HOUSE OF REPRESENTATIVES
THIRTY-SECOND LEGISLATURE, 2024
STATE OF HAWAII

H.B. NO. 1952 H.D. 1

A BILL FOR AN ACT

RELATING TO MEDICAL CANNABIS.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF HAWAII:

SECTION 1. The legislature finds that amendments to the State's medical cannabis dispensary program law are necessary to facilitate the administration of the medical cannabis dispensary program and resolve matters that have arisen since the passage of Act 309, Session Laws of Hawaii 2022, and Act 108, Session Laws of Hawaii 2023.

The purpose of this Act is to:

(1) Provide that medical cannabis dispensaries may purchase cannabis and manufactured cannabis products from other dispensaries without any showing that such purchase is:

(A) Necessary for a qualifying patient's continuous access; or

(B) For medical, scientific, or other legitimate purposes; and

(2) Require the department of health to adopt rules regarding medical cannabis products within a certain time.

SECTION 2. Section 329D-6, Hawaii Revised Statutes, is amended by amending subsection (r) to read as follows:

"(r) A dispensary may purchase cannabis and manufactured cannabis products from another dispensary. The department [~~may~~] shall authorize a dispensary to purchase cannabis and manufactured cannabis products from another dispensary in a manner prescribed by the department by rules adopted pursuant to section 329D-27; provided that:

[~~(1) The purchasing dispensary establishes to the department's satisfaction that:~~

~~(A) The purchase is necessary to ensure that qualifying patients have continuous access to cannabis for medical use; or~~

~~(B) The cannabis and manufactured cannabis products are for medical, scientific, or other legitimate purposes approved by the State;~~

~~(2)~~] (1) The selling dispensary may transport no more than eight hundred ounces, or other amounts with prior

approval by the department, of cannabis or manufactured cannabis products to the purchasing dispensary within a thirty-day period;

[~~(3)~~] (2) The cannabis and manufactured cannabis products are transported between the dispensaries for medical[~~,~~] sales, scientific[~~,~~] use, or other legitimate purposes approved by the State; and

[~~(4)~~] (3) Nothing in this subsection shall relieve any dispensary of its responsibilities and obligations under this chapter and chapter 329."

SECTION 3. Section 329D-10, Hawaii Revised Statutes, is amended as follows:

1. By amending subsection (a) to read:

"(a) The types of medical cannabis products that a dispensary may [~~be manufactured and distributed~~] manufacture and distribute pursuant to this chapter shall be limited to:

(1) Capsules;

(2) Lozenges;

(3) Pills;

(4) Oils and oil extracts;

(5) Tinctures;

(6) Ointments and skin lotions;

(7) Transdermal patches;

(8) Pre-filled and sealed containers used to aerosolize and deliver cannabis orally or by inhalation, such as an inhaler, nebulizer, or device that provides safe pulmonary administration; provided that:

(A) Containers need not be manufactured by the licensed dispensary but shall be filled with cannabis, cannabis oils, or cannabis extracts manufactured by the licensed dispensary or purchased from another dispensary pursuant to section 329D-6(r); but shall not contain nicotine, tobacco-related products, or any other non-cannabis derived products; and

(B) For devices that provide safe pulmonary administration:

(i) The heating element of the device, if any, shall be made of inert materials such as glass, ceramic, or stainless steel, and not of plastic or rubber;

(ii) The device shall be distributed solely for use with single-use, pre-filled, tamper-resistant, sealed containers that do not contain nicotine or other tobacco products;

(iii) There shall be a temperature control on the device that is regulated to prevent the combustion of cannabis oil; and

(iv) The device need not be manufactured by the licensed dispensary;

(9) Pre-rolled cannabis flower products, as specified by the department;

(10) Edible cannabis products, as specified by the department; and

(11) Other products as specified by the department."

2. By amending subsection (d) to read:

"(d) Any medical cannabis product manufactured and distributed pursuant to this chapter shall be regulated and approved by the department and meet all requirements of rules adopted pursuant to this chapter; provided that the department shall establish requirements for child-resistant packaging and accurate and proper labeling. All rules adopted pursuant to

this section shall be adopted no later than nine months after a product is permitted to be manufactured and distributed pursuant to subsection (a)."

SECTION 4. Statutory material to be repealed is bracketed and stricken. New statutory material is underscored.

SECTION 5. This Act shall take effect on July 1, 3000.

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Report Title:
DOH; Medical Cannabis; Dispensaries; Rules

Description:
Specifically authorizes a medical cannabis dispensary to purchase cannabis and manufactured cannabis products from another dispensary. Requires the Department of Health to adopt rules regarding medical cannabis products within a certain time. Effective 7/1/3000. (HD1)

The summary description of legislation appearing on this page is for informational purposes only and is not legislation or evidence of legislative intent.