029	(11)(a) A medical cannabis pharmacy may create a medical cannabis disposal program
1030	that allows an individual to deposit unused or excess medical cannabis or cannabis
1031	residue from a medical cannabis device in a locked box or other secure receptacle
1032	within the medical cannabis pharmacy.
1033	(b) A medical cannabis pharmacy with a disposal program described in Subsection
1034	(11)(a) shall ensure that only a medical cannabis pharmacy agent or pharmacy
1035	medical provider can access deposited medical cannabis.
1036	(c) A medical cannabis pharmacy shall dispose of any deposited medical cannabis by:
1037	(i) rendering the deposited medical cannabis unusable and unrecognizable before
1038	transporting deposited medical cannabis from the medical cannabis pharmacy; and
1039	(ii) disposing of the deposited medical cannabis in accordance with:
1040	(A) federal and state law, rules, and regulations related to hazardous waste;
1041	(B) the Resource Conservation and Recovery Act, 42 U.S.C. Sec. 6991 et seq.;
1042	(C) Title 19, Chapter 6, Part 5, Solid Waste Management Act; and
1043	(D) other regulations that the department makes in accordance with Title 63G,
1044	Chapter 3, Utah Administrative Rulemaking Act.
1045	(12) A medical cannabis pharmacy:
1046	(a) shall employ a pharmacist who is licensed under Title 58, Chapter 17b, Pharmacy
1047	Practice Act, as a pharmacy medical provider;
1048	(b) may employ a physician who has the authority to write a prescription and is licensed

1049	under Title 58, Chapter 67, Utah Medical Practice Act, or Title 58, Chapter 68, V	Utah
1050	Osteopathic Medical Practice Act, as a pharmacy medical provider;	
1051	(c) shall ensure that a pharmacy medical provider described in Subsection (12)(a) w	orks
1052	onsite during all business hours;	
1053	(d) shall designate one pharmacy medical provider described in Subsection (12)(a) a	as the
1054	pharmacist-in-charge to oversee the operation of and generally supervise the me	dical
1055	cannabis pharmacy;[-and]	
1056	(e) shall allow the pharmacist-in-charge to determine which [eannabis and cannabis	
1057	products] medical cannabis products the medical cannabis pharmacy maintains i	n the
1058	medical cannabis pharmacy's inventory[-];	
1059	(f) if a patient product information insert is available, shall provide a patient who	
1060	purchases a medical cannabis product the medical cannabis product's patient pro-	duct
1061	information insert using any of the following methods:	
1062	(i) a physical document;	
1063	(ii) an email message;	
1064	(iii) a text message; or	
1065	(iv) a quick response code;	
1066	(g) for each medical cannabis product sold by the medical cannabis pharmacy, shall	<u>:</u>
1067	(i) allow a medical cannabis cardholder located in the pharmacy to view the bac	<u>k</u>
1068	panel of the product when requested; and	
1069	(ii) beginning July 1, 2025, include a picture of the back panel of the product or	the
1070	medical cannabis pharmacy's website; and	
1071	(h) may not allow a recommending medical provider to recommend medical cannab	ois .
1072	within 500 feet of the medical cannabis pharmacy's property line.	
1073	(13) The department shall establish by rule, in accordance with Title 63G, Chapter 3, Ut	ah
1074	Administrative Rulemaking Act, protocols for a recall of cannabis and cannabis produced	ducts
1075	by a medical cannabis pharmacy.	
1076	Section 15. Section <b>4-41a-1201</b> is amended to read:	
1077	4-41a-1201. Medical cannabis home delivery designation.	
1078	(1) The department may designate a medical cannabis pharmacy as a home delivery	
1079	medical cannabis pharmacy if the department determines that the medical cannabis	
1080	pharmacy's operating plan demonstrates the functional and technical ability to:	
1081	(a) safely conduct transactions for medical cannabis shipments;	
1082	(b) accept electronic medical cannabis orders[that the state central patient portal	

1083	facilitates]; and
1084	(c) accept payments through:
1085	(i) a payment provider that the Division of Finance approves, in consultation with the
1086	state treasurer, in accordance with Section 26-61a-603; or
1087	(ii) a financial institution in accordance with Subsection 26-61a-603(4).
1088	(2) An applicant seeking a designation as a home delivery medical cannabis pharmacy shall
1089	identify in the applicant's operating plan any information relevant to the department's
1090	evaluation described in Subsection (1), including:
1091	(a) the name and contact information of the payment provider;
1092	(b) the nature of the relationship between the prospective licensee and the payment
1093	provider;
1094	(c) the processes of the following to safely and reliably conduct transactions for medical
1095	cannabis shipments:
1096	(i) the prospective licensee; and
1097	(ii) the electronic payment provider or the financial institution described in
1098	Subsection (1)(c); and
1099	(d) the ability of the licensee to comply with the department's rules regarding the secure
1100	transportation and delivery of medical cannabis [or medical cannabis product] to a
1101	medical cannabis cardholder.
1102	(3) Notwithstanding any county or municipal ordinance, a medical cannabis pharmacy that
1103	the department designates as a home delivery medical cannabis pharmacy may deliver
1104	medical cannabis shipments in accordance with this part.
1105	Section 16. Section 4-41a-1202 is amended to read:
1106	4-41a-1202 . Home delivery of medical cannabis shipments Medical cannabis
1107	couriers License.
1108	(1) The department shall make rules, in accordance with Title 63G, Chapter 3, Utah
1109	Administrative Rulemaking Act, to ensure the safety, security, and efficiency of a home
1110	delivery medical cannabis pharmacy's fulfillment of electronic medical cannabis orders[
1111	that the state central patient portal facilitates], including rules regarding the safe and
1112	controlled delivery of medical cannabis shipments.
1113	(2) A person may not operate as a medical cannabis courier without a license that the
1114	department issues under this section.
1115	(3)(a) Subject to Subsections (5) and (6), the department shall issue a license to operate
1116	as a medical cannabis courier to an applicant who is eligible for a license under this

1117	section.
1118	(b) An applicant is eligible for a license under this section if the applicant submits to the
1119	department:
1120	(i) the name and address of an individual who:
1121	(A) has a financial or voting interest of 10% or greater in the proposed medical
1122	cannabis courier; or
1123	(B) has the power to direct or cause the management or control of a proposed
1124	cannabis production establishment;
1125	(ii) an operating plan that includes operating procedures to comply with the operating
1126	requirements for a medical cannabis courier described in this chapter; and
1127	(iii) an application fee in an amount that, subject to Subsection 4-41a-104(5), the
1128	department sets in accordance with Section 63J-1-504.
1129	(4) If the department determines that an applicant is eligible for a license under this section,
1130	the department shall:
1131	(a) charge the applicant an initial license fee in an amount that, subject to Subsection
1132	4-41a-104(5), the department sets in accordance with Section 63J-1-504; and
1133	(b) notify the Department of Public Safety of the license approval and the names of each
1134	individual described in Subsection (3)(b)(i).
1135	(5) The department may not issue a license to operate as a medical cannabis courier to an
1136	applicant if an individual described in Subsection (3)(b)(i):
1137	(a) has been convicted under state or federal law of:
1138	(i) a felony in the preceding 10 years; or
1139	(ii) after September 23, 2019, a misdemeanor for drug distribution; or
1140	(b) is younger than 21 years old.
1141	(6) The department may revoke a license under this part if:
1142	(a) the medical cannabis courier does not begin operations within one year after the day
1143	on which the department issues the initial license;
1144	(b) the medical cannabis courier makes the same violation of this chapter three times;
1145	(c) an individual described in Subsection (3)(b)(i) is convicted, while the license is
1146	active, under state or federal law of:
1147	(i) a felony; or
1148	(ii) after September 23, 2019, a misdemeanor for drug distribution; or
1149	(d) after a change of ownership described in Subsection (14)(c), the department
1150	determines that the medical cannabis courier no longer meets the minimum standards

1151	for licensure and operation of the medical cannabis courier described in this chapter.
1152	(7) The department shall deposit the proceeds of a fee imposed by this section in the
1153	Qualified Production Enterprise Fund.
1154	(8) The department's authority to issue a license under this section is plenary and is not
1155	subject to review.
1156	(9) Each applicant for a license as a medical cannabis courier shall submit, at the time of
1157	application, from each individual who has a financial or voting interest of 10% or
1158	greater in the applicant or who has the power to direct or cause the management or
1159	control of the applicant:
1160	(a) a fingerprint card in a form acceptable to the Department of Public Safety;
1161	(b) a signed waiver in accordance with Subsection 53-10-108(4) acknowledging the
1162	
	registration of the individual's fingerprints in the Federal Bureau of Investigation
1163	Next Generation Identification System's Rap Back Service; and
1164	(c) consent to a fingerprint background check by:
1165	(i) the Bureau of Criminal Identification; and
1166	(ii) the Federal Bureau of Investigation.
1167	(10) The Bureau of Criminal Identification shall:
1168	(a) check the fingerprints the applicant submits under Subsection (9) against the
1169	applicable state, regional, and national criminal records databases, including the
1170	Federal Bureau of Investigation Next Generation Identification System;
1171	(b) report the results of the background check to the department;
1172	(c) maintain a separate file of fingerprints that applicants submit under Subsection (9)
1173	for search by future submissions to the local and regional criminal records databases
1174	including latent prints;
1175	(d) request that the fingerprints be retained in the Federal Bureau of Investigation Next
1176	Generation Identification System's Rap Back Service for search by future
1177	submissions to national criminal records databases, including the Next Generation
1178	Identification System and latent prints; and
1179	(e) establish a privacy risk mitigation strategy to ensure that the department only
1180	receives notifications for an individual with whom the department maintains an
1181	authorizing relationship.
1182	(11) The department shall:
1183	(a) assess an individual who submits fingerprints under Subsection (9) a fee in an
1184	amount that the department sets in accordance with Section 63J-1-504 for the

1185	services that the Bureau of Criminal Identification or another authorized agency
1186	provides under this section; and
1187	(b) remit the fee described in Subsection (11)(a) to the Bureau of Criminal Identification.
1188	(12) The department shall renew a license under this section every year if, at the time of
1189	renewal:
1190	(a) the licensee meets the requirements of this section; and
1191	(b) the licensee pays the department a license renewal fee in an amount that, subject to
1192	Subsection 4-41a-104(5), the department sets in accordance with Section 63J-1-504.
1193	(13) A person applying for a medical cannabis courier license shall submit to the
1194	department a proposed operating plan that complies with this section and that includes:
1195	(a) a description of the physical characteristics of any proposed facilities, including a
1196	floor plan and an architectural elevation, and delivery vehicles;
1197	(b) a description of the credentials and experience of each officer, director, or owner of
1198	the proposed medical cannabis courier;
1199	(c) the medical cannabis courier's employee training standards;
1200	(d) a security plan; and
1201	(e) storage and delivery protocols, both short and long term, to ensure that medical
1202	cannabis shipments are stored and delivered in a manner that is sanitary and
1203	preserves the integrity of the cannabis.
1204	(14)(a) A medical cannabis courier license is not transferable or assignable.
1205	(b) A medical cannabis courier shall report in writing to the department no later than 10
1206	business days before the date of any change of ownership of the medical cannabis
1207	courier.
1208	(c) If the ownership of a medical cannabis courier changes by 50% or more:
1209	(i) concurrent with the report described in Subsection (14)(b), the medical cannabis
1210	courier shall submit a new application described in Subsection (3)(b);
1211	(ii) within 30 days of the submission of the application, the department shall:
1212	(A) conduct an application review; and
1213	(B) award a license to the medical cannabis courier for the remainder of the term
1214	of the medical cannabis courier's license before the ownership change if the
1215	medical cannabis courier meets the minimum standards for licensure and
1216	operation of the medical cannabis courier described in this chapter; and
1217	(iii) if the department approves the license application, notwithstanding Subsection
1218	(4) the medical cannabis courier shall pay a license fee that the department sets in

1219	accordance with Section 63J-1-504 in an amount that covers the board's cost of
1220	conducting the application review.
1221	(15)(a) Except as provided in Subsection(15)(b), a person may not advertise regarding
1222	the transportation of medical cannabis.
1223	(b) Notwithstanding Subsection (14)(a) and subject to Section 4-41a-109, a licensed
1224	home delivery medical cannabis pharmacy or a licensed medical cannabis courier
1225	may advertise:
1226	(i) a green cross;
1227	(ii) the pharmacy's or courier's name and logo; and
1228	(iii) that the pharmacy or courier is licensed to transport medical cannabis shipments
1229	Section 17. Section 4-41a-1203 is amended to read:
1230	4-41a-1203. Medical cannabis shipment transportation.
1231	(1) The department shall ensure that each home delivery medical cannabis pharmacy is
1232	capable of delivering, directly or through a medical cannabis courier, medical cannabis
1233	shipments in a secure manner.
1234	(2)(a) A home delivery medical cannabis pharmacy may contract with a licensed
1235	medical cannabis courier to deliver medical cannabis shipments to fulfill electronic
1236	medical cannabis orders[ that the state central patient portal facilitates].
1237	(b) If a home delivery medical cannabis pharmacy enters into a contract described in
1238	Subsection (2)(a), the pharmacy shall:
1239	(i) impose security and personnel requirements on the medical cannabis courier
1240	sufficient to ensure the security and safety of medical cannabis shipments; and
1241	(ii) provide regular oversight of the medical cannabis courier.
1242	(3) Notwithstanding Subsection 4-41a-404(1), an individual may transport a medical
1243	cannabis shipment if the individual is:
1244	(a) a registered pharmacy medical provider;
1245	(b) a registered medical cannabis pharmacy agent; or
1246	(c) a registered agent of the medical cannabis courier described in Subsection (2).
1247	(4) An individual transporting a medical cannabis shipment under Subsection (3) shall
1248	comply with the requirements of Subsection 4-41a-404(3).
1249	(5) In addition to the requirements in Subsections (3) and (4), the department may establish
1250	by rule, in collaboration with the Division of Professional Licensing and the Board of
1251	Pharmacy and in accordance with Title 63G, Chapter 3, Utah Administrative
1252	Rulemaking Act, requirements for transporting medical cannabis shipments that are

1253	related to safety for human consumption of [cannabis or a cannabis product] medical
1254	cannabis.
1255	(6)(a) It is unlawful for an individual to transport a medical cannabis shipment with a
1256	manifest that does not meet the requirements of Subsection (4).
1257	(b) Except as provided in Subsection (6)(d), an individual who violates Subsection (6)(a)
1258	is:
1259	(i) guilty of an infraction; and
1260	(ii) subject to a \$100 fine.
1261	(c) An individual who is guilty of a violation described in Subsection (6)(b) is not guilty
1262	of a violation of Title 58, Chapter 37, Utah Controlled Substances Act, for the
1263	conduct underlying the violation described in Subsection (6)(b).
1264	(d) If the individual described in Subsection (6)(a) is transporting more cannabis,
1265	cannabis product, or medical cannabis devices than the manifest identifies, except for
1266	a de minimis administrative error:
1267	(i) this chapter does not apply; and
1268	(ii) the individual is subject to penalties under Title 58, Chapter 37, Utah Controlled
1269	Substances Act.
1270	Section 18. Section <b>4-41a-1206</b> is amended to read:
1271	4-41a-1206. Closed-door medical cannabis pharmacy.
1272	(1)(a) Subject to Subsections (1)(b) and (c), a home delivery medical cannabis pharmacy
1273	may open a single closed-door medical cannabis pharmacy.
1274	(b) A home delivery medical cannabis pharmacy may not open a closed-door medical
1275	cannabis pharmacy unless the home delivery medical cannabis pharmacy:
1276	(i) has an operating plan that includes a closed-door medical cannabis pharmacy; and
1277	(ii) obtains a license issued by the department for a closed-door medical cannabis
1278	pharmacy.
1279	(c) An entity that owns multiple home delivery medical cannabis pharmacies may open
1280	only one closed-door medical cannabis pharmacy.
1281	(d) The department may institute a fee in accordance with Section 63J-1-504 to
1282	administer this section.
1283	(2) A home delivery medical cannabis pharmacy that opens a closed-door medical cannabis
1284	pharmacy under Subsection (1) shall ensure:
1285	(a) that a pharmacy medical provider who is a licensed pharmacist:
1286	(i) is directly supervising the packaging of an order; and

1287	(ii) is present in the closed-door medical cannabis pharmacy when an order is
1288	packaged for delivery; and
1289	(b) all record keeping requirements, labeling requirements, and patient counseling
1290	requirements described in this chapter and Title 26B, Chapter 4, Part 2, Cannabinoid
1291	Research and Medical Cannabis, are satisfied before sending out an order.
1292	(3) An individual who prepares an order at a closed-door medical cannabis pharmacy under
1293	this section shall be registered as:
1294	(a) a pharmacy medical provider; or
1295	(b) a medical cannabis pharmacy agent.
1296	(4)(a) A closed-door medical cannabis pharmacy shall operate:
1297	(i) except as provided in Subsection (4)(b), in a facility that is accessible only by an
1298	individual who is a pharmacy medical provider or a medical cannabis pharmacy
1299	agent; and
1300	(ii) at a physical address in accordance with Subsection (6).
1301	(b) A closed-door medical cannabis pharmacy may authorize an individual who is at
1302	least 18 years old and is not a pharmacy medical provider or a cannabis pharmacy
1303	agent to access the closed-door medical cannabis pharmacy if the closed-door
1304	medical cannabis pharmacy:
1305	(i) tracks and monitors the individual at all times while the individual is at the
1306	closed-door medical cannabis pharmacy; and
1307	(ii) maintains a record of the individual's access, including arrival and departure.
1308	(c) A closed-door medical cannabis pharmacy shall operate in a facility that has:
1309	(i) a single, secure public entrance; and
1310	(ii) a security system with a backup power source that:
1311	(A) detects and records entry into the closed-door medical cannabis pharmacy;
1312	(B) provides notice of an unauthorized entry to law enforcement when the
1313	closed-door medical cannabis pharmacy is closed; and
1314	(C) a lock or equivalent restrictive security feature on any area where the
1315	closed-door medical cannabis pharmacy stores a cannabis product.
1316	(d) A closed-door medical cannabis pharmacy shall ensure that any cannabis or cannabis
1317	products in the closed-door medical cannabis pharmacy that are intended for home
1318	delivery are separated in a manner that is readily distinguishable from any other
1319	cannabis or cannabis product in the facility.
1320	(5) A closed-door medical cannabis pharmacy may only provide cannabis or a cannabis

1321	product to an individual through a delivery that complies with this part.
1322	(6)(a) A person may not locate a closed-door medical cannabis pharmacy:
1323	(i) within 1,000 feet of a community location; or
1324	(ii) in or within 600 feet of a district that the relevant municipality or county has
1325	zoned as primarily residential.
1326	(b) The proximity requirements described in Subsection (6)(a) shall be measured from
1327	the nearest entrance to the closed-door medical cannabis pharmacy by following the
1328	shortest route of ordinary pedestrian travel to the property boundary of the
1329	community location or residential area.
1330	(c) The licensing board may grant a waiver to reduce the proximity requirements in
1331	Subsection (6)(a) by up to 20% if the licensing board determines that it is not
1332	reasonably feasible for the applicant to site the proposed closed-door medical
1333	cannabis pharmacy without the waiver.
1334	(d) An applicant for a license under this section shall provide evidence of compliance
1335	with the proximity requirements described in Subsection (6)(a).
1336	(7) When determining where a closed-door medical cannabis pharmacy may open, the
1337	licensing board:
1338	(a) shall utilize geographic regions created by the department through rule;
1339	(b) shall prioritize allowing entities that do not have a medical cannabis pharmacy in a
1340	region to open a closed-door medical cannabis pharmacy in the region;
1341	(c) of the total amount of closed-door medical cannabis pharmacies, may allow only
1342	three closed-door medical cannabis pharmacies to operate in counties of the first and
1343	second class as described in Section 17-50-501; and
1344	(d) for determining the three closed-door medical cannabis pharmacies described in
1345	Subsection (7)(c), consider the following:
1346	(i) the history of compliance with state law and rules for all licenses issued under this
1347	chapter;
1348	(ii) the medical cannabis pharmacy's willingness to offer a variety of brands and
1349	products;
1350	(iii) the ability of the operating plan to ensure the safety and security of the
1351	community;
1352	(iv) the suitability of the proposed location and the location's ability to serve the local
1353	community; and
1354	(v) any other relevant information determined through rule.

1355	(8) A closed-door medical cannabis pharmacy may not account for more than:
1356	(a) for an entity that holds a single medical cannabis pharmacy license, the greater of:
1357	(i) 35% of the medical cannabis pharmacy's total revenue; or
1358	(ii) \$2,000,000 in total revenue; or
1359	(b) for an entity that holds more than one medical cannabis pharmacy license, the greater
1360	of:
1361	(i) 35% of the total revenue of the entity's medical cannabis pharmacy that generates
1362	the most revenue; or
1363	(ii) \$2,000,000 in total revenue.
1364	(9) Notwithstanding any other provision of this section, the [department] licensing board
1365	may issue only [three] one closed-door medical cannabis pharmacy [licenses] license
1366	before July 1, 2027.
1367	(10) In accordance with Title 63G, Chapter 3, Utah Administrative Rulemaking Act, the
1368	department shall make rules to implement this section.
1369	Section 19. Section <b>26B-1-310</b> is amended to read:
1370	26B-1-310 . Qualified Patient Enterprise Fund Creation Revenue neutrality
1371	Uniform fee.
1372	(1) There is created an enterprise fund known as the "Qualified Patient Enterprise Fund."
1373	(2) The fund created in this section is funded from:
1374	(a) money the department deposits into the fund under Chapter 4, Part 2, Cannabinoid
1375	Research and Medical Cannabis;
1376	(b) appropriations the Legislature makes to the fund; and
1377	(c) the interest described in Subsection (3).
1378	(3) Interest earned on the fund shall be deposited into the fund.
1379	(4) Money deposited into the fund may only be used by:
1380	(a) the department to accomplish the department's responsibilities described in Chapter
1381	4, Part 2, Cannabinoid Research and Medical Cannabis; and
1382	(b) the Center for Medical Cannabis Research created in Section 53B-17-1402 to
1383	accomplish the Center for Medical Cannabis Research's responsibilities[-]; and
1384	(c) if there is remaining money in the fund balance on June 30 of each fiscal year after
1385	financial obligations under Subsections (4)(a) through (b) are met, an amount up to
1386	\$300,000, the medical cannabis ombudsman and available for expenditure the next
1387	fiscal year for the program described in Subsection 26B-4-248(4) and, subject to
1388	Subsection (7), the program's associated administrative costs.

1389	(5) The department shall set fees authorized under Chapter 4, Part 2, Cannabinoid Research
1390	and Medical Cannabis, in amounts that the department anticipates are necessary, in total,
1391	to cover the department's cost to implement Chapter 4, Part 2, Cannabinoid Research
1392	and Medical Cannabis.
1393	(6) The department may impose a uniform fee on each medical cannabis transaction in a
1394	medical cannabis pharmacy in an amount that, subject to Subsection (5), the department
1395	sets in accordance with Section 63J-1-504.
1396	(7) No more than 20% of the amount transferred under Subsection (4)(c) may be used for
1397	administrative costs.
1398	Section 20. Section <b>26B-1-435</b> is amended to read:
1399	26B-1-435 . Medical Cannabis Policy Advisory Board creation Membership
1400	Duties.
1401	(1) There is created within the department the Medical Cannabis Policy Advisory Board.
1402	(2)(a) The advisory board shall consist of the following members:
1403	(i) appointed by the executive director:
1404	(A) a qualified medical provider who has recommended medical cannabis to at
1405	least 100 patients before being appointed;
1406	[(B) a medical research professional;]
1407	[(C)] (B) a mental health specialist;
1408	[(D)] (C) an individual who represents an organization that advocates for medical
1409	cannabis patients;
1410	[(E)] (D) [an individual] a member of the general public who holds a medical
1411	cannabis patient card; and
1412	[(F)] (E) a member of the general public who does not hold a medical cannabis
1413	card;[ <del>-and</del> ]
1414	(ii) appointed by the commissioner of the Department of Agriculture and Food:
1415	(A) an individual who owns or operates a licensed cannabis cultivation facility, as
1416	defined in Section 4-41a-102;
1417	(B) an individual who owns or operates a licensed medical cannabis pharmacy;
1418	and
1419	(C) a law enforcement officer[-]; and
1420	(iii) a representative from the Center for Medical Cannabis Research created in
1421	Section 53B-14-1402, appointed by the Center for Medical Cannabis Research.
1422	(b) The commissioner of the Department of Agriculture and Food shall ensure that at

1423	least one individual appointed under Subsection (2)(a)(ii)(A) or (B) also owns or
1424	operates a licensed cannabis processing facility.
1425	(3)(a) Subject to Subsection (3)(b), a member of the advisory board shall serve for a four
1426	year term.
1427	(b) When appointing the initial membership of the advisory board, the executive director
1428	and the commissioner of the Department of Agriculture and Food shall coordinate to
1429	appoint four advisory board members to serve a term of two years to ensure that
1430	approximately half of the board is appointed every two years.
1431	(4)(a) If an advisory board member is no longer able to serve as a member, a new
1432	member shall be appointed in the same manner as the original appointment.
1433	(b) A member appointed in accordance with Subsection (4)(a) shall serve for the
1434	remainder of the unexpired term of the original appointment.
1435	(5)(a) A majority of the advisory board members constitutes a quorum.
1436	(b) The action of a majority of a quorum constitutes an action of the advisory board.
1437	(c) For a term lasting one year, the advisory board shall annually designate members of
1438	the advisory board to serve as chair and vice-chair.
1439	(d) When designating the chair and vice-chair, the advisory board shall ensure that at
1440	least one individual described Subsection (2)(a)(i) is appointed as chair or vice-chair.
1441	(6) An advisory board member may not receive compensation or benefits for the member's
1442	service on the advisory board but may receive per diem and reimbursement for travel
1443	expenses incurred as an advisory board member in accordance with:
1444	(a) Sections 63A-3-106 and 63A-3-107; and
1445	(b) rules made by the Division of Finance pursuant to Sections 63A-3-106 and
1446	63A-3-107.
1447	(7) The department shall:
1448	(a) provide staff support for the advisory board; and
1449	(b) assist the advisory board in conducting meetings.
1450	(8) The advisory board may recommend:
1451	(a) to the department or the Department of Agriculture and Food changes to current or
1452	proposed medical cannabis rules or statutes; and
1453	(b) to the appropriate legislative committee whether the advisory board supports a
1454	change to medical cannabis statutes.
1455	(9) The advisory board shall:
1456	(a) review any draft rule that is authorized under [this chapter] Chapter 4. Part 2.

1457 Cannabinoid Research and Medical Cannabis, or Title 4, Chapter 41a, Cannabis 1458 Production Establishments and Pharmacies; 1459 (b) consult with the Department of Agriculture and Food regarding the issuance of an additional: 1460 1461 (i) cultivation facility license under Section 4-41a-205; or 1462 (ii) pharmacy license under Section 4-41a-1005; 1463 (c) consult with the department regarding cannabis patient education; 1464 (d) consult regarding the reasonableness of any fees set by the department or the 1465 Department of Agriculture and Food that pertain to the medical cannabis program; 1466 and 1467 (e) consult regarding any issue pertaining to medical cannabis when asked by the 1468 department or the Utah Department of Agriculture and Food. 1469 Section 21. Section **26B-4-201** is amended to read: 1470 26B-4-201. Definitions. 1471 As used in this part: 1472 (1) "Active tetrahydrocannabinol" means THC, any THC analog, and 1473 tetrahydrocannabinolic acid. 1474 (2) "Administration of criminal justice" means the performance of detection, apprehension, 1475 detention, pretrial release, post-trial release, prosecution, and adjudication. 1476 (3) "Advertise" means information provided by a person in any medium: 1477 (a) to the public; and 1478 (b) that is not age restricted to an individual who is at least 21 years old. 1479 (4) "Advisory board" means the Medical Cannabis Policy Advisory Board created in 1480 Section 26B-1-435. 1481 (5) "Cannabis Research Review Board" means the Cannabis Research Review Board 1482 created in Section 26B-1-420. 1483 (6) "Cannabis" means marijuana. 1484 (7) "Cannabis processing facility" means the same as that term is defined in Section 1485 4-41a-102. 1486 (8) "Cannabis product" means a product that: 1487 (a) is intended for human use; and (b) contains cannabis or any tetrahydrocannabinol or THC analog in a total 1488 1489 concentration of 0.3% or greater on a dry weight basis. 1490 (9) "Cannabis production establishment" means the same as that term is defined in Section

1491	4-41a-102.
1492	(10) "Cannabis production establishment agent" means the same as that term is defined in
1493	Section 4-41a-102.
1494	(11) "Cannabis production establishment agent registration card" means the same as that
1495	term is defined in Section 4-41a-102.
1496	(12) "Conditional medical cannabis card" means an electronic medical cannabis card that
1497	the department issues in accordance with Subsection 26B-4-213(1)(b) to allow an
1498	applicant for a medical cannabis card to access medical cannabis during the department's
1499	review of the application.
1500	(13) "Controlled substance database" means the controlled substance database created in
1501	Section 58-37f-201.
1502	(14) "Delivery address" means the same as that term is defined in Section 4-41a-102.
1503	(15) "Department" means the Department of Health and Human Services.
1504	(16) "Designated caregiver" means:
1505	(a) an individual:
1506	(i) whom an individual with a medical cannabis patient card or a medical cannabis
1507	guardian card designates as the patient's caregiver; and
1508	(ii) who registers with the department under Section 26B-4-214; or
1509	(b)(i) a facility that an individual designates as a designated caregiver in accordance
1510	with Subsection 26B-4-214(1)(b); or
1511	(ii) an assigned employee of the facility described in Subsection 26B-4-214(1)(b)(ii).
1512	(17) "Directions of use" means recommended routes of administration for a medical
1513	cannabis treatment and suggested usage guidelines.
1514	(18) "Dosing guidelines" means a quantity range and frequency of administration for a
1515	recommended treatment of medical cannabis.
1516	(19) "Government issued photo identification" means any of the following forms of
1517	identification:
1518	(a) a valid state-issued driver license or identification card;
1519	(b) a valid United States federal-issued photo identification, including:
1520	(i) a United States passport;
1521	(ii) a United States passport card;
1522	(iii) a United States military identification card; or
1523	(iv) a permanent resident card or alien registration receipt card; or
1524	(c) a foreign passport.

1525	(20) "Home delivery medical cannabis pharmacy" means a medical cannabis pharmacy that
1526	the department authorizes, as part of the pharmacy's license, to deliver medical cannabis
1527	shipments to a delivery address to fulfill electronic orders[-that the state central patient
1528	portal facilitates].
1529	(21) "Inventory control system" means the system described in Section 4-41a-103.
1530	(22) "Legal dosage limit" means an amount that:
1531	(a) is sufficient to provide 30 days of treatment based on the dosing guidelines that the
1532	relevant recommending medical provider or [the state central patient portal or ]
1533	pharmacy medical provider, in accordance with Subsection 26B-4-230(5),
1534	recommends; and
1535	(b) may not exceed:
1536	(i) for unprocessed cannabis in a medicinal dosage form, 113 grams by weight; and
1537	(ii) for a cannabis product in a medicinal dosage form, a quantity that contains, in
1538	total, greater than 20 grams of active tetrahydrocannabinol.
1539	(23) "Legal use termination date" means a date on the label of a container of unprocessed
1540	cannabis flower:
1541	(a) that is 60 days after the date of purchase of the cannabis; and
1542	(b) after which, the cannabis is no longer in a medicinal dosage form outside of the
1543	primary residence of the relevant medical cannabis patient cardholder.
1544	(24) "Limited medical provider" means an individual who:
1545	(a) meets the recommending qualifications; and
1546	(b) has no more than 15 patients with a valid medical cannabis patient card as a result of
1547	the individual's recommendation, in accordance with Subsection 26B-4-204(1)(b).
1548	(25) "Marijuana" means the same as that term is defined in Section 58-37-2.
1549	(26) "Medical cannabis" or "medical cannabis product" means cannabis in a medicinal
1550	dosage form or a cannabis product in a medicinal dosage form.
1551	(27) "Medical cannabis card" means a medical cannabis patient card, a medical cannabis
1552	guardian card, a medical cannabis caregiver card, or a conditional medical cannabis card.
1553	(28) "Medical cannabis cardholder" means:
1554	(a) a holder of a medical cannabis card; or
1555	(b) a facility or assigned employee, described in Subsection (16)(b), only:
1556	(i) within the scope of the facility's or assigned employee's performance of the role of
1557	a medical cannabis patient cardholder's caregiver designation under Subsection
1558	26B-4-214(1)(b); and

1559	(ii) while in possession of documentation that establishes:
1560	(A) a caregiver designation described in Subsection 26B-4-214(1)(b);
1561	(B) the identity of the individual presenting the documentation; and
1562	(C) the relation of the individual presenting the documentation to the caregiver
1563	designation.
1564	(29) "Medical cannabis caregiver card" means an electronic document that a cardholder
1565	may print or store on an electronic device or a physical card or document that:
1566	(a) the department issues to an individual whom a medical cannabis patient cardholder
1567	or a medical cannabis guardian cardholder designates as a designated caregiver; and
1568	(b) is connected to the electronic verification system.
1569	(30) "Medical cannabis courier" means the same as that term is defined in Section
1570	4-41a-102.
1571	(31)(a) "Medical cannabis device" means a device that an individual uses to ingest or
1572	inhale [eannabis in a medicinal dosage form or a cannabis product in a medicinal
1573	dosage form] medical cannabis.
1574	(b) "Medical cannabis device" does not include a device that:
1575	(i) facilitates cannabis combustion; or
1576	(ii) an individual uses to ingest substances other than cannabis.
1577	(32) "Medical cannabis guardian card" means an electronic document that a cardholder may
1578	print or store on an electronic device or a physical card or document that:
1579	(a) the department issues to the parent or legal guardian of a minor with a qualifying
1580	condition; and
1581	(b) is connected to the electronic verification system.
1582	(33) "Medical cannabis ombudsman" means the same as that term is defined in Section
1583	<u>26B-4-248.</u>
1584	[(33)] (34) "Medical cannabis patient card" means an electronic document that a cardholder
1585	may print or store on an electronic device or a physical card or document that:
1586	(a) the department issues to an individual with a qualifying condition; and
1587	(b) is connected to the electronic verification system.
1588	[(34)] (35) "Medical cannabis pharmacy" means a person that:
1589	(a)(i) acquires or intends to acquire medical cannabis [or a cannabis product in a
1590	medicinal dosage form ]from a cannabis processing facility or another medical
1591	cannabis pharmacy or a medical cannabis device; or
1592	(ii) possesses medical cannabis or a medical cannabis device; and

1593	(b) sells or intends to sell medical cannabis or a medical cannabis device to a medical
1594	cannabis cardholder.
1595	[(35)] (36) "Medical cannabis pharmacy agent" means an individual who holds a valid
1596	medical cannabis pharmacy agent registration card issued by the department.
1597	[(36)] (37) "Medical cannabis pharmacy agent registration card" means a registration card
1598	issued by the department that authorizes an individual to act as a medical cannabis
1599	pharmacy agent.
1600	[(37)] (38) "Medical cannabis shipment" means the same as that term is defined in Section
1601	4-41a-102.
1602	[(38)] (39) "Medical cannabis treatment" means [eannabis in a medicinal dosage form, a
1603	cannabis product in a medicinal dosage form, or] medical cannabis or a medical cannabis
1604	device.
1605	[(39)] (40)(a) "Medicinal dosage form" means:
1606	(i) for processed medical cannabis, the following with a specific and consistent
1607	cannabinoid content:
1608	(A) a tablet;
1609	(B) a capsule;
1610	(C) a concentrated liquid or viscous oil;
1611	(D) a liquid suspension that does not exceed 30 milliliters;
1612	(E) a topical preparation;
1613	(F) a transdermal preparation;
1614	(G) a sublingual preparation;
1615	(H) a gelatinous cube, gelatinous rectangular cuboid, or lozenge in a cube or
1616	rectangular cuboid shape;
1617	(I) a resin or wax;
1618	(J) an aerosol;
1619	(K) a suppository preparation; or
1620	(L) a soft or hard confection that is a uniform rectangular cuboid or uniform
1621	spherical shape, is homogeneous in color and texture, and each piece is a single
1622	serving; or
1623	(ii) for unprocessed cannabis flower, a container described in Section 4-41a-602 that:
1624	(A) contains cannabis flower in a quantity that varies by no more than 10% from
1625	the stated weight at the time of packaging;
1626	(B) at any time the medical cannabis cardholder transports or possesses the

1627	container in public, is contained within an opaque bag or box that the medical
1628	cannabis pharmacy provides; and
1629	(C) is labeled with the container's content and weight, the date of purchase, the
1630	legal use termination date, and a barcode that provides information connected
1631	to an inventory control system.
1632	(b) "Medicinal dosage form" includes a portion of unprocessed cannabis flower that:
1633	(i) the medical cannabis cardholder has recently removed from the container
1634	described in Subsection [(39)(a)(ii)] (40)(a)(ii) for use; and
1635	(ii) does not exceed the quantity described in Subsection [(39)(a)(ii)] (40)(a)(ii).
1636	(c) "Medicinal dosage form" does not include:
1637	(i) any unprocessed cannabis flower outside of the container described in Subsection
1638	(39)(a)(ii)] $(40)(a)(ii)$ , except as provided in Subsection $[(39)(b)]$ $(40)(b)$ ;
1639	(ii) any unprocessed cannabis flower in a container described in Subsection [
1640	$\frac{(39)(a)(ii)}{(40)(a)(ii)}$ after the legal use termination date;
1641	(iii) a process of vaporizing and inhaling concentrated cannabis by placing the
1642	cannabis on a nail or other metal object that is heated by a flame, including a
1643	blowtorch;
1644	(iv) a liquid suspension that is branded as a beverage;
1645	(v) a substance described in Subsection $[(39)(a)(i)]$ $(40)(a)(i)$ or (ii) if the substance is
1646	not measured in grams, milligrams, or milliliters; or
1647	(vi) a substance that contains or is covered to any degree with chocolate.
1648	[(40)] (41) "Nonresident patient" means an individual who:
1649	(a) is not a resident of Utah or has been a resident of Utah for less than 45 days;
1650	(b) has a currently valid medical cannabis card or the equivalent of a medical cannabis
1651	card under the laws of another state, district, territory, commonwealth, or insular
1652	possession of the United States; and
1653	(c) has been diagnosed with a qualifying condition as described in Section 26B-4-203.
1654	[(41)] (42) "Patient product information insert" means a single page document or webpage
1655	that contains information about a medical cannabis product regarding:
1656	(a) how to use the product;
1657	(b) common side effects;
1658	(c) serious side effects;
1659	(d) dosage;
1660	(e) contraindications:

1661	(f) safe storage;
1662	(g) information on when a product should not be used; and
1663	(h) other information the department deems appropriate in consultation with the
1664	cannabis processing facility that created the product.
1665	(43) "Pharmacy medical provider" means the medical provider required to be on site at a
1666	medical cannabis pharmacy under Section 26B-4-219.
1667	[(42)] (44) "Provisional patient card" means a card that:
1668	(a) the department issues to a minor with a qualifying condition for whom:
1669	(i) a recommending medical provider has recommended a medical cannabis
1670	treatment; and
1671	(ii) the department issues a medical cannabis guardian card to the minor's parent or
1672	legal guardian; and
1673	(b) is connected to the electronic verification system.
1674	[(43)] (45) "Qualified medical provider" means an individual:
1675	(a) who meets the recommending qualifications; and
1676	(b) whom the department registers to recommend treatment with cannabis in a medicinal
1677	dosage form under Section 26B-4-204.
1678	[(44)] (46) "Qualified Patient Enterprise Fund" means the enterprise fund created in Section
1679	26B-1-310.
1680	[(45)] (47) "Qualifying condition" means a condition described in Section 26B-4-203.
1681	[(46)] (48) "Recommend" or "recommendation" means, for a recommending medical
1682	provider, the act of suggesting the use of medical cannabis treatment, which:
1683	(a) certifies the patient's eligibility for a medical cannabis card; and
1684	(b) may include, at the recommending medical provider's discretion, directions of use,
1685	with or without dosing guidelines.
1686	[(47)] (49) "Recommending medical provider" means a qualified medical provider or a
1687	limited medical provider.
1688	[(48)] (50) "Recommending qualifications" means that an individual:
1689	(a)(i) has the authority to write a prescription;
1690	(ii) is licensed to prescribe a controlled substance under Title 58, Chapter 37, Utah
1691	Controlled Substances Act; and
1692	(iii) possesses the authority, in accordance with the individual's scope of practice, to
1693	prescribe a Schedule II controlled substance; and
1694	(b) is licensed as:

1695	(i) a podiatrist under Title 58, Chapter 5a, Podiatric Physician Licensing Act;
1696	(ii) an advanced practice registered nurse under Title 58, Chapter 31b, Nurse Practice
1697	Act;
1698	(iii) a physician under Title 58, Chapter 67, Utah Medical Practice Act, or Title 58,
1699	Chapter 68, Utah Osteopathic Medical Practice Act; or
1700	(iv) a physician assistant under Title 58, Chapter 70a, Utah Physician Assistant Act.
1701	[(49) "State central patient portal" means the website the department creates, in accordance
1702	with Section 26B-4-236, to facilitate patient safety, education, and an electronic medical
1703	cannabis order.]
1704	[(50)] (51) "State electronic verification system" means the system described in Section
1705	26B-4-202.
1706	[(51)] (52) "Targeted marketing" means the promotion by a qualified medical provider,
1707	medical clinic, or medical office that employs a qualified medical provider of a medical
1708	cannabis recommendation service using any of the following methods:
1709	(a) electronic communication to an individual who is at least 21 years old and has
1710	requested to receive promotional information;
1711	(b) an in-person marketing event that is held in an area where only an individual who is
1712	at least 21 years old may access the event;
1713	(c) other marketing material that is physically or digitally displayed in the office of the
1714	medical clinic or office that employs a qualified medical provider; or
1715	(d) a leaflet that a qualified medical provider, medical clinic, or medical office that
1716	employs a qualified medical provider shares with an individual who is at least 21
1717	years old.
1718	[(52)] (53) "Tetrahydrocannabinol" or "THC" means a substance derived from cannabis or a
1719	synthetic equivalent as described in Subsection 58-37-4(2)(a)(iii)(AA).
1720	[(53)] (54) "THC analog" means the same as that term is defined in Section 4-41-102.
1721	Section 22. Section <b>26B-4-202</b> is amended to read:
1722	26B-4-202 . Electronic verification system.
1723	(1) The Department of Agriculture and Food, the department, the Department of Public
1724	Safety, and the Division of Technology Services shall:
1725	(a) enter into a memorandum of understanding in order to determine the function and
1726	operation of the state electronic verification system in accordance with Subsection (2);
1727	(b) coordinate with the Division of Purchasing, under Title 63G, Chapter 6a, Utah
1728	Procurement Code, to develop a request for proposals for a third-party provider to

1729	develop and maintain the state electronic verification system in coordination with the
1730	Division of Technology Services; and
1731	(c) select a third-party provider who:
1732	(i) meets the requirements contained in the request for proposals issued under
1733	Subsection (1)(b); and
1734	(ii) may not have any commercial or ownership interest in a cannabis production
1735	establishment or a medical cannabis pharmacy.
1736	(2) The Department of Agriculture and Food, the department, the Department of Public
1737	Safety, and the Division of Technology Services shall ensure that the state electronic
1738	verification system described in Subsection (1):
1739	(a) allows an individual to apply for a medical cannabis patient card or, if applicable, a
1740	medical cannabis guardian card, provided that the card may not become active until:
1741	(i) the relevant qualified medical provider completes the associated medical cannabis
1742	recommendation; or
1743	(ii) for a medical cannabis card related to a limited medical provider's
1744	recommendation, the medical cannabis pharmacy completes the recording
1745	described in Subsection (2)(d);
1746	(b) allows an individual to apply to renew a medical cannabis patient card or a medical
1747	cannabis guardian card in accordance with Section 26B-4-213;
1748	(c) allows a qualified medical provider, or an employee described in Subsection (3)
1749	acting on behalf of the qualified medical provider, to:
1750	(i) access dispensing and card status information regarding a patient:
1751	(A) with whom the qualified medical provider has a provider-patient relationship;
1752	and
1753	(B) for whom the qualified medical provider has recommended or is considering
1754	recommending a medical cannabis card;
1755	(ii) electronically recommend treatment with [cannabis in a medicinal dosage form or
1756	a cannabis product in a medicinal dosage form] medical cannabis and optionally
1757	recommend dosing guidelines;
1758	(iii) electronically renew a recommendation to a medical cannabis patient cardholder
1759	or medical cannabis guardian cardholder:
1760	(A) using telehealth services, for the qualified medical provider who originally
1761	recommended a medical cannabis treatment during a face-to-face visit with the
1762	patient; or

1763	(B) during a face-to-face visit with the patient, for a qualified medical provider
1764	who did not originally recommend the medical cannabis treatment during a
1765	face-to-face visit; and
1766	(iv) submit an initial application, renewal application, or application payment on
1767	behalf of an individual applying for any of the following:
1768	(A) a medical cannabis patient card;
1769	(B) a medical cannabis guardian card; or
1770	(C) a medical cannabis caregiver card;
1771	(d) allows a medical cannabis pharmacy medical provider or medical cannabis pharmacy
1772	agent, in accordance with Subsection 4-41a-1101(10)(a), to:
1773	(i) access the electronic verification system to review the history within the system of
1774	a patient with whom the provider or agent is interacting, limited to read-only
1775	access for medical cannabis pharmacy agents unless the medical cannabis
1776	pharmacy's pharmacist in charge authorizes add and edit access;
1777	(ii) record a patient's recommendation from a limited medical provider, including any
1778	directions of use, dosing guidelines, or caregiver indications from the limited
1779	medical provider;
1780	(iii) record a limited medical provider's renewal of the provider's previous
1781	recommendation; and
1782	(iv) submit an initial application, renewal application, or application payment on
1783	behalf of an individual applying for any of the following:
1784	(A) a medical cannabis patient card;
1785	(B) a medical cannabis guardian card; or
1786	(C) a medical cannabis caregiver card;
1787	(e) connects with:
1788	(i) an inventory control system that a medical cannabis pharmacy uses to track in real
1789	time and archive purchases of any [eannabis in a medicinal dosage form, eannabis
1790	product in a medicinal dosage form,] medical cannabis or a medical cannabis
1791	device, including:
1792	(A) the time and date of each purchase;
1793	(B) the quantity and type of [eannabis, eannabis product,] medical cannabis or
1794	medical cannabis device purchased;
1795	(C) any cannabis production establishment, any medical cannabis pharmacy, or
1796	any medical cannabis courier associated with the [cannabis, cannabis product,]

1797	medical cannabis or medical cannabis device; and
1798	(D) the personally identifiable information of the medical cannabis cardholder
1799	who made the purchase; and
1800	(ii) any commercially available inventory control system that a cannabis production
1801	establishment utilizes in accordance with Section 4-41a-103 to use data that the
1802	Department of Agriculture and Food requires by rule, in accordance with Title
1803	63G, Chapter 3, Utah Administrative Rulemaking Act, from the inventory
1804	tracking system that a licensee uses to track and confirm compliance;
1805	(f) provides access to:
1806	(i) the department to the extent necessary to carry out the department's functions and
1807	responsibilities under this part;
1808	(ii) the Department of Agriculture and Food to the extent necessary to carry out the
1809	functions and responsibilities of the Department of Agriculture and Food under
1810	Title 4, Chapter 41a, Cannabis Production Establishments and Pharmacies; and
1811	(iii) the Division of Professional Licensing to the extent necessary to carry out the
1812	functions and responsibilities related to the participation of the following in the
1813	recommendation and dispensing of medical cannabis:
1814	(A) a podiatrist licensed under Title 58, Chapter 5a, Podiatric Physician Licensing
1815	Act;
1816	(B) a pharmacist licensed under Title 58, Chapter 17b, Pharmacy Practice Act;
1817	(C) an advanced practice registered nurse licensed under Title 58, Chapter 31b,
1818	Nurse Practice Act;
1819	(D) a physician licensed under Title 58, Chapter 67, Utah Medical Practice Act, or
1820	Title 58, Chapter 68, Utah Osteopathic Medical Practice Act; or
1821	(E) a physician assistant licensed under Title 58, Chapter 70a, Utah Physician
1822	Assistant Act;
1823	[(g) provides access to and interaction with the state central patient portal;]
1824	[(h)] (g) communicates dispensing information from a record that a medical cannabis
1825	pharmacy submits to the state electronic verification system under Subsection
1826	4-41a-1102(3)(a)(ii) to the controlled substance database;
1827	[(i)] (h) provides access to state or local law enforcement only to verify the validity of an
1828	individual's medical cannabis card for the administration of criminal justice and
1829	through a database used by law enforcement; and
1830	(i) creates a record each time a person accesses the system that identifies the person

1831	who accesses the system and the individual whose records the person accesses.
1832	(3)(a) An employee of a qualified medical provider may access the electronic
1833	verification system for a purpose described in Subsection (2)(c) on behalf of the
1834	qualified medical provider if:
1835	(i) the qualified medical provider has designated the employee as an individual
1836	authorized to access the electronic verification system on behalf of the qualified
1837	medical provider;
1838	(ii) the qualified medical provider provides written notice to the department of the
1839	employee's identity and the designation described in Subsection (3)(a)(i); and
1840	(iii) the department grants to the employee access to the electronic verification
1841	system.
1842	(b) An employee of a business that employs a qualified medical provider may access the
1843	electronic verification system for a purpose described in Subsection (2)(c) on behalf
1844	of the qualified medical provider if:
1845	(i) the qualified medical provider has designated the employee as an individual
1846	authorized to access the electronic verification system on behalf of the qualified
1847	medical provider;
1848	(ii) the qualified medical provider and the employing business jointly provide written
1849	notice to the department of the employee's identity and the designation described
1850	in Subsection (3)(b)(i); and
1851	(iii) the department grants to the employee access to the electronic verification
1852	system.
1853	(4)(a) As used in this Subsection (4), "prescribing provider" means:
1854	(i) a podiatrist licensed under Title 58, Chapter 5a, Podiatric Physician Licensing Act
1855	(ii) an advanced practice registered nurse licensed under Title 58, Chapter 31b, Nurse
1856	Practice Act;
1857	(iii) a physician licensed under Title 58, Chapter 67, Utah Medical Practice Act, or
1858	Title 58, Chapter 68, Utah Osteopathic Medical Practice Act; or
1859	(iv) a physician assistant licensed under Title 58, Chapter 70a, Utah Physician
1860	Assistant Act.
1861	(b) A prescribing provider may access information in the electronic verification system
1862	regarding a patient the prescribing provider treats.
1863	(5) The department may release limited data that the system collects for the purpose of:
1864	(a) conducting medical and other department approved research:

1865	(b) providing the report required by Section 26B-4-222; and
1866	(c) other official department purposes.
1867	(6) The department shall make rules, in accordance with Title 63G, Chapter 3, Utah
1868	Administrative Rulemaking Act, to establish:
1869	(a) the limitations on access to the data in the state electronic verification system as
1870	described in this section; and
1871	(b) standards and procedures to ensure accurate identification of an individual requesting
1872	information or receiving information in this section.
1873	(7) Any person who negligently or recklessly releases any information in the state
1874	electronic verification system in violation of this section is guilty of a class C
1875	misdemeanor.
1876	(8) Any person who obtains or attempts to obtain information from the state electronic
1877	verification system by misrepresentation or fraud is guilty of a third degree felony.
1878	(9)(a) Except as provided in Subsections (9)(c) and (9)(e), a person may not knowingly
1879	and intentionally use, release, publish, or otherwise make available to any other
1880	person information obtained from the state electronic verification system for any
1881	purpose other than a purpose specified in this section.
1882	(b) Each separate violation of this Subsection (9) is:
1883	(i) a third degree felony; and
1884	(ii) subject to a civil penalty not to exceed \$5,000.
1885	(c) A law enforcement officer who uses the database used by law enforcement to access
1886	information in the electronic verification system for a reason that is not the
1887	administration of criminal justice is guilty of a class B misdemeanor.
1888	(d) The department shall determine a civil violation of this Subsection (9) in accordance
1889	with Title 63G, Chapter 4, Administrative Procedures Act.
1890	(e) Civil penalties assessed under this Subsection (9) shall be deposited into the General
1891	Fund.
1892	(f) This Subsection (9) does not prohibit a person who obtains information from the state
1893	electronic verification system under Subsection (2)(a), (c), or (f) from:
1894	(i) including the information in the person's medical chart or file for access by a
1895	person authorized to review the medical chart or file;
1896	(ii) providing the information to a person in accordance with the requirements of the
1897	Health Insurance Portability and Accountability Act of 1996; or
1898	(iii) discussing or sharing that information about the patient with the patient.

1899	Section 23. Section <b>26B-4-214</b> is amended to read:
1900	26B-4-214. Medical cannabis caregiver card Registration Renewal
1901	Revocation.
1902	(1)(a) A cardholder described in Section 26B-4-213 may designate[, through the state
1903	central patient portal,] up to two individuals, or an individual and a facility in
1904	accordance with Subsection (1)(b), to serve as a designated caregiver for the
1905	cardholder.
1906	(b)(i) A cardholder described in Section 26B-4-213 may designate one of the
1907	following types of facilities as one of the caregivers described in Subsection (1)(a):
1908	(A) for a patient or resident, an assisted living facility, as that term is defined in
1909	Section 26B-2-201;
1910	(B) for a patient or resident, a nursing care facility, as that term is defined in
1911	Section 26B-2-201; or
1912	(C) for a patient, a general acute hospital, as that term is defined in Section
1913	26B-2-201.
1914	(ii) A facility may:
1915	(A) assign one or more employees to assist patients with medical cannabis
1916	treatment under the caregiver designation described in this Subsection (1)(b);
1917	and
1918	(B) receive a medical cannabis shipment from a medical cannabis pharmacy or a
1919	medical cannabis courier on behalf of the medical cannabis cardholder within
1920	the facility who designated the facility as a caregiver.
1921	(iii) The department shall make rules to regulate the practice of facilities and facility
1922	employees serving as designated caregivers under this Subsection (1)(b).
1923	(c) A parent or legal guardian described in Subsection 26B-4-213(2)(d), in consultation
1924	with the minor and the minor's qualified medical provider, may designate[, through
1925	the state central patient portal,] up to two individuals to serve as a designated
1926	caregiver for the minor, if the department determines that the parent or legal guardian
1927	is not eligible for a medical cannabis guardian card under Section 26B-4-213.
1928	(d)(i) Upon the entry of a caregiver designation under this Subsection (1) by a patient
1929	with a terminal illness described in Section 26B-4-203, the department shall issue
1930	to the designated caregiver an electronic conditional medical cannabis caregiver
1931	card, in accordance with this Subsection (1)(d).
1932	(ii) A conditional medical cannabis caregiver card is valid for the lesser of:

1933	(A) 60 days; or
1934	(B) the day on which the department completes the department's review and issues
1935	a medical cannabis caregiver card under Subsection (1)(a), denies the patient's
1936	medical cannabis caregiver card application, or revokes the conditional
1937	medical cannabis caregiver card under Section 26B-4-246.
1938	(iii) The department may issue a conditional medical cannabis card to an individual
1939	applying for a medical cannabis patient card for which approval of the
1940	Compassionate Use Board is not required.
1941	(iv) An individual described in Subsection (1)(b)(iii) has the rights, restrictions, and
1942	obligations under law applicable to a holder of the medical cannabis card for
1943	which the individual applies and for which the department issues the conditional
1944	medical cannabis card.
1945	(2) An individual that the department registers as a designated caregiver under this section
1946	and a facility described in Subsection (1)(b):
1947	(a) for an individual designated caregiver, may carry a valid medical cannabis caregiver
1948	card;
1949	(b) in accordance with this part, may purchase, possess, transport, or assist the patient in
1950	the use of [eannabis in a medicinal dosage form, a cannabis product in a medicinal
1951	dosage form,] medical cannabis or a medical cannabis device on behalf of the
1952	designating medical cannabis cardholder;
1953	(c) may not charge a fee to an individual to act as the individual's designated caregiver
1954	or for a service that the designated caregiver provides in relation to the role as a
1955	designated caregiver; and
1956	(d) may accept reimbursement from the designating medical cannabis cardholder for
1957	direct costs the designated caregiver incurs for assisting with the designating
1958	cardholder's medicinal use of cannabis.
1959	(3)(a) The department shall:
1960	(i) within 15 days after the day on which an individual submits an application in
1961	compliance with this section, issue a medical cannabis card to the applicant if the
1962	applicant:
1963	(A) is designated as a caregiver under Subsection (1);
1964	(B) is eligible for a medical cannabis caregiver card under Subsection (4); and
1965	(C) complies with this section; and
1966	(ii) notify the Department of Public Safety of each individual that the department

1967	registers as a designated caregiver.
1968	(b) The department shall ensure that a medical cannabis caregiver card contains the
1969	information described in Subsections (5)(b) and (3)(c)(i).
1970	(c) If a cardholder described in Section 26B-4-213 designates an individual as a
1971	caregiver who already holds a medical cannabis caregiver card, the individual with
1972	the medical cannabis caregiver card:
1973	(i) shall report to the department the information required of applicants under
1974	Subsection (5)(b) regarding the new designation;
1975	(ii) if the individual makes the report described in Subsection (3)(c)(i), is not required
1976	to file an application for another medical cannabis caregiver card;
1977	(iii) may receive an additional medical cannabis caregiver card in relation to each
1978	additional medical cannabis patient who designates the caregiver; and
1979	(iv) is not subject to an additional background check.
1980	(4) An individual is eligible for a medical cannabis caregiver card if the individual:
1981	(a) is at least 21 years old;
1982	(b) is a Utah resident;
1983	(c) pays to the department a fee in an amount that, subject to Subsection 26B-1-310(5),
1984	the department sets in accordance with Section 63J-1-504, plus the cost of the
1985	criminal background check described in Section 26B-4-215; and
1986	(d) signs an acknowledgment stating that the applicant received the information
1987	described in Subsection 26B-4-213(9)[-].
1988	(5) An eligible applicant for a medical cannabis caregiver card shall:
1989	(a) submit an application for a medical cannabis caregiver card to the department
1990	through an electronic application connected to the state electronic verification
1991	system; and
1992	(b) submit the following information in the application described in Subsection (5)(a):
1993	(i) the applicant's name, gender, age, and address;
1994	(ii) the name, gender, age, and address of the cardholder described in Section
1995	26B-4-213 who designated the applicant;
1996	(iii) if a medical cannabis guardian cardholder designated the caregiver, the name,
1997	gender, and age of the minor receiving a medical cannabis treatment in relation to
1998	the medical cannabis guardian cardholder; and
1999	(iv) any additional information that the department requests to assist in matching the
2000	application with the designating medical cannabis patient.

2001 (6) Except as provided in Subsection (6)(b), a medical cannabis caregiver card that the 2002 department issues under this section is valid for the lesser of: 2003 (a) an amount of time that the cardholder described in Section 26B-4-213 who 2004 designated the caregiver determines; or 2005 (b) the amount of time remaining before the card of the cardholder described in Section 2006 26B-4-213 expires. 2007 (7)(a) If a designated caregiver meets the requirements of Subsection (4), the designated 2008 caregiver's medical cannabis caregiver card renews automatically at the time the 2009 cardholder described in Section 26B-4-213 who designated the caregiver: 2010 (i) renews the cardholder's card; and 2011 (ii) renews the caregiver's designation, in accordance with Subsection (7)(b). 2012 (b) The department shall provide a method in the card renewal process to allow a 2013 cardholder described in Section 26B-4-213 who has designated a caregiver to: 2014 (i) signify that the cardholder renews the caregiver's designation; 2015 (ii) remove a caregiver's designation; or 2016 (iii) designate a new caregiver. 2017 (8) The department shall record the issuance or revocation of a medical cannabis card under 2018 this section in the controlled substance database. 2019 Section 24. Section **26B-4-222** is amended to read: 2020 26B-4-222 . Report. 2021 (1) By the November interim meeting each year, the department shall report to the Health 2022 and Human Services Interim Committee on: 2023 (a) the number of applications and renewal applications filed for medical cannabis cards; 2024 (b) the number of qualifying patients and designated caregivers; (c) the nature of the debilitating medical conditions of the qualifying patients: 2025 2026 (d) the age and county of residence of cardholders; 2027 (e) the number of medical cannabis cards revoked: 2028 (f) the number of practitioners providing recommendations for qualifying patients; 2029 (g) the number of license applications and renewal license applications received; 2030 (h) the number of licenses the department has issued in each county; 2031 (i) the number of licenses the department has revoked; 2032 (j) the quantity of medical cannabis shipments that the state central patient portal 2033 facilitates]; 2034 (k) the number of overall purchases of medical cannabis [and medical cannabis products]

2035	from each medical cannabis pharmacy;
2036	(l) the expenses incurred and revenues generated from the medical cannabis program;
2037	and
2038	(m) an analysis of product availability in medical cannabis pharmacies in consultation
2039	with the Department of Agriculture and Food.
2040	(2) The report shall include information provided by the Center for Medical Cannabis
2041	Research described in Section 53B-17-1402.
2042	(3) The department may not include personally identifying information in the report
2043	described in this section.
2044	(4) The department shall report to the working group described in Section 36-12-8.2 as
2045	requested by the working group.
2046	Section 25. Section 26B-4-243 is amended to read:
2047	26B-4-243 . Guidance for treatment with medical cannabis.
2048	The department, in consultation with the Center for Medical Cannabis Research created
2049	in Section 53B-17-1402, shall:
2050	(1) develop evidence-based guidance for treatment with medical cannabis based on the
2051	latest medical research that shall include:
2052	(a) for each qualifying condition, a summary of the latest medical research regarding the
2053	treatment of the qualifying condition with medical cannabis;
2054	(b) risks, contraindications, side effects, and adverse reactions that are associated with
2055	medical cannabis use; and
2056	(c) potential drug interactions between medical cannabis and medications that have been
2057	approved by the United States Food and Drug Administration;[-and]
2058	(2) educate recommending medical providers, pharmacy medical providers, medical
2059	cannabis cardholders, and the public regarding:
2060	(a) the evidence-based guidance for treatment with medical cannabis described in
2061	Subsection (1)(a);
2062	(b) relevant warnings and safety information related to medical cannabis use; and
2063	(c) other topics related to medical cannabis use as determined by the department[-] ; and
2064	(3) develop patient product information inserts for medical cannabis products in
2065	consultation with the cannabis processing facility that created the product and does not
2066	contain proprietary information about the product.
2067	Section 26. Section 26B-4-247 is amended to read:
2068	26R-4-247 Department coordination

2069	The department shall:
2070	(1) provide draft rules made under this chapter to the:
2071	(a) advisory board for the advisory board's review; and
2072	(b) medical cannabis ombudsman;
2073	(2) consult with the advisory board regarding:
2074	(a) patient education; and
2075	(b) fees set by the department that pertain to the medical cannabis program; and
2076	(3) when appropriate, consult with the advisory board regarding issues that arise in the
2077	medical cannabis program.
2078	Section 27. Section 26B-4-248 is enacted to read:
2079	26B-4-248 . Medical cannabis ombudsman Duties Appeals.
2080	(1)(a) There is created a medical cannabis ombudsman within the Office of Ombuds
2081	within the department.
2082	(b) The department shall consult with the Department of Agriculture and Food regarding
2083	the selection of the medical cannabis ombudsman.
2084	(c) The medical cannabis ombudsman or an immediate family member of the medical
2085	cannabis ombudsman may not have an ownership interest in a cannabis production
2086	establishment or medical cannabis pharmacy.
2087	(2) The ombudsman shall:
2088	(a) provide training and information to private citizens, civic groups, governmental
2089	entities, and other interested parties across the state regarding the role and duties of
2090	the ombudsman;
2091	(b) develop a website to provide the information described in Subsection (2)(b) in a form
2092	that is easily accessible;
2093	(c) consult on proposed rules that are created under Title 4, Chapter 41a, Cannabis
2094	Production Establishments and Pharmacies, and Title 26B, Chapter 4, Part 2,
2095	Cannabinoid Research and Medical Cannabis;
2096	(d) cooperate and coordinate with governmental entities and other organizations in the
2097	community in exercising the duties under this section; and
2098	(e) as appropriate, make recommendations to the Department of Agriculture and Food
2099	and the department regarding the creation or modification of rules that the
2100	ombudsman considers necessary to carry out the ombudsman's duties under this
2101	section.
2102	(3)(a) The ombudsman shall:

2103	(i) determine which entities receive licenses:
2104	(A) under Section 4-41a-1005 in consultation with the Department of Agriculture
2105	and Food and in accordance with Section 4-41a-1005; and
2106	(B) described in this Subsection (3); and
2107	(ii) inform the Department of Agriculture and Food of the selections.
2108	(b)(i) Subject to the requirements of this Subsection (3) and the criteria established
2109	for obtaining a medical cannabis pharmacy license under Title 4, Chapter 41a,
2110	Cannabis Production Establishments and Pharmacies, the ombudsman shall:
2111	(A) before January 1, 2026, select one entity to receive a medical cannabis
2112	pharmacy license; and
2113	(B) before January 1, 2027, but not before January 1, 2026, select one entity to
2114	receive a medical cannabis pharmacy license.
2115	(ii) When selecting entities under this Subsection (3), if there is a conflict between
2116	the criteria established for obtaining a medical cannabis pharmacy license under
2117	Title 4, Chapter 41a, Cannabis Production Establishments and Pharmacies, and
2118	this section, this section controls.
2119	(c) For the license described in Subsection (3)(b)(i)(B), the ombudsman may not select
2120	an entity:
2121	(i) that owns any interest in or operates a medical cannabis production establishment;
2122	<u>or</u>
2123	(ii) that is owned, partially or entirely, or operated by a medical cannabis production
2124	establishment.
2125	(d) The ombudsman:
2126	(i) may not select an entity to receive a license under this Subsection (3) if the entity
2127	owns a financial interest in a medical cannabis pharmacy or is owned by an entity
2128	that owns a financial interest in a medical cannabis pharmacy; and
2129	(ii) shall select an entity that will site a medical cannabis pharmacy license issued
2130	under this Subsection (3) in an area:
2131	(A) designated as a medically underserved area as determined by the federal
2132	Health Resources and Services Administration; and
2133	(B) located in a county of the third, fourth, fifth, or sixth class.
2134	(e) A license described in this Subsection (3) may not be transferred to another entity
2135	unless that entity meets the requirements of Subsections (3)(c) and (3)(d) that the
2136	transferring entity met when obtaining the license.

2170

2137	(4)(a) The ombudsman shall contract with a nonprofit entity that provides assistance to
2138	medical cannabis cardholders for purchasing medical cannabis or a medical cannabis
2139	device.
2140	(b) Subject to available funds, the contracted nonprofit entity may provide monthly \$150
2141	vouchers to a medical cannabis pharmacy as part of the program described in this
2142	Subsection (4).
2143	(c) A medical cannabis patient is eligible for the program if the individual is:
2144	(i) an active medical cannabis cardholder patient; and
2145	(ii) enrolled in Medicaid or Medicare.
2146	(d) The ombudsman may make rules to effectuate the program described in this
2147	Subsection (4) in accordance with Title 63G, Chapter 4, Administrative Procedures
2148	Act.
2149	(e) A contracted nonprofit entity shall provide the ombudsman an accounting each
2150	quarter of:
2151	(i) how money was used; and
2152	(ii) other metrics determined relevant by the ombudsman.
2153	(5)(a) The ombudsman shall hear all appeals for administrative action taken under Title
2154	4, Chapter 41a, Cannabis Production Establishments and Pharmacies as an informal
2155	proceeding under Title 63G, Chapter 4, Administrative Procedures Act.
2156	(b) The ombudsman shall create rules for hearing appeals in accordance with Title 63G,
2157	Chapter 3, Utah Administrative Rulemaking Act.
2158	(6) Before August 1, 2026, and each year thereafter, the ombudsman shall provide a report
2159	to the Medical Cannabis Governance Structure Working Group created in Section
2160	<u>36-12-8.2 regarding:</u>
2161	(a) the number of appeals heard under Subsection (5);
2162	(b) the number of patients served under Subsection (4); and
2163	(c) policy recommendations related to the medical cannabis program.
2164	Section 28. Section 63I-2-204 is amended to read:
2165	63I-2-204 . Repeal dates: Title 4.
2166	(1) Section 4-11-117, Beekeeping working group Development of standards, is repealed
2167	May 1, 2025.
2168	(2) Subsection 4-41a-102(6), regarding the Cannabis Research Review Board, is repealed
2169	July 1, [ <del>2026</del> ] <u>2025</u> .

(3) Section 4-46-104, Transition, is repealed July 1, 2024.

- 2171 Section 29. Section **63I-2-226** is amended to read:
- 2172 **63I-2-226** . Repeal dates: Titles 26 through 26B.
- 2173 (1) Section 26B-1-241, Tardive dyskinesia, is repealed July 1, 2024.
- 2174 (2) Section 26B-1-302, National Professional Men's Basketball Team Support of Women
- and Children Issues Restricted Account, is repealed July 1, 2024.
- 2176 (3) Section 26B-1-309, Medicaid Restricted Account, is repealed July 1, 2024.
- 2177 (4) Section 26B-1-313, Cancer Research Restricted Account, is repealed July 1, 2024.
- 2178 (5) Section 26B-1-420, Cannabis Research Review Board, is repealed July 1, [2026] 2025.
- 2179 (6) Subsection 26B-1-421(9)(a), regarding a report to the Cannabis Research Review
- 2180 Board, is repealed July 1, [2026] 2025.
- 2181 (7) Section 26B-1-423, Rural Physician Loan Repayment Program Advisory Committee --
- 2182 Membership -- Compensation -- Duties, is repealed July 1, 2026.
- 2183 (8) Section 26B-2-243, Data collection and reporting requirements concerning incidents of
- abuse, neglect, or exploitation, is repealed July 1, 2027.
- 2185 (9) Section 26B-3-142, Long-acting injectables, is repealed July 1, 2024.
- 2186 (10) Subsection 26B-3-215(5), regarding reporting on coverage for in vitro fertilization and
- genetic testing, is repealed July 1, 2030.
- 2188 (11) Subsection 26B-4-201(5), regarding the Cannabis Research Review Board, is repealed
- 2189 July 1, [<del>2026</del>] 2025.
- 2190 (12) Subsection 26B-4-212(1)(b), regarding the Cannabis Research Review Board, is
- 2191 repealed July 1, [2026] 2025.
- 2192 (13) Section 26B-4-702, Creation of Utah Health Care Workforce Financial Assistance
- 2193 Program, is repealed July 1, 2027.
- 2194 (14) Subsection 26B-4-703(3)(b), regarding per diem and expenses for the Rural Physician
- Loan Repayment Program Advisory Committee, is repealed July 1, 2026.
- 2196 (15) Subsection 26B-4-703(3)(c), regarding expenses for the Rural Physician Loan
- 2197 Repayment Program, is repealed July 1, 2026.
- 2198 (16) Subsection 26B-4-703(6)(b), regarding recommendations from the Rural Physician
- Loan Repayment Program Advisory Committee, is repealed July 1, 2026.
- 2200 (17) Section 26B-5-117, Early childhood mental health support grant program, is repealed
- 2201 January 2, 2025.
- 2202 (18) Section 26B-5-302.5, Study concerning civil commitment and the Utah State Hospital,
- 2203 is repealed July 1, 2025.
- 2204 (19) Section 26B-6-414, Respite care services, is repealed July 1, 2025.

- 2205 (20) Section 26B-7-120, Invisible condition alert program education and outreach, is
- 2206 repealed July 1, 2025.
- Section 30. Section **63I-2-236** is amended to read:
- 2208 **63I-2-236** . Repeal dates: Title 36.
- 2209 (1) Section 36-12-8.2, Medical cannabis governance structure working group, is repealed
- 2210 July 1, [<del>2025</del>] <u>2026</u>.
- 2211 (2) Section 36-29-107.5, Murdered and Missing Indigenous Relatives Task Force --
- 2212 Creation -- Membership -- Quorum -- Compensation -- Staff -- Vacancies -- Duties --
- Interim report, is repealed November 30, 2024.
- 2214 (3) Section 36-29-109, Utah Broadband Center Advisory Commission, is repealed
- 2215 November 30, 2027.
- 2216 (4) Section 36-29-110, Blockchain and Digital Innovation Task Force, is repealed
- 2217 November 30, 2024.
- Section 31. **Repealer.**
- This bill repeals:
- Section 4-41a-801.1, Enforcement for medical cannabis pharmacies and couriers -- Fine
- 2221 **-- Citation.**
- Section 26B-4-236, State central patient portal -- Department duties.
- Section 32. **Effective Date.**
- This bill takes effect on May 7, 2025.