House



LEGISLATIVE ACTION

Senate Comm: RCS 02/20/2019

The Committee on Rules (Brandes) recommended the following: Senate Amendment (with title amendment) Delete everything after the enacting clause and insert: Section 1. Paragraphs (g) and (j) of subsection (1), subsection (4), paragraph (e) of subsection (8), and subsections (14) and (15) of section 381.986, Florida Statutes, are amended to read: 381.986 Medical use of marijuana.-(1) DEFINITIONS.-As used in this section, the term: (a) "Marijuana delivere denive" means on shiret word

(g) "Marijuana delivery device" means an object used,

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12	intended for use, or designed for use in preparing, storing,
13	ingesting, inhaling, or otherwise introducing marijuana into the
14	human body, and which is dispensed from a medical marijuana
15	treatment center for medical use by a qualified patient, except
16	that delivery devices intended for the medical use of marijuana
17	by smoking need not be dispensed from a medical marijuana
18	treatment center in order to qualify as marijuana delivery
19	devices.
20	(j) "Medical use" means the acquisition, possession, use,
21	delivery, transfer, or administration of marijuana authorized by
22	a physician certification. The term does not include:
23	1. Possession, use, or administration of marijuana that was
24	not purchased or acquired from a medical marijuana treatment
25	center.
26	2. Possession, use, or administration of marijuana in a
27	form for smoking, in the form of commercially produced food
28	items other than edibles, or of marijuana seeds or flower,
29	except for flower in a sealed, tamper-proof receptacle for
30	vaping.
31	3. Use or administration of any form or amount of marijuana
32	in a manner that is inconsistent with the qualified physician's
33	directions or physician certification.
34	4. Transfer of marijuana to a person other than the
35	qualified patient for whom it was authorized or the qualified
36	patient's caregiver on behalf of the qualified patient.
37	5. The smoking of marijuana in an enclosed indoor workplace
38	as defined in s. 386.203(5).
39	6.5. Use or administration of marijuana in the following
40	locations:

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41	a. On any form of public transportation, except for low-THC
42	cannabis.
43	b. In any public place, except for low-THC cannabis.
44	c. In a qualified patient's place of employment, except
45	when permitted by his or her employer.
46	d. In a state correctional institution, as defined in s.
47	944.02, or a correctional institution, as defined in s. 944.241.
48	e. On the grounds of a preschool, primary school, or
49	secondary school, except as provided in s. 1006.062.
50	f. In a school bus, a vehicle, an aircraft, or a motorboat,
51	except for low-THC cannabis.
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53	For the purposes of this subparagraph, the exceptions for low-
54	THC cannabis do not include the smoking of low-THC cannabis.
55	(4) PHYSICIAN CERTIFICATION
56	(a) A qualified physician may issue a physician
57	certification only if the qualified physician:
58	1. Conducted a physical examination while physically
59	present in the same room as the patient and a full assessment of
60	the medical history of the patient.
61	2. Diagnosed the patient with at least one qualifying
62	medical condition.
63	3. Determined that the medical use of marijuana would
64	likely outweigh the potential health risks for the patient, and
65	such determination must be documented in the patient's medical
66	record. If a patient is younger than 18 years of age, a second
67	physician must concur with this determination, and such
68	concurrence must be documented in the patient's medical record.
69	4. Determined whether the patient is pregnant and
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70 documented such determination in the patient's medical record. A 71 physician may not issue a physician certification, except for 72 low-THC cannabis, to a patient who is pregnant.

73 5. Reviewed the patient's controlled drug prescription
74 history in the prescription drug monitoring program database
75 established pursuant to s. 893.055.

6. Reviews the medical marijuana use registry and confirmed that the patient does not have an active physician certification from another qualified physician.

7. Registers as the issuer of the physician certification for the named qualified patient on the medical marijuana use registry in an electronic manner determined by the department, and:

a. Enters into the registry the contents of the physician certification, including the patient's qualifying condition and the dosage not to exceed the daily dose amount determined by the department, the amount and forms of marijuana authorized for the patient, and any types of marijuana delivery devices needed by the patient for the medical use of marijuana.

b. Updates the registry within 7 days after any change is
made to the original physician certification to reflect such
change.

92 c. Deactivates the registration of the qualified patient
93 and the patient's caregiver when the physician no longer
94 recommends the medical use of marijuana for the patient.

95 8. Obtains the voluntary and informed written consent of 96 the patient for medical use of marijuana each time the qualified 97 physician issues a physician certification for the patient, 98 which shall be maintained in the patient's medical record. The

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99	patient, or the patient's parent or legal guardian if the
100	patient is a minor, must sign the informed consent acknowledging
101	that the qualified physician has sufficiently explained its
102	content. The qualified physician must use a standardized
103	informed consent form adopted in rule by the Board of Medicine
104	and the Board of Osteopathic Medicine, which must include, at a
105	minimum, information related to:
106	a. The Federal Government's classification of marijuana as
107	a Schedule I controlled substance.
108	b. The approval and oversight status of marijuana by the
109	Food and Drug Administration.
110	c. The current state of research on the efficacy of
111	marijuana to treat the qualifying conditions set forth in this
112	section.
113	d. The potential for addiction.
114	e. The potential effect that marijuana may have on a
115	patient's coordination, motor skills, and cognition, including a
116	warning against operating heavy machinery, operating a motor
117	vehicle, or engaging in activities that require a person to be
118	alert or respond quickly.
119	f. The potential side effects of marijuana use.
120	g. The risks, benefits, and drug interactions of marijuana.
121	h. The risks specifically associated with smoking
122	marijuana.
123	<u>i.</u> h. That the patient's de-identified health information
124	contained in the physician certification and medical marijuana
125	use registry may be used for research purposes.
126	
127	A physician may not certify the medical use of marijuana by
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128 smoking for a patient under 18 years of age unless the patient 129 is diagnosed with a terminal condition, the certifying physician 130 determines that smoking is the most effective means of 131 administering medical marijuana for the patient, and a second 132 physician who is a pediatrician concurs with that determination. Such determination and concurrence must be documented in the 133 134 patient's medical record. 135 (b) If a qualified physician issues a physician 136 certification for a qualified patient diagnosed with a 137 qualifying medical condition pursuant to paragraph (2)(k), the 138 physician must submit the following to the applicable board 139 within 14 days after issuing the physician certification: 140 1. Documentation supporting the qualified physician's 141 opinion that the medical condition is of the same kind or class 142 as the conditions in paragraphs (2)(a) - (j). 143 2. Documentation that establishes the efficacy of marijuana as treatment for the condition. 144 145 3. Documentation supporting the qualified physician's opinion that the benefits of medical use of marijuana would 146 147 likely outweigh the potential health risks for the patient. 148 4. Any other documentation as required by board rule. 149 150 The department must submit such documentation to the Consortium 151 Coalition for Medical Marijuana Clinical Outcomes Research and 152 Education established pursuant to s. 1004.4351. 153 (c) The Board of Medicine and the Board of Osteopathic 154 Medicine shall each, by July 1, 2021, adopt by rule practice 155 standards for the certification of smoking as a route of 156 administration. The department shall provide the Board of

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Medicine and the Board of Osteopathic Medicine information from the medical marijuana use registry as necessary for the adoption of practice standards under this paragraph. Such information may not include a qualified physician's, a qualified patient's, or a caregiver's personal identifying information.

162 (d) (c) A qualified physician may not issue a physician certification for more than three 70-day supply limits of 163 164 marijuana or six 35-day supply limits of marijuana in a form for smoking. The department shall quantify by rule a daily dose 165 166 amount with equivalent dose amounts for each allowable form of 167 marijuana dispensed by a medical marijuana treatment center. The 168 department shall use the daily dose amount to calculate a 70-day 169 supply or a 35-day supply, as appropriate.

1. A qualified physician may request an exception to the daily dose amount limit. The request shall be made electronically on a form adopted by the department in rule and must include, at a minimum:

a. The qualified patient's qualifying medical condition.

b. The dosage and route of administration that was insufficient to provide relief to the qualified patient.

c. A description of how the patient will benefit from an increased amount.

179 d. The minimum daily dose amount of marijuana that would be 180 sufficient for the treatment of the qualified patient's 181 qualifying medical condition.

182 2. A qualified physician must provide the qualified183 patient's records upon the request of the department.

3. The department shall approve or disapprove the request within 14 days after receipt of the complete documentation

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186 required by this paragraph. The request shall be deemed approved 187 if the department fails to act within this time period.

<u>(e)</u> (d) A qualified physician must evaluate an existing qualified patient at least once every 30 weeks before issuing a new physician certification. A physician must:

1. Determine if the patient still meets the requirements to be issued a physician certification under paragraph (a).

2. Identify and document in the qualified patient's medical records whether the qualified patient experienced either of the following related to the medical use of marijuana:

a. An adverse drug interaction with any prescription or nonprescription medication; or

b. A reduction in the use of, or dependence on, other types of controlled substances as defined in s. 893.02.

3. Submit a report with the findings required pursuant to subparagraph 2. to the department. The department shall submit such reports to the <u>Consortium</u> Coalition for Medical Marijuana <u>Clinical Outcomes</u> Research and Education established pursuant to s. 1004.4351.

(f) (c) An active order for low-THC cannabis or medical cannabis issued pursuant to former s. 381.986, Florida Statutes 2016, and registered with the compassionate use registry before June 23, 2017, is deemed a physician certification, and all patients possessing such orders are deemed qualified patients until the department begins issuing medical marijuana use registry identification cards.

212 (g) (f) The department shall monitor physician registration 213 in the medical marijuana use registry and the issuance of 214 physician certifications for practices that could facilitate

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215 unlawful diversion or misuse of marijuana or a marijuana 216 delivery device and shall take disciplinary action as 217 appropriate.

218 (h) (q) The Board of Medicine and the Board of Osteopathic 219 Medicine shall jointly create a physician certification pattern 220 review panel that shall review all physician certifications 221 submitted to the medical marijuana use registry. The panel shall 222 track and report the number of physician certifications and the 223 qualifying medical conditions, dosage, supply amount, and form 224 of marijuana certified. The panel shall report the data both by 225 individual qualified physician and in the aggregate, by county, 226 and statewide. The physician certification pattern review panel 227 shall, beginning January 1, 2018, submit an annual report of its 228 findings and recommendations to the Governor, the President of 229 the Senate, and the Speaker of the House of Representatives.

(i) (h) The department, the Board of Medicine, and the Board of Osteopathic Medicine may adopt rules pursuant to ss. 120.536(1) and 120.54 to implement this subsection.

(8) MEDICAL MARIJUANA TREATMENT CENTERS.-

234 (e) A licensed medical marijuana treatment center shall 235 cultivate, process, transport, and dispense marijuana for 236 medical use. A licensed medical marijuana treatment center may 237 not contract for services directly related to the cultivation, processing, and dispensing of marijuana or marijuana delivery 238 239 devices, except that a medical marijuana treatment center 240 licensed pursuant to subparagraph (a)1. may contract with a 241 single entity for the cultivation, processing, transporting, and 242 dispensing of marijuana and marijuana delivery devices. A licensed medical marijuana treatment center must, at all times, 243

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244 maintain compliance with the criteria demonstrated and 245 representations made in the initial application and the criteria established in this subsection. Upon request, the department may 246 247 grant a medical marijuana treatment center a variance from the 248 representations made in the initial application. Consideration 249 of such a request shall be based upon the individual facts and 250 circumstances surrounding the request. A variance may not be 251 granted unless the requesting medical marijuana treatment center 252 can demonstrate to the department that it has a proposed 253 alternative to the specific representation made in its application which fulfills the same or a similar purpose as the 254 255 specific representation in a way that the department can 256 reasonably determine will not be a lower standard than the 257 specific representation in the application. A variance may not 258 be granted from the requirements in subparagraph 2. and 259 subparagraphs (b)1. and 2.

1. A licensed medical marijuana treatment center may transfer ownership to an individual or entity who meets the requirements of this section. A publicly traded corporation or publicly traded company that meets the requirements of this section is not precluded from ownership of a medical marijuana treatment center. To accommodate a change in ownership:

a. The licensed medical marijuana treatment center shall notify the department in writing at least 60 days before the anticipated date of the change of ownership.

269 b. The individual or entity applying for initial licensure 270 due to a change of ownership must submit an application that 271 must be received by the department at least 60 days before the 272 date of change of ownership.

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273 c. Upon receipt of an application for a license, the 274 department shall examine the application and, within 30 days 275 after receipt, notify the applicant in writing of any apparent 276 errors or omissions and request any additional information 277 required.

d. Requested information omitted from an application for licensure must be filed with the department within 21 days after the department's request for omitted information or the application shall be deemed incomplete and shall be withdrawn from further consideration and the fees shall be forfeited.

Within 30 days after the receipt of a complete application, the department shall approve or deny the application.

2. A medical marijuana treatment center, and any individual or entity who directly or indirectly owns, controls, or holds with power to vote 5 percent or more of the voting shares of a medical marijuana treatment center, may not acquire direct or indirect ownership or control of any voting shares or other form of ownership of any other medical marijuana treatment center.

3. A medical marijuana treatment center may not enter into any form of profit-sharing arrangement with the property owner or lessor of any of its facilities where cultivation, processing, storing, or dispensing of marijuana and marijuana delivery devices occurs.

4. All employees of a medical marijuana treatment center must be 21 years of age or older and have passed a background screening pursuant to subsection (9).

300 5. Each medical marijuana treatment center must adopt and 301 enforce policies and procedures to ensure employees and

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302 volunteers receive training on the legal requirements to 303 dispense marijuana to qualified patients.

6. When growing marijuana, a medical marijuana treatment 304 305 center:

306 a. May use pesticides determined by the department, after 307 consultation with the Department of Agriculture and Consumer Services, to be safely applied to plants intended for human 308 309 consumption, but may not use pesticides designated as 310 restricted-use pesticides pursuant to s. 487.042.

b. Must grow marijuana within an enclosed structure and in a room separate from any other plant.

313 c. Must inspect seeds and growing plants for plant pests that endanger or threaten the horticultural and agricultural interests of the state in accordance with chapter 581 and any 316 rules adopted thereunder.

d. Must perform fumigation or treatment of plants, or remove and destroy infested or infected plants, in accordance with chapter 581 and any rules adopted thereunder.

7. Each medical marijuana treatment center must produce and make available for purchase at least one low-THC cannabis product.

8. Each medical marijuana treatment center must produce and make available for purchase at least one type of pre-rolled marijuana cigarette.

326 9.8. A medical marijuana treatment center that produces 327 edibles must hold a permit to operate as a food establishment 328 pursuant to chapter 500, the Florida Food Safety Act, and must 329 comply with all the requirements for food establishments 330 pursuant to chapter 500 and any rules adopted thereunder.

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331 Edibles may not contain more than 200 milligrams of 332 tetrahydrocannabinol, and a single serving portion of an edible 333 may not exceed 10 milligrams of tetrahydrocannabinol. Edibles 334 may have a potency variance of no greater than 15 percent. 335 Edibles may not be attractive to children; be manufactured in 336 the shape of humans, cartoons, or animals; be manufactured in a 337 form that bears any reasonable resemblance to products available 338 for consumption as commercially available candy; or contain any 339 color additives. To discourage consumption of edibles by children, the department shall determine by rule any shapes, 340 341 forms, and ingredients allowed and prohibited for edibles. 342 Medical marijuana treatment centers may not begin processing or 343 dispensing edibles until after the effective date of the rule. 344 The department shall also adopt sanitation rules providing the 345 standards and requirements for the storage, display, or 346 dispensing of edibles.

10.9. Within 12 months after licensure, a medical marijuana 347 348 treatment center must demonstrate to the department that all of 349 its processing facilities have passed a Food Safety Good 350 Manufacturing Practices, such as Global Food Safety Initiative 351 or equivalent, inspection by a nationally accredited certifying 352 body. A medical marijuana treatment center must immediately stop 353 processing at any facility which fails to pass this inspection until it demonstrates to the department that such facility has 355 met this requirement.

356 11.10. When processing marijuana, a medical marijuana 357 treatment center must:

358 a. Process the marijuana within an enclosed structure and 359 in a room separate from other plants or products.

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b. Comply with department rules when processing marijuana with hydrocarbon solvents or other solvents or gases exhibiting potential toxicity to humans. The department shall determine by rule the requirements for medical marijuana treatment centers to use such solvents or gases exhibiting potential toxicity to humans.

366 c. Comply with federal and state laws and regulations and 367 department rules for solid and liquid wastes. The department 368 shall determine by rule procedures for the storage, handling, 369 transportation, management, and disposal of solid and liquid 370 waste generated during marijuana production and processing. The 371 Department of Environmental Protection shall assist the 372 department in developing such rules.

373 d. Test the processed marijuana using a medical marijuana 374 testing laboratory before it is dispensed. Results must be 375 verified and signed by two medical marijuana treatment center 376 employees. Before dispensing, the medical marijuana treatment 377 center must determine that the test results indicate that low-378 THC cannabis meets the definition of low-THC cannabis, the 379 concentration of tetrahydrocannabinol meets the potency 380 requirements of this section, the labeling of the concentration 381 of tetrahydrocannabinol and cannabidiol is accurate, and all 382 marijuana is safe for human consumption and free from 383 contaminants that are unsafe for human consumption. The 384 department shall determine by rule which contaminants must be 385 tested for and the maximum levels of each contaminant which are 386 safe for human consumption. The Department of Agriculture and 387 Consumer Services shall assist the department in developing the 388 testing requirements for contaminants that are unsafe for human

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389 consumption in edibles. The department shall also determine by 390 rule the procedures for the treatment of marijuana that fails to 391 meet the testing requirements of this section, s. 381.988, or 392 department rule. The department may select a random sample from 393 edibles available for purchase in a dispensing facility which 394 shall be tested by the department to determine that the edible meets the potency requirements of this section, is safe for 395 396 human consumption, and the labeling of the tetrahydrocannabinol 397 and cannabidiol concentration is accurate. A medical marijuana 398 treatment center may not require payment from the department for 399 the sample. A medical marijuana treatment center must recall 400 edibles, including all edibles made from the same batch of 401 marijuana, which fail to meet the potency requirements of this 402 section, which are unsafe for human consumption, or for which 403 the labeling of the tetrahydrocannabinol and cannabidiol 404 concentration is inaccurate. The medical marijuana treatment 405 center must retain records of all testing and samples of each 406 homogenous batch of marijuana for at least 9 months. The medical 407 marijuana treatment center must contract with a marijuana 408 testing laboratory to perform audits on the medical marijuana 409 treatment center's standard operating procedures, testing 410 records, and samples and provide the results to the department 411 to confirm that the marijuana or low-THC cannabis meets the 412 requirements of this section and that the marijuana or low-THC 413 cannabis is safe for human consumption. A medical marijuana 414 treatment center shall reserve two processed samples from each 415 batch and retain such samples for at least 9 months for the 416 purpose of such audits. A medical marijuana treatment center may 417 use a laboratory that has not been certified by the department

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418 under s. 381.988 until such time as at least one laboratory 419 holds the required certification, but in no event later than July 1, 2018. 420 421 e. Package the marijuana in compliance with the United 422 States Poison Prevention Packaging Act of 1970, 15 U.S.C. ss. 423 1471 et seq. 424 f. Package the marijuana in a receptacle that has a firmly 425 affixed and legible label stating the following information: 42.6 (I) The marijuana or low-THC cannabis meets the 427 requirements of sub-subparagraph d. 428 (II) The name of the medical marijuana treatment center 429 from which the marijuana originates. 430 (III) The batch number and harvest number from which the 431 marijuana originates and the date dispensed. 432 (IV) The name of the physician who issued the physician 433 certification. 434 (V) The name of the patient. 435 (VI) The product name, if applicable, and dosage form, 436 including concentration of tetrahydrocannabinol and cannabidiol. 437 The product name may not contain wording commonly associated 438 with products marketed by or to children. 439 (VII) The recommended dose. 440 (VIII) A warning that it is illegal to transfer medical 441 marijuana to another person. 442 (IX) A marijuana universal symbol developed by the 443 department. 444 12.11. The medical marijuana treatment center shall include in each package a patient package insert with information on the 445 specific product dispensed related to: 446

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447	a. Clinical pharmacology.
448	b. Indications and use.
449	c. Dosage and administration.
450	d. Dosage forms and strengths.
451	e. Contraindications.
452	f. Warnings and precautions.
453	g. Adverse reactions.
454	13. In addition to the packaging and labeling requirements
455	in subparagraphs 11. and 12., marijuana in a form for smoking
456	must be packaged in a sealed receptacle with a legible and
457	prominent warning to keep away from children and a warning that
458	states marijuana smoke contains carcinogens and may negatively
459	affect health. Such receptacles for marijuana in a form for
460	smoking must be plain, opaque, and white without depictions of
461	the product or images other than the medical marijuana treatment
462	center's department-approved logo and the marijuana universal
463	symbol.
464	14. Before dispensing a marijuana delivery device, a
465	medical marijuana treatment center must ensure that the
466	marijuana delivery device:
467	a. Has a firmly affixed, legible, and permanent label
468	showing the medical marijuana treatment center's department-
469	approved logo, including each individual marijuana cigarette or
470	wrapping paper.
471	b. Does not incorporate colors, shapes, forms, or designs
472	that are intended to make the marijuana delivery device
473	attractive to children or are likely, by their nature, to be
474	attractive to children. The department shall adopt rules
475	specifying allowable colors, shapes, forms, and designs for

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476 marijuana delivery devices.

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477 15.12. Each edible shall be individually sealed in plain, opaque wrapping marked only with the marijuana universal symbol. 478 479 Where practical, each edible shall be marked with the marijuana 480 universal symbol. In addition to the packaging and labeling 481 requirements in subparagraphs 11. and 12. subparagraphs 10. and 482 11., edible receptacles must be plain, opaque, and white without 483 depictions of the product or images other than the medical marijuana treatment center's department-approved logo and the 484 485 marijuana universal symbol. The receptacle must also include a 486 list all of the edible's ingredients, storage instructions, an 487 expiration date, a legible and prominent warning to keep away 488 from children and pets, and a warning that the edible has not 489 been produced or inspected pursuant to federal food safety laws.

<u>16.13.</u> When dispensing marijuana or a marijuana delivery device, a medical marijuana treatment center:

a. May dispense any active, valid order for low-THC cannabis, medical cannabis and cannabis delivery devices issued pursuant to former s. 381.986, Florida Statutes 2016, which was entered into the medical marijuana use registry before July 1, 2017.

b. May not dispense more than a 70-day supply of marijuana or more than a 35-day supply of marijuana in a form for smoking to a qualified patient or caregiver. <u>A 35-day supply of</u> marijuana in a form for smoking may not exceed four ounces.

501 c. Must have the medical marijuana treatment center's 502 employee who dispenses the marijuana or a marijuana delivery 503 device enter into the medical marijuana use registry his or her 504 name or unique employee identifier.

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505 d. Must verify that the qualified patient and the 506 careqiver, if applicable, each have an active registration in 507 the medical marijuana use registry and an active and valid 508 medical marijuana use registry identification card, the amount 509 and type of marijuana dispensed matches the physician 510 certification in the medical marijuana use registry for that 511 qualified patient, and the physician certification has not 512 already been filled. e. May not dispense marijuana to a qualified patient who is 513 younger than 18 years of age. If the qualified patient is 514 515 younger than 18 years of age, marijuana may only be dispensed 516 only to the qualified patient's caregiver. 517 f. May not dispense or sell any other type of cannabis, 518 alcohol, or illicit drug-related product, including pipes, 519 bongs, or wrapping papers, other than a marijuana delivery 520 device required for the medical use of marijuana and which is 521 specified in a physician certification. 522 g. Must, upon dispensing the marijuana or marijuana 523 delivery device, record in the registry the date, time, 524 quantity, and form of marijuana dispensed; the type of marijuana 525 delivery device dispensed; and the name and medical marijuana 526 use registry identification number of the qualified patient or 527 caregiver to whom the marijuana delivery device was dispensed. 528 h. Must ensure that patient records are not visible to 529 anyone other than the qualified patient, his or her caregiver,

and authorized medical marijuana treatment center employees. (14) EXCEPTIONS TO OTHER LAWS.-

(a) Notwithstanding s. 893.13, s. 893.135, s. 893.147, or
any other provision of law, but subject to the requirements of

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534	this section, a qualified patient and the qualified patient's
535	caregiver may purchase from a medical marijuana treatment center
536	for the patient's medical use a marijuana delivery device and up
537	to the amount of marijuana authorized in the physician
538	certification, but may not possess more than a 70-day supply of
539	marijuana at any given time and all marijuana purchased must
540	remain in its original packaging.
541	(b) Notwithstanding paragraph (a), s. 893.13, s. 893.135,
542	s. 893.147, or any other provision of law, a qualified patient
543	and the qualified patient's caregiver may purchase and possess a
544	marijuana delivery device intended for the medical use of
545	marijuana by smoking from a vendor other than a medical
546	marijuana treatment center.
547	<u>(c)(b) Notwithstanding</u> s. 893.13, s. 893.135, s. 893.147,
548	or any other provision of law, but subject to the requirements
549	of this section, an approved medical marijuana treatment center
550	and its owners, managers, and employees may manufacture,
551	possess, sell, deliver, distribute, dispense, and lawfully
552	dispose of marijuana or a marijuana delivery device as provided
553	in this section, s. 381.988, and by department rule. For the
554	purposes of this subsection, the terms "manufacture,"
555	"possession," "deliver," "distribute," and "dispense" have the
556	same meanings as provided in s. 893.02.
557	<u>(d)(c) Notwithstanding</u> s. 893.13, s. 893.135, s. 893.147,
558	or any other provision of law, but subject to the requirements
559	of this section, a certified marijuana testing laboratory,
560	including an employee of a certified marijuana testing

561 laboratory acting within the scope of his or her employment, may 562 acquire, possess, test, transport, and lawfully dispose of

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563 marijuana as provided in this section, in s. 381.988, and by 564 department rule.

(e) (d) A licensed medical marijuana treatment center and 566 its owners, managers, and employees are not subject to licensure or regulation under chapter 465 or chapter 499 for manufacturing, possessing, selling, delivering, distributing, 569 dispensing, or lawfully disposing of marijuana or a marijuana delivery device, as provided in this section, in s. 381.988, and 571 by department rule.

(f) (e) This subsection does not exempt a person from prosecution for a criminal offense related to impairment or intoxication resulting from the medical use of marijuana or relieve a person from any requirement under law to submit to a breath, blood, urine, or other test to detect the presence of a controlled substance.

(g) (f) Notwithstanding s. 893.13, s. 893.135, s. 893.147, or any other provision of law, but subject to the requirements of this section and pursuant to policies and procedures established pursuant to s. 1006.62(8), school personnel may possess marijuana that is obtained for medical use pursuant to this section by a student who is a qualified patient.

584 (h) (q) Notwithstanding s. 893.13, s. 893.135, s. 893.147, 585 or any other provision of law, but subject to the requirements 586 of this section, a research institute established by a public 587 postsecondary educational institution, such as the H. Lee 588 Moffitt Cancer Center and Research Institute, Inc., established 589 under s. 1004.43, or a state university that has achieved the 590 preeminent state research university designation under s. 1001.7065 may possess, test, transport, and lawfully dispose of 591

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592 marijuana for research purposes as provided by this section. 593 (15) APPLICABILITY.-594 (a) This section does not limit the ability of an employer 595 to establish, continue, or enforce a drug-free workplace program 596 or policy. 597 (b) This section does not require an employer to 598 accommodate the medical use of marijuana in any workplace or any 599 employee working while under the influence of marijuana. 600 (c) This section does not create a cause of action against 601 an employer for wrongful discharge or discrimination. 602 (d) This section does not impair the ability of any party 603 to restrict or limit smoking on his or her private property. 604 (e) This section does not prohibit the medical use of 605 marijuana, or a caregiver assisting with the medical use of 606 marijuana, in a nursing home licensed under part II of chapter 607 400; in a hospice facility licensed under part IV of chapter 608 400; or in an assisted living facility licensed under part I of chapter 429, if the medical use of marijuana is not prohibited 609 610 in the facility's policies. 611 (f) Marijuana, as defined in this section, is not 612 reimbursable under chapter 440. Section 2. Section 1004.4351, Florida Statutes, is amended 613 614 to read: 615 1004.4351 Medical marijuana research and education.-616 (1) SHORT TITLE.-This section shall be known and may be 617 cited as the "Medical Marijuana Research and Education Act." 618 (2) LEGISLATIVE FINDINGS. - The Legislature finds that: 619 (a) The present state of knowledge concerning the use of 620 marijuana to alleviate pain and treat illnesses is limited

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621 because permission to perform clinical studies on marijuana is 622 difficult to obtain, with access to research-grade marijuana so 623 restricted that little or no unbiased studies have been 624 performed.

(b) Under the State Constitution, marijuana is available for the treatment of certain debilitating medical conditions.

(c) Additional clinical studies are needed to ensure that the residents of this state obtain the correct dosing, formulation, route, modality, frequency, quantity, and quality of marijuana for specific illnesses.

(d) An effective medical marijuana research and education program would mobilize the scientific, educational, and medical resources that presently exist in this state to determine the appropriate and best use of marijuana to treat illness.

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638 639 (3) DEFINITIONS.—As used in this section, the term:

(a) "Board" means the Medical Marijuana Research andEducation Board.

(b) <u>"Consortium"</u> <u>"Coalition"</u> means the <u>Consortium</u> Coalition for Medical Marijuana <u>Clinical Outcomes</u> Research and Education.

640 (c) "Marijuana" has the same meaning as provided in s. 29,641 Art. X of the State Constitution.

642 (4) <u>CONSORTIUM</u> COALITION FOR MEDICAL MARIJUANA <u>CLINICAL</u>
 643 <u>OUTCOMES</u> RESEARCH AND EDUCATION. –

(a) There is established within the H. Lee Moffitt Cancer
Center and Research Institute, Inc., the <u>Consortium</u> Coalition
for Medical Marijuana <u>Clinical Outcomes</u> Research <u>consisting of</u>
<u>public and private universities</u> and Education. The purpose of
the <u>consortium</u> coalition is to conduct rigorous scientific
research <u>and</u>, provide education, disseminate <u>such</u> research, and

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650 guide policy for the adoption of a statewide policy on ordering 651 and dosing practices for the medical use of marijuana. The 652 <u>consortium</u> coalition shall be physically located at the H. Lee 653 Moffitt Cancer Center and Research Institute, Inc.

654 (b) The Medical Marijuana Research and Education Board is 655 established to direct the operations of the consortium 656 coalition. The board shall be composed of a chairperson 657 appointed by the H. Lee Moffitt Cancer Center and Research 658 Institute, Inc., a member appointed by the University of 659 Florida, and a member representing each other participating 660 university seven members appointed by the president of the university the chief executive officer of the H. Lee Moffitt 661 662 Cancer Center and Research Institute, Inc. Board members must 663 have experience in a variety of scientific and medical fields, 664 including, but not limited to, oncology, neurology, psychology, 665 pediatrics, nutrition, and addiction. Members shall be appointed 666 to 4-year terms and may be reappointed to serve additional terms. The chair shall be elected by the board from among its 667 668 members to serve a 2-year term. The board shall meet at least 669 semiannually at the call of the chair or, in his or her absence or incapacity, the vice chair. Four members constitute a quorum. 670 671 A majority vote of the members present is required for all 672 actions of the board. The board may prescribe, amend, and repeal 673 a charter governing the manner in which it conducts its 674 business. A board member shall serve without compensation but is 675 entitled to be reimbursed for travel expenses by the consortium 676 coalition or the organization he or she represents in accordance 677 with s. 112.061.

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(c) The $\underline{\text{consortium}}$ $\underline{\text{coalition}}$ shall be administered by a

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679	coalition director, who shall be appointed by the H. Lee Moffitt
680	Cancer Center and Research Institute, Inc and serve at the
681	pleasure of the board. The coalition director shall, subject to
682	the approval of the board:
683	1. Propose a budget for the <u>consortium</u> coalition .
684	2. Foster the collaboration of scientists, researchers, and
685	other appropriate personnel in accordance with the <u>consortium's</u>
686	coalition's charter.
687	3. Engage individuals in public and private university
688	programs relevant to the consortium's work to participate in the
689	consortium.
690	4.3. Identify and prioritize the research to be conducted
691	by the <u>consortium</u> coalition .
692	<u>5.</u> 4. Prepare <u>a plan for medical marijuana research</u> the
693	Medical Marijuana Research and Education Plan for submission to
694	the board.
695	<u>6.</u> 5. Apply for grants to obtain funding for research
696	conducted by the consortium coalition.
697	7.6. Perform other duties as determined by the board.
698	(d) The board shall advise the Board of Governors, the
699	State Surgeon General, the Governor, and the Legislature with
700	respect to medical marijuana research and education in this
701	state. The board shall explore methods of implementing and
702	enforcing medical marijuana laws in relation to cancer control,
703	research, treatment, and education.
704	(d) (e) The board shall annually adopt a plan for medical
705	marijuana research. The plan shall organize a program of
706	research that contributes to the body of scientific knowledge on
707	the effects of the medical use of marijuana and informs both

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708 policy and medical practice related to the treatment of 709 debilitating medical conditions with marijuana. Research shall 710 include tracking clinical outcomes, certification standards, 711 dosing standards, routes of administration, efficacy, and side 712 effects. Research must also include the study of the effects of 713 smoking marijuana to treat debilitating medical conditions. The 714 board must award funds to members of the consortium to perform research consistent with the plan, known as the "Medical 715 716 Marijuana Research and Education Plan," which must be in 717 accordance with state law and coordinate with existing programs 718 in this state. The plan must include recommendations for the 719 coordination and integration of medical, pharmacological, 720 nursing, paramedical, community, and other resources connected 721 with the treatment of debilitating medical conditions; research 722 related to the treatment of such medical conditions; and 723 education.

(e) (f) By February 15 of each year, the board shall issue a report to the Governor, the President of the Senate, and the Speaker of the House of Representatives on research projects, research findings, community outreach initiatives, and future plans for the consortium coalition.

(f) (g) Beginning August 1, 2019 January 15, 2018, and 729 730 quarterly thereafter, the Department of Health shall submit to the board a data set that includes, for each patient registered 7.31 732 in the medical marijuana use registry, the patient's qualifying 733 medical condition and the daily dose amount, routes of 734 administration, and forms of marijuana certified for the 735 patient. The department shall also submit to the board a data 736 set for all patients registered in the medical marijuana use

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737 registry before August 1, 2019. (5) RESPONSIBILITIES OF THE H. LEE MOFFITT CANCER CENTER 738 739 AND RESEARCH INSTITUTE, INC.-The H. Lee Moffitt Cancer Center 740 and Research Institute, Inc., shall allocate staff and provide 741 information and assistance, as the consortium's coalition's 742 budget permits, to assist the board in fulfilling its 743 responsibilities. 744 Section 3. Paragraph (h) of subsection (2) and paragraph (b) of subsection (3) of section 381.987, Florida Statutes, are 745 746 amended to read: 747 381.987 Public records exemption for personal identifying 748 information relating to medical marijuana held by the 749 department.-750 (2) The department shall allow access to the confidential 751 and exempt information in the medical marijuana use registry to: (h) The Consortium Coalition for Medical Marijuana Clinical 752 753 Outcomes Research and Education established in s. 1004.4351(4). 754 (3) The department shall allow access to the confidential 755 and exempt information pertaining to the physician certification 756 for marijuana and the dispensing thereof, whether in the 757 registry or otherwise held by the department, to: 758 (b) The Consortium Coalition for Medical Marijuana Clinical 759 Outcomes Research and Education pursuant to s. 381.986 for the 760 purpose of conducting research regarding the medical use of 761 marijuana. 762 Section 4. The proviso following Specific Appropriation 422 763 in section 3 of chapter 2018-9, Laws of Florida, and the proviso 764 following Specific Appropriation 424 in section 3 of chapter

765 2018-9, Laws of Florida, are repealed and the funds appropriated

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766	by those specific appropriations which were affected by those
767	provisos are released from reserve.
768	Section 5. This act shall take effect upon becoming a law.
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770	=========== T I T L E A M E N D M E N T =================================
771	And the title is amended as follows:
772	Delete everything before the enacting clause
773	and insert:
774	A bill to be entitled
775	An act relating to the medical use of marijuana;
776	amending s. 381.986, F.S.; redefining the term
777	"marijuana delivery device" to eliminate the
778	requirement that such devices must be purchased from a
779	medical marijuana treatment center; redefining the
780	term "medical use" to include the possession, use, or
781	administration of marijuana in a form for smoking;
782	restricting the smoking of marijuana in enclosed
783	indoor workplaces; conforming a provision to changes
784	made by the act; requiring a patient's informed
785	consent form to include the risks specifically
786	associated with smoking marijuana; prohibiting a
787	physician from certifying a patient under 18 years of
788	age to smoke marijuana for medical use unless the
789	patient is diagnosed with a terminal condition and the
790	physician makes a certain determination in concurrence
791	with a second physician who is a pediatrician;
792	conforming a provision to changes made by the act;
793	requiring the Board of Medicine and the Board of
794	Osteopathic Medicine to adopt certain practice

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795 standards by rule; requiring the Department of Health 796 to provide the boards with certain information from 797 the medical marijuana use registry, as necessary; 798 establishing supply limits for physician certifications for marijuana in a form for smoking; 799 requiring each medical marijuana treatment center to 800 801 produce and make available for purchase at least one 802 type of pre-rolled marijuana cigarette; requiring that 803 marijuana in a form for smoking meet certain packaging 804 and labeling requirements; requiring a medical 805 marijuana treatment center to ensure that a marijuana 806 delivery device meets certain packaging and labeling 807 requirements; requiring the department to adopt rules 808 specifying certain packaging and labeling requirements 809 for marijuana delivery devices; prohibiting a medical 810 marijuana treatment center from dispensing more than a 811 specified supply limit of marijuana in a form for 812 smoking; deleting a provision prohibiting a medical 813 marijuana treatment center from dispensing or selling 814 specified products; allowing marijuana delivery 815 devices to be purchased from a vendor other than a 816 medical marijuana treatment center; providing 817 applicability; amending s. 1004.4351, F.S.; renaming 818 the Coalition for Medical Marijuana Research and 819 Education as the Consortium for Medical Marijuana 820 Clinical Outcomes Research; establishing the 821 consortium for a specified purpose; renaming the 822 Medical Marijuana Research and Education Board as the 823 Medical Marijuana Research Board; requiring the board

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824 to direct the operations of the consortium; providing 825 membership of the board; providing for the appointment of a consortium director; providing duties of the 826 827 consortium director; requiring the board to annually 828 adopt a plan for medical marijuana research; requiring 829 the plan to include specified information; providing 830 research requirements for the plan; requiring the 831 board to issue an annual report to the Governor and 8.32 Legislature by a specified date; requiring the 833 department to submit certain data sets to the board; 834 amending s. 381.987, F.S.; conforming provisions to 835 changes made by the act; repealing proviso language in 836 s. 3, ch. 2018-9, Laws of Florida, relating to 837 salaries and benefits positions and other personnel 838 services of the department; providing an effective 839 date.