

A BILL FOR AN ACT

RELATING TO MEDICAL MARIJUANA.

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

1	SECTION 1. The legislature finds that in the interest of
2	public health and safety, regulatory oversight is needed
3	regarding the State's medical use of marijuana statutes. While
4	the legislature affirmed the need of qualified patients to
5	utilize this controlled substance for medical purposes, the lac
6	of regulation regarding potency, dosage, quality control, and
7	distribution leaves those same citizens lacking in effective
8	care.
9	The purpose of this Act is to:
10	(1) Further define marijuana for medical use;
11	(2) Create standards for dosage and potency regarding
12	qualifying conditions; and
13	(3) Create licensing and operational regulations for
14	manufacturers and dispensaries for medical marijuana.
15	SECTION 2. Section 329-121, Hawaii Revised Statutes, is
16	amended to read as follows:
17	"[+]PART IX.[+] MEDICAL USE OF MARIJUANA

1	§329-121 Definitions. As used in this part:				
2	"Adequate supply" means an amount of marijuana jointly				
3	possessed between the qualifying patient and the primary				
4	caregiver that is not more than is reasonably necessary to				
5	assure the uninterrupted availability of marijuana for the				
6	purpose of alleviating the symptoms or effects of a qualifying				
7	patient's debilitating medical condition; provided that an				
8	"adequate supply" shall not exceed [seven marijuana plants,				
9	whether immature or mature, and] four ounces of usable marijuana				
10	at any given time.				
11	"Debilitating medical condition" means:				
12	(1) Cancer, glaucoma, positive status for human				
13	immunodeficiency virus, acquired immune deficiency				
14	syndrome, or the treatment of these conditions;				
15	(2) A chronic or debilitating disease or medical condition				
16	or its treatment that produces one or more of the				
17	following:				
18	(A) Cachexia or wasting syndrome;				
19	(B) Severe pain;				
20	(C) Severe nausea;				

1		(D)	Seizures, including those characteristic of
2			epilepsy; or
3		(E)	Severe and persistent muscle spasms, including
4			those characteristic of multiple sclerosis or
5			Crohn's disease; or
6	(3)	Any	other medical condition approved by the department
7		of h	ealth pursuant to administrative rules in response
8		to a	request from a physician or potentially
9		qual	ifying patient.
10	<u>"Dis</u>	pensa	ry" means an entity registered with the department
11	of public	safe	ety and licensed by the department of health to
12	dispense	marij	uana for medical use.
13	<u>"Man</u>	ufact	urer" means an entity registered with the
14	departmen	t of	public safety and licensed by the department of
15	health to	cult	ivate, acquire, manufacture, possess, prepare,
16	transfer,	tran	sport, or supply marijuana for medical use.
17	<u>"Mar</u>	ijuan	a for medical use" or "medical marijuana" means
18	any speci	es of	the genus cannabis plant, or any mixture or
19	preparati	on of	them, including whole plant extracts and resins,
20	and is de	liver	ed in the form of:
21	(1)	Liqu	id, including, but not limited to, oil;

1	(2)	<pre>Pill;</pre>				
2	<u>(3)</u>	Vaporized delivery method with use of liquid or oil				
3		but which does not require the use of dried leaves or				
4		plant form; or				
5	(4)	Any other method, excluding smoking, as approved by				
6		the department of health.				
7	["Ma	rijuana" shall have the same meaning as "marijuana" and				
8	"marijuan	a concentrate" as provided in sections 329-1 and 712-				
9	1240.]					
10	["Me	dical use" means the acquisition, possession,				
11	cultivati	on, use, distribution, or transportation of marijuana				
12	or paraph	ernalia relating to the administration of marijuana to				
13	alleviate the symptoms or effects of a qualifying patient's					
14	debilitat	ing medical condition. For the purposes of "medical				
15	use", the	term distribution is limited to the transfer of				
16	marijuana	and paraphernalia.				
17	"Phy	sician" means a person who is licensed to practice				
18	under cha	pter 453 and is licensed with authority to prescribe				
19	drugs and	is registered under section 329-32. "Physician" does				
20	not inclu	de physician's assistant or advanced practice				

1 registered nurse with prescriptive authority as described in 2 section 453-5.3 or 457-8.6. 3 "Primary caregiver" means a person eighteen years of age or 4 older, other than the qualifying patient and the qualifying 5 patient's physician, who has agreed to undertake responsibility 6 for managing the well-being of the qualifying patient with 7 respect to the medical use of marijuana. In the case of a minor 8 or an adult lacking legal capacity, the primary caregiver shall 9 be a parent, guardian, or person having legal custody. 10 "Qualifying patient" means a person who has been diagnosed 11 by a physician as having a debilitating medical condition. 12 ["Usable marijuana" means the dried leaves and flowers of 13 the plant Cannabis family Moraceae, and any mixture or 14 preparation thereof, that are appropriate for the medical use of 15 marijuana. "Usable marijuana" does not include the seeds, 16 stalks, and roots of the plant.] 17 "Written certification" means the qualifying patient's 18 medical records or a statement signed by a qualifying patient's 19 physician, stating that in the physician's professional opinion, 20 the qualifying patient has a debilitating medical condition and

the potential benefits of the medical use of marijuana would

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- 1 likely outweigh the health risks for the qualifying patient.
- 2 The department of health may require, through its rulemaking
- 3 authority, that all written certifications comply with a
- 4 designated form. "Written certifications" are valid for only
- 5 one year from the time of signing."
- 6 SECTION 3. Chapter 329, Hawaii Revised Statutes, is
- 7 amended by adding 4 new sections to be appropriately designated
- 8 and to read as follows:
- 9 "§329-A Potency and Dosage. The department of health
- 10 shall review and publicly report the medical and scientific
- 11 findings regarding potency requirements, dosage recommendations,
- 12 and chemical compositions of any plant of the genus cannabis
- 13 that could be beneficial for each debilitating medical
- 14 condition. The department shall submit its first report to the
- 15 legislature no later than twenty days before the convening of
- 16 the regular session of 2016. The information shall then be made
- 17 available and distributed to qualifying patients, primary
- 18 caregivers, physicians, dispensaries, and manufacturers by
- 19 January 31, 2016. The information in this report shall be
- 20 updated and distributed annually.

1	<u>§329-</u>	B Manufacturer	Licensing.	<u>(a) In</u>	accordance	<u>with</u>
2	sections 3	29-32 and 329-33	3, a manufac	cturer of	marijuana	<u>for</u>
3	medical us	e must be regist	ered with t	the depar	tment of p	ublic
4	safety. D	ue to the nature	e of marijua	ana as a	controlled	
5	substance,	a manufacturer	must also h	oe licens	sed by the	
6	department	of health. No	more than t	two manuf	acturers s	hall be
7	<u>licensed a</u>	t any one time i	in the State	e, and or	ne manufact	urer must
8	be license	d prior to Decem	mber 31, 201	L5. The	department	shall
9	consider f	or licensing onl	Ly those ent	ities me	eting, at	<u>a</u>
10	minimum, t	he following red	quirements:			
11	(1)	A manufacturer s	shall contra	act with	a laborato	ry for
12		purposes of test	ing its pro	oduct for	content,	
13		contamination, a	and consiste	ency to v	verify that	<u>the</u>
14		marijuana fulfil	ls the guid	delines o	of the pote	ncy and
15		dosage report as	defined by	y section	n 329-A and	
16		distributed by t	he departme	ent of he	ealth. The	cost of
17		laboratory testi	ng shall be	e paid by	the manuf	acturer.
18	(2)	A manufacturer m	nust be able	e to cons	sistently s	upply
19	_	marijuana for me	edical use i	n the ag	proved for	ms as
20		defined in this	section.			

1	(3)	A manufacturer is subject to sections 329-35, 329-36,		
2		329-37, 329-38, and 329-39, and accordingly, must		
3		develop and maintain:		
4		(A) Procedures for internal oversight;		
5		(B) Procedures to ensure accurate record keeping;		
6		(C) Procedures for prescription tracking and product		
7		labeling; and		
8		(D) Procedures for the implementation of appropriate		
9		security measures to deter and prevent the theft		
10		of marijuana for medical use and unauthorized		
11		entrance into areas containing the controlled		
12		substance, including a fully operational security		
13		alarm system, facility access controls, perimeter		
14		intrusion detection systems, and a personnel		
15		identification system.		
16	(4)	A manufacturer shall not share office space with,		
17		refer patients to a physician, or have any financial		
18		relationship with a physician.		
19	(5)	A manufacturer may not employ any person who is under		
20		twenty-one years of age or who has been convicted of a		
21		felony offense under section 329, or part IV of		

1	chapter 712. Each employee must submit a completed
2	criminal history record check consent form and a full
3	set of classifiable fingerprints for submission to the
4	department of health according to the criminal
5	background check requirements of section 846-2.7. The
6	department must conduct a Hawaii criminal history
7	record check and is authorized to exchange the
8	fingerprints with the Federal Bureau of Investigation
9	to obtain the applicant's national criminal history
10	record information.
11	(b) A manufacturer shall operate a single location for
12	cultivation, harvesting, manufacturing, packaging, and
13	processing of marijuana for medical use. The facility must be
14	operational and begin distribution of marijuana for medical use
15	by July 1, 2016.
16	(c) Each manufacturer shall report to the department of
17	health on a monthly basis:
18	(1) The amount, dosages, and chemical composition of
19	marijuana for medical use that the manufacturer
20	distributed; and

1	(2) The tracking number assigned to the marijuana for
2	medical use that the manufacturer distributed.
3	(d) Each manufacturer shall be reviewed annually for
4	license renewal by the department of health to ensure regulatory
5	compliance.
6	(e) A manufacturer shall pay to the department of health
7	fees of \$5,000 for its initial licensing application and \$2,500
8	per year thereafter to retain its license.
9	§329-C Dispensary Licensing. (a) In accordance with
10	sections 329-32 and 329-33, a dispensary of marijuana for
11	medical use must be registered with the department of public
12	safety. Due to the nature of marijuana as a controlled
13	substance, a dispensary must also be specially licensed as a
14	dispenser of medical marijuana by the department of health. No
15	more than three dispensaries shall be licensed per county at any
16	one time, and no more than ten dispensaries licensed at any one
17	time in the State. No less than two dispensaries must be
18	licensed prior to December 31, 2015. The department shall
19	consider for licensing only those entities meeting, at a
20	minimum, the following requirements:

1	<u>(</u>	A dispensary shall have functioned as a pharmacy in
2		accordance with chapter 461, or as a clinic in
3		accordance with section 329-31.5, for no less than
4		five years.
5	(2)	A dispensary shall have in their employ no more than
6		four registered pharmacists to be listed in their
7		license as authorized dispensers. No other employee
8		of the dispensary is permitted to handle, process,
9		dispense, or distribute marijuana for medical use.
10	(3)	A dispensary may not employ a registered pharmacist
11		who is under twenty-one years of age or who has been
12		convicted of a felony offense under section 329, or
13		part IV of chapter 712. Each pharmacist must submit a
14		completed criminal history record check consent form
15		and a full set of classifiable fingerprints for
16		submission to the department of health according to
17		the criminal background check requirements of section
18		846-2.7. The department must conduct a Hawaii
19		criminal history record check and is authorized to
20		exchange the fingerprints with the Federal Bureau of

1		Inve	stigation to obtain the applicant's national
2		crim	inal history record information.
3	(4)	A di	spensary is subject to sections 329-35, 329-36,
4		329-	37, 329-38, 329-39 and, in accordance, must show
5		proo	f of:
6		<u>(A)</u>	Procedures for internal oversight;
7		<u>(B)</u>	Procedures to ensure accurate record keeping;
8		<u>(C)</u>	Procedures for prescription tracking and product
9			labeling; and
10		<u>(D)</u>	Procedures for the implementation of appropriate
11			security measures to deter and prevent the theft
12			of marijuana for medical use and unauthorized
13			entrance into areas containing the controlled
14			substance, including a fully operational security
15			alarm system, facility access controls, perimeter
16			intrusion detection systems, and a personnel
17			identification system.
18	(b)	A di	spensary shall utilize the guidelines of the
19	potency a	nd do	sage report as defined by section 329-A to
20	determine	the	correct treatment for the qualifying patient
21	according	to t	he diagnosis of the physician.

1	(c) A dispensary shall only dispense marijuana for medical
2	use to a registered qualifying patient or their registered
3	primary caregiver.
4	(d) Each dispensary shall report to the department of
5	health on a monthly basis:
6	(1) The amount, dosages, and chemical composition of
7	marijuana for medical use dispensed; and
8	(2) The tracking number assigned to the marijuana for
9	medical use dispensed.
10	(e) Each dispensary shall be reviewed annually for license
11	renewal by the department of health to ensure compliance with
12	all of the above regulations.
13	(f) A dispensary shall pay to the department of health
14	fees of \$2,500 for its initial licensing application and \$1,000
15	per year thereafter to retain its license.
16	§329-D Intentional diversion; penalty. (a)
17	Notwithstanding any law to the contrary, a manufacturer or
18	dispensary, or any agent thereof, who intentionally sells,
19	provides, dispenses, distributes, or otherwise transfers
20	marijuana for medical use to any individual or entity other than
21	a registered qualifying patient or primary caregiver, or



- 1 licensed manufacturer or dispensary shall be guilty of a class C
- 2 felony. The manufacturer or dispensary shall immediately lose
- 3 its license with the department of health.
- 4 (b) Notwithstanding any law to the contrary, any
- 5 qualifying patient, primary caregiver, or connected individual
- 6 who intentionally sells, provides, dispenses, distributes, or
- 7 otherwise transfers marijuana for medical use to any individual
- 8 or entity other than a registered qualifying patient or primary
- 9 caregiver shall be quilty of a class C felony. The connected
- 10 patient's qualifying status will immediately be reviewed for
- 11 possible revocation."
- 12 SECTION 4. The department of public safety and department
- 13 of health shall adopt rules pursuant to chapter 91 to effectuate
- 14 the purpose of this Act.
- 15 SECTION 5. Statutory material to be repealed is bracketed
- 16 and stricken. New statutory material is underscored.
- 17 SECTION 6. This Act shall take effect on July 1, 2015;
- 18 provided that the amendments made to section 329-121, Hawaii
- 19 Revised Statutes, by section 2 of this Act shall take effect
- **20** July 1, 2016.

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INTRODUCED BY:

JAN 2 9 2015

Report Title:

Medical Marijuana; Licensing of Manufacturers and Dispensaries

Description:

Establishes duties and oversight of manufacturers and dispensaries for distribution of marijuana for medical use. Amends the definition of marijuana for medical use and creates requirements for potency and dosages.

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