

A BILL FOR AN ACT

RELATING TO MEDICAL CANNABIS PRODUCTS.

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

- 1 SECTION 1. Section 329D-1, Hawaii Revised Statutes, is
- 2 amended by amending the definition of "manufactured cannabis
- 3 product" to read as follows:
- 4 ""Manufactured cannabis product" means any capsule,
- 5 lozenge, oil or oil extract, tincture, ointment or skin lotion,
- 6 pill, transdermal patch, or pre-filled and sealed container used
- 7 to aerosolize and deliver cannabis orally, such as an inhaler or
- 8 nebulizer, that has been manufactured using cannabis, or any
- 9 other products as specified by the department pursuant to
- 10 section [329D-10(a)(10).] 329D-10(a)."
- 11 SECTION 2. Section 329D-10, Hawaii Revised Statutes, is
- 12 amended to read as follows:
- 13 "§329D-10 Types of manufactured cannabis products. (a)
- 14 The types of medical cannabis products that may be manufactured
- 15 and distributed pursuant to this chapter shall be limited to:
- 16 (1) Capsules;
- 17 (2) Lozenges;



1	(3)	Pills;		
2	(4)	Oils and oil extracts;		
3	(5)	Tinctures;		
4	(6)	Ointments and skin lotions;		
5	(7)	Transdermal patches;		
6	(8)	Pre-filled and sealed containers used to aerosolize		
7		and deliver cannabis orally, such as with an inhaler		
8		or nebulizer; provided that containers need not be		
9		manufactured by the licensed dispensary but shall be		
10		filled with cannabis, cannabis oils, or cannabis		
11		extracts manufactured by the licensed dispensary;		
12		shall not contain nicotine, tobacco-related products,		
13		or any other non-cannabis derived products; and shall		
14		be designed to be used with devices used to provide		
15	·	safe pulmonary administration of manufactured cannabis		
16		products;		
17	(9)	Devices that provide safe pulmonary administration;		
18		provided that:		
19		(A) The heating element of the device, if any, is		
20		made of inert materials such as glass, ceramic,		

or stainless steel, and not of plastic or rubber;

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1		(B)	The device is distributed solely for use with
2			single-use, pre-filled, tamper-resistant, sealed
3			containers that do not contain nicotine or other
4			tobacco products;
5		(C)	The device is used to aerosolize and deliver
6			cannabis by inhalation, such as an inhaler,
7			medical-grade nebulizer, or other similar medical
8			grade volitization device;
9		(D)	There is a temperature control on the device that
10			is regulated to prevent the combustion of
11			cannabis oil; and
12		(E)	The device need not be manufactured by the
13			licensed dispensary; [and]
14	(10)	Cann	abis seeds;
15	(11)	Cann	abis clones; and
16	[(10)]	(12)	Other products as specified by the department.
17	(b)	As u	sed in this section[, "lozenge"] <u>:</u>
18	"Can	nabis	clone" means a cutting or other specimen of a
19	cannabis	plant	that is genetically identical to the plant from
20	which it	was t	aken and can be replanted or developed to produce
21	a mature	canna	bis plant.

- 1 "Cannabis seed" means the seed of the plant (genus)
- 2 Cannabis.
- 3 "Lozenge" means a small tablet manufactured in a manner to
- 4 allow for the dissolving of its medicinal or therapeutic
- 5 component slowly in the mouth."
- 6 SECTION 3. Statutory material to be repealed is bracketed
- 7 and stricken. New statutory material is underscored.
- 8 SECTION 4. This Act shall take effect upon its approval.

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HB HMS 2020-0093

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Report Title:

Medical Cannabis; Cannabis Seeds; Cannabis Clones; Dispensaries

Description:

Authorizes the manufacture and distribution of cannabis seeds and cannabis clones by medical cannabis dispensary licensees.

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