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JAN 1 6 2020

A BILL FOR AN ACT

RELATING TO INDUSTRIAL HEMP DERIVED PRODUCTS.

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

SECTION 1. The legislature finds that Act 228, Session
 Laws of Hawaii 2016, established the industrial hemp pilot
 program within the department of agriculture and has created the
 promise of a new form of diversified agriculture in Hawaii.
 Since the inception of the hemp pilot program, thirty-six
 industrial hemp farmers have registered with the department and
 are currently cultivating hemp for commercial use.

8 The legislature further finds that Congress passed the 9 Agricultural Improvement Act of 2018, otherwise known as the 10 farm bill, which removed hemp derived extracts, derivatives, and 11 cannabinoids, such as cannabidiol (CBD) as schedule 1 substances 12 in the Controlled Substances Act from hemp plants that contain 13 no more than 0.3 percent tetrahydrocannabinol. This effectively 14 legalized the sale of CBD products from the commercial 15 cultivation of hemp in the United States.

16 The legislature finds that with the passage of the farm17 bill, over sixteen thousand hemp growers have emerged throughout



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1 the United States. Industrial hemp is currently being used 2 nationally in hundreds of different applications including 3 consumer textiles, personal care, industrial components, and 4 dietary supplements containing CBD. The hemp industry across 5 the country has grown rapidly, and hemp-derived products are 6 used by a wide range of consumers.

7 The legislature also finds that, while the United States 8 Department of Agriculture has opened the industrial hemp market, 9 the Food and Drug Administration (FDA) has continued to exercise 10 jurisdiction over the regulation of ingestible and topical hemp 11 products. In 2019, the FDA, in its continuation of evaluating 12 regulatory frameworks for hemp-derived compounds, held a public 13 hearing, and opened a public docket for data gathering. The FDA 14 has also issued non-legally binding public statements arguing 15 that it is illegal to market CBD as a food additive or dietary 16 supplement because it is an active ingredient in a 17 pharmaceutical drug.

18 The legislature additionally finds that, with the existence 19 of competing federal frameworks, several states, such as 20 Florida, Ohio, and Texas have already acted to pass laws or 21 regulations that explicitly allow hemp-derived CBD products to



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be produced and sold and to provide certainty for businesses and 1 2 consumers. While it is expected that the FDA will eventually 3 use its authority to regulate hemp-derived products, the only 4 enforcement action it has taken to date is to send warning letters against improper disease remediation claims made by food 5 6 and supplement companies. In Hawaii, the department of health, 7 has adhered to the FDA public guidance that products containing CBD are adulterated food, beverage, or cosmetic products and 8 9 therefore their sale in Hawaii is prohibited. Despite this 10 suggested prohibition, CBD products continue to be sold across 11 Hawaii, with no regulatory oversight.

12 The legislature finds that, given the time expected for the 13 FDA to act, and the existing confusion among consumers and the 14 industry, it is important that a timely regulatory framework be 15 established around CBD, both to provide consumer safety 16 requirements, and certainty for Hawaii hemp farmers to continue 17 to viably operate their industrial hemp operations in the State. 18 The purpose of this Act is to:

19 (1) Establish a regulatory framework for consumer products
 20 containing CBD that were manufactured legally through
 21 approved government programs and to clarify that these



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1		products shall not be considered adulterated food,		
2		beverage, or cosmetics;		
3	(2)	Prohibit manufacturers of these products from making		
4		health related claims; and		
5	(3)	Require these products to be properly labeled to be		
6		legally allowed for sale in the State.		
7	SECTION 2. Chapter 328, Hawaii Revised Statutes, is			
8	amended by adding a new part to be appropriately designated and			
9	to read as follows:			
10		"PART . INDUSTRIAL HEMP DERIVED PRODUCTS		
11	§328	- Definitions. As used in this part:		
11 12		- Definitions. As used in this part: ustrial hemp" has the same meaning as defined in		
		ustrial hemp" has the same meaning as defined in		
12	"Ind section 1	ustrial hemp" has the same meaning as defined in		
12 13	"Ind section 1 "Ind	ustrial hemp" has the same meaning as defined in 41-31.		
12 13 14	"Ind section 1 "Ind	ustrial hemp" has the same meaning as defined in 41-31. ustrial hemp product" means a finished product g industrial hemp that meets the following conditions:		
12 13 14 15	"Ind section 1 "Ind containin	ustrial hemp" has the same meaning as defined in 41-31. ustrial hemp product" means a finished product g industrial hemp that meets the following conditions:		
12 13 14 15 16	"Ind section 1 "Ind containin	ustrial hemp" has the same meaning as defined in 41-31. ustrial hemp product" means a finished product g industrial hemp that meets the following conditions: Is a cosmetic, food, food additive, dietary		
12 13 14 15 16 17	"Ind section 1 "Ind containin (1)	ustrial hemp" has the same meaning as defined in 41-31. ustrial hemp product" means a finished product g industrial hemp that meets the following conditions: Is a cosmetic, food, food additive, dietary supplement, or herb;		

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1	concentrates, extracts, isolates, resins, or				
2	derivatives; and				
3	(4) Contains no more than 0.3 percent				
4	tetrahydrocannabinol.				
5	"Industrial hemp product" does not include industrial hemp				
6	or a hemp product that is a drug that has been approved as a				
7	drug by the United States Food and Drug Administration.				
8	§328- Manufacture, distribution, or sale of industrial				
9	hemp products. Nothing in this part shall prohibit an entity				
10	licensed pursuant chapter 329D from manufacturing, distributing,				
11	or selling products that contain industrial hemp, cannabinoids,				
12	extracts, or derivatives from industrial hemp grown in				
13	compliance with section 141-32.				
14	§328- Labeling. The label of any package of a food,				
15	beverage, or cosmetic containing cannabidiol derived from				
16	industrial hemp shall include the following statement or a				
17	substantially similar statement:				
18	"CANNABIDIOL USE WHILE PREGNANT OR BREASTFEEDING MAY BE HARMFUL.				
19	KEEP OUT OF REACH OF CHILDREN."				
20	§328- Health-related statements. A manufacturer,				
21	distributor, or seller of an industrial hemp product shall not				



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1 include on the label of the product, or publish or disseminate 2 in advertising or marketing, any health-related statement that 3 is untrue in any particular manner or that tends to create a 4 misleading impression as to the health effects of consuming 5 products containing industrial hemp or cannabinoids, extracts, 6 or derivatives from industrial hemp.

For the purposes of this section, "health-related statement" means a statement related to health and includes a statement of a curative or therapeutic nature that, expressly or impliedly, suggests a relationship between the consumption of industrial hemp or industrial hemp products and health benefits or effects on health.

13 §328- Use in food products. In order for industrial 14 hemp to be used in food products, a manufacturer shall comply 15 with the following:

16 (1) All parts of the hemp plant used in food shall come
17 from a state or country that has an established and
18 approved industrial hemp program that meets all of the
19 federal requirements regarding the lawful and safe
20 cultivation of industrial hemp, and inspects or
21 regulates hemp under a food safety program or



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1		equi	valent criteria to ensure safety for human	
2		consumption;		
3	(2)	The	industrial hemp cultivator or grower shall be in	
4		good	standing and in compliance with the governing	
5		laws of the state or country of origin; and		
6	(3)	A raw hemp product shall not be distributed or sold in		
7		the	State without a certificate of analysis from an	
8		independent testing laboratory that confirms the		
9		following:		
10		(A)	The raw hemp product is the product of a batch of	
11			industrial hemp that was tested by the	
12			independent testing laboratory in accordance with	
13			section 141-32;	
14		(B)	A tested random sample of the batch of industrial	
15			hemp contained a total	
16			delta-9-tetrahydrocannabinol concentration that	
17			did not exceed 0.3 percent on a dry-weight basis;	
18			and	
19		(C)	The tested sample of the batch did not contain	
20			contaminants that are unsafe for human	
21			consumption.	



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1 For the purposes of this section, "manufacture" means to 2 compound, blend, extract, infuse, or otherwise make or prepare a 3 "Manufacture" does not include planting, growing, product. 4 harvesting, drying, curing, grading, or trimming a plant or part 5 of a plant.

6 §328-Hemp products; when adulterated or misbranded. A food, beverage, or cosmetic product shall not be considered 7 8 adulterated pursuant to sections 328-9 and 328-18 or misbranded 9 pursuant to sections 328-10 and 328-19 solely by the inclusion of industrial hemp or cannabinoids, extracts, or derivatives 10 from industrial hemp. The sale of food, beverages, or cosmetics 11 12 that include industrial hemp or cannabinoids, extracts, or derivatives from industrial hemp shall not be restricted or 13 14 prohibited based solely on the inclusion of industrial hemp or 15 cannabinoids, extracts, or derivatives from industrial hemp." 16 SECTION 3. This Act shall take effect on July 1, 2020.

INTRODUCED BY: Resoly & Baken By Request



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Report Title:

Industrial Hemp; Derived Products; Adulterated Product; Labeling

Description:

Establishes a regulatory framework for products containing CBD that were manufactured legally through approved government programs. Clarifies that these products are not considered adulterated food, beverage, or cosmetic products. Prohibits manufacturers from making health related claims. Requires product labeling for the products to be legally allowed in the State.

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