A BILL FOR AN ACT

RELATING TO MEDICAL CANNABIS PRODUCTS.

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

- 1 SECTION 1. The legislature finds that, for twenty years,
- 2 medical cannabis patients have been allowed to self-cultivate
- 3 cannabis. However, there has been no legal way to acquire
- 4 cannabis seeds or clones, which would allow qualifying patients
- 5 to obtain strains with medicinal properties while minimizing the
- 6 risk of unwanted pests and pathogens from being introduced into
- 7 a homegrown environment. Similarly, qualifying patients and
- 8 primary caregivers who cultivate medical cannabis at a
- 9 registered grow site under the State's medical cannabis registry
- 10 program have no legal means of obtaining safe, laboratory-tested
- 11 genetic material from which to propagate their plants.
- 12 The legislature further finds that state law on the medical
- 13 use of cannabis currently authorizes qualifying patients to
- 14 cultivate up to ten medical cannabis plants at a grow site
- 15 listed on the patient's registry card but does not specify by
- 16 what means propagules, cuttings, or other cannabis genetic
- material necessary to produce these plants may be obtained. 17



- 1 practice, existing law often forces patients or their primary
- 2 caregivers to obtain propagules, cuttings, or other cannabis
- 3 genetic material from the illicit market.
- 4 The legislature also finds that, in a number of other
- 5 states that authorize the medical use of cannabis, propagules
- 6 and cuttings are available for purchase through state-licensed
- 7 dispensaries. These laws ensure that patients who choose to
- 8 cultivate their own cannabis plants have a legal channel from
- 9 which to obtain safe, quality-assured genetic material with
- 10 verified therapeutic properties.
- 11 The State's medical cannabis dispensary system law was
- 12 enacted, in part, to improve qualifying patients' access to safe
- 13 and quality-assured medical cannabis and medical cannabis
- 14 products. However, state law does not explicitly authorize
- 15 licensed dispensaries to distribute cannabis seeds, clones,
- 16 propagules, or cuttings. Accordingly, the purpose of this Act
- 17 is to:
- (1) Authorize the manufacture and distribution of cannabis
- seeds and cannabis clones by medical cannabis
- 20 dispensary licensees;

1	(2)	Include cannabis propagules and cuttings in the
2		definition of the term "cannabis", thereby authorizing
3		dispensaries to distribute medical cannabis propagules
4		and cuttings;

- 5 (3) Authorize only persons who may cultivate medical
 6 cannabis to purchase the propagules and cuttings; and
- 7 (4) Impose quantity limits and quality requirements on the distributed propagules and cuttings.
- 9 SECTION 2. Section 329D-1, Hawaii Revised Statutes, is 10 amended as follows:
- 11 1. By inserting two new definitions to read:
- ""Cutting" means the stem of a cannabis plant that is taken
- or cut off for the purpose of being rooted and grown into a new
- 14 cannabis plant.
- "Propagule" means any part of a cannabis plant that can be
- used to grow a new cannabis plant."
- 17 2. By amending the definition of "cannabis" to read:
- 18 ""Cannabis" shall have the same meaning as in section
- 19 329-121. "Cannabis" includes propagules and cuttings."
- 20 3. By amending the definition of "manufactured cannabis
- 21 product" to read:



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          ""Manufactured cannabis product" means any capsule,
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    lozenge, oil or oil extract, tincture, ointment or skin lotion,
 3
    pill, transdermal patch, or pre-filled and sealed container used
 4
    to aerosolize and deliver cannabis orally, such as an inhaler or
 5
    nebulizer, that has been manufactured using cannabis, or any
 6
    other products as specified by the department pursuant to
 7
    section [<del>329D-10(a)-(11).</del>] 329D-10(a)."
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         SECTION 3. Section 329D-10, Hawaii Revised Statutes, is
 9
    amended to read as follows:
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          "§329D-10 Types of manufactured cannabis products.
                                                                 (a)
11
    The types of medical cannabis products that may be manufactured
12
    and distributed pursuant to this chapter shall be limited to:
13
         (1)
              Capsules;
14
         (2)
              Lozenges;
         (3)
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              Pills;
16
         (4)
              Oils and oil extracts;
17
         (5)
              Tinctures;
18
         (6)
              Ointments and skin lotions;
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              Transdermal patches;
         (7)
20
         (8)
              Pre-filled and sealed containers used to aerosolize
21
               and deliver cannabis orally, such as with an inhaler
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1		or n	ebulizer; provided that containers need not be	
2		manu	factured by the licensed dispensary but shall be	
3		fill	ed with cannabis, cannabis oils, or cannabis	
4		extr	acts manufactured by the licensed dispensary;	
5		shal	l not contain nicotine, tobacco-related products,	
6		or a	ny other non-cannabis derived products; and shall	
7		be d	esigned to be used with devices used to provide	
8		safe	pulmonary administration of manufactured cannabis	
9		prod	ucts;	
10	(9)	Devices that provide safe pulmonary administration;		
11		prov	ided that:	
12		(A)	The heating element of the device, if any, is	
13			made of inert materials such as glass, ceramic,	
14			or stainless steel, and not of plastic or rubber;	
15		(B)	The device is distributed solely for use with	
16			single-use, pre-filled, tamper-resistant, sealed	
17			containers that do not contain nicotine or other	
18			tobacco products;	
19		(C)	The device is used to aerosolize and deliver	

cannabis by inhalation, such as an inhaler,

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1			medical-grade nebulizer, or other similar medical
2			grade volitization device;
3		(D)	There is a temperature control on the device that
4			is regulated to prevent the combustion of
5			cannabis oil; and
6		(E)	The device need not be manufactured by the
7			licensed dispensary;
8	(10)	(Oth	er products, including edible] Edible cannabis
9	1	produ	ucts[$_{ au}$] as specified by the department; [$_{ ext{and}}$]
10	(11)	Canna	abis seeds;
11	(12)	Canna	abis clones; and
12	[(11)]	(13)	Other products as specified by the department.
13	[-(d) -	As-1	used in this section, "lozenge" means a small
14	tablet man	u£ac	tured in a manner to allow for the dissolving of
15	its medici	nal «	or therapeutic component slowly in the mouth.
16	(c)	As u	sed in this section, "edible cannabis products"
17	means manu	facti	ared cannabis products intended for
18	gastrointe	stina	al-administration of any cannabinoid extracted
19	from the ea	annal	ois plant and regulated as manufactured cannabis
20	products a	n d n e	ot as a "drug" or "food" as defined and regulated

- 1 in chapter 328, or as "bottled water" as defined and regulated
- 2 in chapter 328D.
- 3 (d) (b) Any medical cannabis product manufactured
- 4 pursuant to this chapter shall be regulated and approved by the
- 5 department and meet all requirements of rules adopted pursuant
- 6 to this chapter; provided that the department shall establish
- 7 requirements for child-resistant packaging and accurate and
- 8 proper labeling.
- 9 (c) As used in this section:
- "Cannabis clone" means a cutting or other specimen of a
- 11 cannabis plant that is genetically identical to the plant from
- 12 which it was taken and can be replanted or developed to produce
- 13 a mature cannabis plant.
- "Cannabis seed" means the seed of the plant (genus)
- 15 Cannabis.
- "Edible cannabis product" means any manufactured cannabis
- 17 product intended for gastrointestinal administration of any
- 18 cannabinoid extracted from the cannabis plant and regulated as
- 19 manufactured cannabis products and not as a "drug" or "food" as
- 20 defined and regulated in chapter 328, or as "bottled water" as
- 21 defined and regulated in chapter 328D.

1 "Lozenge" means a small tablet manufactured in a manner to 2 allow for the dissolving of its medicinal or therapeutic 3 component slowly in the mouth." 4 SECTION 4. Section 329D-13, Hawaii Revised Statutes, is 5 amended to read as follows: 6 "§329D-13 Dispensing limits. (a) A qualifying patient, 7 primary caregiver, qualifying out-of-state patient, or caregiver 8 of a qualifying out-of-state patient shall be allowed to 9 purchase no more than four ounces of cannabis, not including propagules, cuttings, cannabis clones, and cannabis seeds, 10 11 within a consecutive period of fifteen days, or no more than 12 eight ounces of cannabis, not including propagules, cuttings, 13 cannabis clones, and cannabis seeds, within a consecutive period 14 of thirty days. 15 (b) A qualifying patient or primary caregiver authorized 16 to cultivate cannabis pursuant to sections 329-122 and 17 329-130(a) shall be allowed to purchase no more than a total of five propagules, cuttings, cannabis clones, and cannabis seeds 18 19 within a consecutive period of fifteen days, or no more than a 20 total of ten propagules, cuttings, cannabis clones, and cannabis 21 seeds within a consecutive period of thirty days; provided that:

1	(1)	Propagules, cuttings, cannabis clones, and cannabis			
2		seeds sold by the licensee shall have undergone			
3		laboratory-based testing for residual pesticides and			
4		heavy metals, and shall not be distributed if			
5		pesticides or heavy metals are detected; and			
6	(2)	This subsection shall not apply to a qualifying out-			
7		of-state patient or a caregiver of a qualifying out-			
8		of-state patient.			
9	[-(b)	A qualifying patient, primary caregiver, qualifying			
10	out-of-st	ate patient, or caregiver of a qualifying out of state			
11	patient may purchase cannabis] (c) Purchases under subsections				
12	(a) and (b) may be made from any dispensary location in the				
13	State, subject to the limits and restrictions set forth in				
14	[subsection] subsections (a) [-] and (b).				
15	[(c)	Beginning on January 1, 2018, this section] (d)			
16	Subsections (a) and (c) may apply to qualifying out-of-state				
17	patients	from other states, territories of the United States, or			
18	the Distr	ict of Columbia[$+$] attempting to purchase cannabis, not			
19	including	including propagules, cuttings, cannabis clones, and cannabis			
20	seeds, provided that the patient meets the registration				
21	requirements of section 329-123.5."				

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- 1 SECTION 5. This Act does not affect rights and duties that
- 2 matured, penalties that were incurred, and proceedings that were
- 3 begun before its effective date.
- 4 SECTION 6. Statutory material to be repealed is bracketed
- 5 and stricken. New statutory material is underscored.
- 6 SECTION 7. This Act shall take effect upon its approval.

S.B. NO. 545 S.D. 1

Report Title:

Medical Cannabis; Cannabis Seeds; Cannabis Clones; Dispensaries

Description:

Authorizes the sale and distribution of cannabis propagules, cuttings, cannabis seeds and cannabis clones by medical cannabis dispensary licensees, subject to quantity limits and quality requirements. (SD1)

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