J1 9lr0651 CF HB 17

By: Senator Zirkin

Introduced and read first time: February 4, 2019

Assigned to: Judicial Proceedings

Committee Report: Favorable with amendments

Senate action: Adopted

Read second time: March 13, 2019

CHAPTER

1 AN ACT concerning

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Natalie M. LaPrade Medical Cannabis Commission – Food Containing Medical Cannabis <u>Edible Cannabis Products</u>

4 FOR the purpose of requiring the Natalie M. LaPrade Medical Cannabis Commission to 5 allow certain dispensaries and dispensary agents to acquire, possess, process, 6 transfer, transport, sell, distribute, or dispense food containing medical cannabis 7 edible cannabis products for use by a qualifying patient or caregiver; requiring the Commission, in consultation with the Maryland Department of Health, to adopt 8 9 certain regulations; requiring the Commission to allow certain processors and 10 processor agents to acquire, possess, process, package, label, transfer, transport, sell, 11 and distribute to a dispensary food containing medical cannabis edible cannabis 12 <u>products</u> for use by a qualifying patient or caregiver; requiring the Commission to 13 allow certain processors and processor agents to transport food containing medical eannabis edible cannabis products to an independent testing laboratory; defining a 14 certain term; making technical corrections; and generally relating to the processing 15 and distribution of food containing medical cannabis edible cannabis products. 16

- 17 BY repealing and reenacting, without amendments,
- 18 Article Health General
- 19 Section 13–3301(a), (e), and (f), 13–3307(a)(1), and 13–3309(a), and 21–101(a) and (i)
- 20 Annotated Code of Maryland
- 21 (2015 Replacement Volume and 2018 Supplement)
- 22 BY repealing and reenacting, with amendments,
 - Article Health General

EXPLANATION: CAPITALS INDICATE MATTER ADDED TO EXISTING LAW.

[Brackets] indicate matter deleted from existing law.

<u>Underlining</u> indicates amendments to bill.

Strike out indicates matter stricken from the bill by amendment or deleted from the law by amendment.



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1 2 3 4	Section 13–3301(e) and (g) through (n), 13–3307(e) through (i), and 13–3309(e) through (h) Annotated Code of Maryland (2015 Replacement Volume and 2018 Supplement)
5 6 7 8 9	BY adding to Article – Health – General Section 13–3301(g), 13–3307(e) and (i), and 13–3309(e) and (i) Annotated Code of Maryland (2015 Replacement Volume and 2018 Supplement)
10 11 12 13 14 15	BY repealing and reenacting, with amendments, Article - Health - General Section 13-3301(g) through (n), 13-3307(e) through (i), and 13-3309(e) through (h) Annotated Code of Maryland (2015 Replacement Volume and 2018 Supplement) SECTION 1. BE IT ENACTED BY THE GENERAL ASSEMBLY OF MARYLAND, That the Laws of Maryland read as follows:
17	Article – Health – General
18	13–3301.
19	(a) In this subtitle the following words have the meanings indicated.
20 21 22 23 24	(e) "Dispensary" means an entity licensed under this subtitle that acquires, possesses, processes, transfers, transports, sells, distributes, dispenses, or administers cannabis, products containing cannabis, related supplies, related products containing cannabis including food EDIBLE CANNABIS PRODUCTS, tinctures, aerosols, oils, or ointments, or educational materials for use by a qualifying patient or caregiver.
25 26	(f) "Dispensary agent" means an owner, a member, an employee, a volunteer, an officer, or a director of a dispensary.
27	(G) "FOOD" HAS THE MEANING STATED IN § 21–101 OF THIS ARTICLE.
28 29 30	(G) (1) "EDIBLE CANNABIS PRODUCT" MEANS A MEDICAL CANNABIS PRODUCT INTENDED FOR HUMAN CONSUMPTION BY ORAL INGESTION, IN WHOLE OR IN PART.
31 32	(2) "EDIBLE CANNABIS PRODUCT" INCLUDES MEDICAL CANNABIS PRODUCTS THAT DISSOLVE OR DISINTEGRATE IN THE MOUTH.

(3) "EDIBLE CANNABIS PRODUCT" DOES NOT INCLUDE ANY:

1 **(I)** MEDICAL CANNABIS CONCENTRATE; OR 2 MEDICAL CANNABIS-INFUSED PRODUCT, INCLUDING AN (II) 3 OIL, A WAX, AN OINTMENT, A SALVE, A TINCTURE, A CAPSULE, A SUPPOSITORY, A 4 DERMAL PATCH, A CARTRIDGE, A CHEWABLE OR DISSOLVABLE GELATINOUS CUBE, OR ANY OTHER PRODUCT CONTAINING MEDICAL CANNABIS CONCENTRATE OR 5 6 USABLE CANNABIS THAT HAS BEEN PROCESSED SO THAT DRIED LEAVES AND FLOWERS ARE INTEGRATED INTO OTHER MATERIAL. 7 "Fund" means the Natalie M. LaPrade Medical Cannabis Commission 8 [(g)] **(H)** 9 Fund established under § 13–3303 of this subtitle. 10 [(h)] (I) "Grower" means an entity licensed under this subtitle that: 11 (1) Cultivates or packages medical cannabis; and 12 Is authorized by the Commission to provide cannabis to a processor, dispensary, or independent testing laboratory. 13 14 "Independent testing laboratory" means a facility, an entity, or a site [(i)] **(J)** that offers or performs tests related to the inspection and testing of cannabis and products 15 containing cannabis. 16 "Medical cannabis grower agent" means an owner, an employee, a 17 [(i)] **(K)** 18 volunteer, an officer, or a director of a grower. [(k)] **(L)** "Processor" means an entity that: 19 20 (1) Transforms medical cannabis into another product or extract; and 21(2)Packages and labels medical cannabis. "Processor agent" means an owner, a member, an employee, a 22[(1)] **(M)** volunteer, an officer, or a director of a processor. 23[(m)](N)24"Qualifying patient" means an individual who: 25Has been provided with a written certification by a certifying provider 26in accordance with a bona fide provider-patient relationship; and 27 If under the age of 18 years, has a caregiver. (2)[(n)] **(O)** "Written certification" means a certification that: 28

Is issued by a certifying provider to a qualifying patient with whom the

provider has a bona fide provider-patient relationship; and

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- 1 (2) Includes a written statement certifying that, in the provider's professional opinion, after having completed an assessment of the patient's medical history and current medical condition, the patient has a condition:
- 4 (i) That meets the inclusion criteria and does not meet the exclusion 5 criteria of the certifying provider's application; and
- 6 (ii) For which the potential benefits of the medical use of cannabis 7 would likely outweigh the health risks for the patient; and
- 8 (3) May include a written statement certifying that, in the provider's professional opinion, a 30-day supply of medical cannabis would be inadequate to meet the medical needs of the qualifying patient.
- 11 13–3307.
- 12 (a) (1) A dispensary shall be licensed by the Commission.
- 13 (E) THE COMMISSION SHALL ALLOW A DISPENSARY LICENSED UNDER THIS
 14 SECTION OR A DISPENSARY AGENT REGISTERED UNDER § 13–3308 OF THIS
 15 SUBTITLE TO ACQUIRE, POSSESS, PROCESS, TRANSFER, TRANSPORT, SELL,
 16 DISTRIBUTE, OR DISPENSE FOOD CONTAINING MEDICAL CANNABIS
 17 CANNABIS PRODUCTS FOR USE BY A QUALIFYING PATIENT OR A CAREGIVER.
- [(e)] (F) A dispensary licensed under this section or a dispensary agent registered under § 13–3308 of this subtitle may not be penalized or arrested under State law for acquiring, possessing, processing, transferring, transporting, selling, distributing, or dispensing MEDICAL cannabis, products containing MEDICAL cannabis, related supplies, or educational materials for use by a qualifying patient or a caregiver.
- [(f)] (G) The Commission shall establish requirements for security and product handling procedures that a dispensary must meet to obtain a license under this section, including a requirement for a product—tracking system.
- 26 **[(g)] (H)** The Commission may inspect a dispensary licensed under this section to ensure compliance with this subtitle.
- 28 (I) THE COMMISSION, IN CONSULTATION WITH THE DEPARTMENT, SHALL
 29 ADOPT REGULATIONS TO REQUIRE A DISPENSARY TO MEET ANY ADDITIONAL
 30 REQUIREMENTS THAT THE COMMISSION DETERMINES IS NECESSARY, INCLUDING
 31 REQUIRING A PERMIT, FOR THE DISPENSING OF EDIBLE CANNABIS PRODUCTS.
- [(h)] (J) The Commission may impose penalties or rescind the license of a dispensary that does not meet the standards for licensure set by the Commission.

- 1 Each dispensary licensed under this section shall submit to the [(i)] (J) **(K)** (1) 2 Commission a quarterly report. 3 (2) The quarterly report shall include: (i) The number of patients served; 4 The county of residence of each patient served; 5 (ii) The medical condition for which medical cannabis was 6 (iii) 7 recommended: 8 (iv) The type and amount of medical cannabis dispensed; and 9 If available, a summary of clinical outcomes, including adverse 10 events and any cases of suspected diversion. 11 The quarterly report may not include any personal information that (3) 12 identifies a patient. 13 13–3309. 14 A processor shall be licensed by the Commission. (a) 15 **(E)** THE COMMISSION SHALL ALLOW A PROCESSOR LICENSED UNDER THIS SECTION OR A PROCESSOR AGENT REGISTERED UNDER § 13-3310 OF THIS SUBTITLE 16 17 TO: 18 **(1)** ACQUIRE, POSSESS, PROCESS, PACKAGE, LABEL, TRANSFER, 19 TRANSPORT, SELL, AND DISTRIBUTE TO A DISPENSARY FOOD CONTAINING MEDICAL 20 CANNABIS EDIBLE CANNABIS PRODUCTS FOR USE BY A QUALIFYING PATIENT OR A 21**CAREGIVER; AND** 22**(2)** TRANSPORT FOOD CONTAINING MEDICAL CANNABIS EDIBLE 23CANNABIS PRODUCTS TO AN INDEPENDENT TESTING LABORATORY. 24A processor licensed under this section or a processor agent registered 25under § 13–3310 of this subtitle may not be penalized or arrested under State law for: 26Acquiring, possessing, processing, packaging, labeling, transferring, (1) transporting, selling, or distributing medical cannabis or products containing medical 27 28cannabis to a dispensary for use by a qualifying patient or a caregiver; or
- 29 (2) Transporting medical cannabis or products containing medical 30 cannabis to an independent testing laboratory.

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October 1, 2019.

1 2 3	[(f)] (G) The Commission shall establish requirements for security and product handling procedures that a processor must meet to obtain a license under this section including a requirement for a product—tracking system.
4 5	[(g)] (H) The Commission may inspect a processor licensed under this section to ensure compliance with this subtitle.
6 7	(I) THE COMMISSION, IN CONSULTATION WITH THE DEPARTMENT, SHALL ADOPT REGULATIONS:
8 9 10	(1) INCLUDING BUT NOT LIMITED TO THE PACKAGING, LABELING MARKETING, AND APPEARANCE OF EDIBLE CANNABIS PRODUCTS, TO ENSURE THE SAFETY OF MINORS; AND
11 12 13	(2) TO REQUIRE A PROCESSOR TO MEET ANY ADDITIONAL REQUIREMENTS THAT THE COMMISSION DETERMINES ARE NECESSARY, INCLUDING REQUIRING A PERMIT, FOR THE PROCESSING OF EDIBLE CANNABIS PRODUCTS.
14 15	[(h)] (J) The Commission may impose penalties or rescind the license of a processor that does not meet the standards for licensure set by the Commission.
16	21–101.
17	(a) In this title the following words have the meanings indicated.
18	(i) "Food" means:
19 20	(1) Any substance that is used as food or drink for human beings or as a component of food or drink for human beings; or
21	(2) Chewing gum or any substance that is used as a component of chewing

SECTION 2. AND BE IT FURTHER ENACTED, That this Act shall take effect