Senator Evan J. Vickers proposes the following substitute bill:

1	CANNABINOID RESEARCH
2	2017 GENERAL SESSION
3	STATE OF UTAH
4	Chief Sponsor: Brad M. Daw
5	Senate Sponsor: Evan J. Vickers
6 7	LONG TITLE
8	General Description:
o 9	-
9 10	This bill enacts provisions related to research of cannabis and cannabinoid products.
	Highlighted Provisions:
11	This bill:
12	 allows a person to possess cannabis, a cannabinoid product, and an expanded
13	cannabinoid product and to distribute the cannabis, a cannabinoid product, or an
14	expanded cannabinoid product to a patient pursuant to an institutional review
15	board-approved study;
16	 allows a person conducting an institutional review board-approved study to import
17	and distribute cannabis, a cannabinoid product, and an expanded cannabinoid
18	product under certain circumstances; and
19	 creates the Cannabinoid Product Board within the Department of Health.
20	Money Appropriated in this Bill:
21	None
22	Other Special Clauses:
23	This bill provides a special effective date.
24	Utah Code Sections Affected:
25	ENACTS:

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	26-59-101, Utah Code Annotated 1953	
	26-59-102 , Utah Code Annotated 1953	
	26-59-103, Utah Code Annotated 1953	
	26-59-201 , Utah Code Annotated 1953	
	26-59-202 , Utah Code Annotated 1953	
	58-37-3.6, Utah Code Annotated 1953	
Bei	it enacted by the Legislature of the state of Utah:	
	Section 1. Section 26-59-101 is enacted to read:	
	CHAPTER 59. CANNABINOID RESEAU	RCH ACT
	<u>26-59-101.</u> Title.	
	This chapter is known as "Cannabinoid Research Act."	
	Section 2. Section 26-59-102 is enacted to read:	
	<u>26-59-102.</u> Definitions.	
	As used in this chapter:	
	(1) "Approved study" means a medical research study:	
	(a) the purpose of which is to investigate the medical benefit	its and risks of cannabinoid
pro	ducts; and	
	(b) that is approved by an IRB.	
	(2) "Board" means the Cannabinoid Product Board created	in Section <u>26-59-201.</u>
	(3) "Cannabinoid product" means the same as that term is d	efined in Section 58-37-3.6.
	(4) "Cannabis" means the same as that term is defined in Se	ection <u>58-37-3.6.</u>
	(5) "Expanded cannabinoid product" means the same as that	t term is defined in Section
<u>58-</u>	<u>37-36.</u>	
	(6) "Institutional review board" or "IRB" means an institution	onal review board that is
regi	istered for human subject research by the United States Departm	ent of Health and Human
Ser	vices.	
	Section 3. Section 26-59-103 is enacted to read:	
	<u>26-59-103.</u> Institutional review board Approved study	r, cannabis, cannabinoid
pro	oduct, or expanded cannabinoid product.	
	(1) A person conducting an approved study may, for the put	poses of the study:

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57	(a) process a cannabinoid product or an expanded cannabinoid product;
58	(b) possess a cannabinoid product or an expanded cannabinoid product; and
59	(c) administer a cannabinoid product, or an expanded cannabinoid product to an
60	individual in accordance with the approved study.
61	(2) A person conducting an approved study may:
62	(a) import cannabis, a cannabinoid product, or an expanded cannabinoid product from
63	another state if:
64	(i) the importation complies with federal law; and
65	(ii) the person uses the cannabis, cannabinoid product, or expanded cannabinoid
66	product in accordance with the approved study; or
67	(b) obtain cannabis, a cannabinoid product, or an expanded cannabinoid product from
68	the National Institute on Drug Abuse.
69	(3) A person conducting an approved study may distribute cannabis, a cannabinoid
70	product, or an expanded cannabinoid product outside the state if:
71	(a) the distribution complies with federal law; and
72	(b) the distribution is for the purposes of, and in accordance with, the approved study.
73	Section 4. Section 26-59-201 is enacted to read:
74	<u>26-59-201.</u> Cannabinoid Product Board.
75	(1) There is created the Cannabinoid Product Board within the department.
76	(2) The department shall appoint, in consultation with a professional association based
77	in the state that represents physicians, seven members to the Cannabinoid Product Board as
78	<u>follows:</u>
79	(a) three individuals who are medical research professionals; and
80	(b) four physicians.
81	(3) The department shall appoint board members under Subsection (2) such that three
82	of the board members are members of the Controlled Substances Advisory Committee created
83	<u>in Section 58-38a-201.</u>
84	(4) (a) Four of the board members appointed under Subsection (2) shall serve an initial
85	term of two years and three of the board members appointed under Subsection (2) shall serve
86	an initial term of four years.
87	(b) Successor board members shall each serve a term of four years.

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88	(5) The department may remove a board member without cause.
89	(6) The board shall nominate a board member to serve as chairperson of the board by a
90	majority vote of the board members.
91	(7) The board shall meet as often as necessary to accomplish the duties assigned to the
92	board under this chapter.
93	(8) Each board member, including the chair, has one vote.
94	(9) (a) A majority of board members constitutes a quorum.
95	(b) A vote of a majority of the quorum at any board meeting is necessary to take action
96	on behalf of the board.
97	(10) A board member may not receive compensation for the member's service on the
98	board, but may, in accordance with rules adopted by the board in accordance with Title 63G,
99	Chapter 3, Utah Administrative Rulemaking Act, receive:
100	(a) per diem at the rate established under Section 63A-3-106; and
101	(b) travel expenses at the rate established under Section 63A-3-107.
102	Section 5. Section 26-59-202 is enacted to read:
103	<u>26-59-202.</u> Cannabinoid Product Board Duties.
104	(1) The board shall evaluate the safety and efficacy of cannabinoid products, including:
105	(a) medical conditions that respond to cannabinoid products;
106	(b) cannabinoid dosage amounts and medical dosage forms; and
107	(c) interaction of cannabinoid products with other treatments.
108	(2) Based on the board's evaluation under Subsection (1), the board shall develop
109	guidelines for a physician recommending treatment with a cannabinoid product that includes a
110	list of medical conditions, if any, that the board determines are appropriate for treatment with
111	cannabinoid medicine.
112	(3) The board shall submit the guidelines described in Subsection (2) to:
113	(a) the director of the Division of Occupational and Professional Licensing; and
114	(b) the Health and Human Services Interim Committee.
115	(4) The board shall report the board's findings before November 1 of each year to the
116	Health and Human Services Interim Committee.
117	Section 6. Section 58-37-3.6 is enacted to read:
118	58-37-3.6. Exemption for possession or distribution of a cannabinoid product

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119	pursuant to an approved study.
120	(1) As used in this section:
121	(a) "Cannabinoid product" means a product intended for human ingestion that:
122	(i) contains an extract or concentrate that is obtained from cannabis;
123	(ii) is prepared in a medicinal dosage form; and
124	(iii) contains at least 10 units of cannabidiol for every one unit of tetrahydrocannabinol.
125	(b) "Cannabis" means any part of the plant cannabis sativa, whether growing or not.
126	(c) "Drug paraphernalia" means the same as that term is defined in Section 58-37a-3.
127	(d) "Expanded cannabinoid product" means a product intended for human ingestion
128	that:
129	(i) contains an extract or concentrate that is obtained from cannabis;
130	(ii) is prepared in a medicinal dosage form; and
131	(iii) contains less than 10 units of cannabidiol for every one unit of
132	tetrahydrocannabinol.
133	(e) "Medicinal dosage form" means:
134	(i) a tablet;
135	(ii) a capsule;
136	(iii) a concentrated oil;
137	(iv) a liquid suspension;
138	(v) a transdermal preparation; or
139	(vi) a sublingual preparation.
140	(f) "Tetrahydrocannabinol" means a substance derived from cannabis that meets the
141	description in Subsection 58-37-4(2)(a)(iii)(AA).
142	(2) Notwithstanding any other provision of this chapter, an individual who possesses or
143	distributes a cannabinoid product or an expanded cannabinoid product is not subject to the
144	penalties described in this title for the possession or distribution of marijuana or
145	tetrahydrocannabinol to the extent that the individual's possession or distribution of the
146	cannabinoid product or expanded cannabinoid product complies with Title 26, Chapter 59,
147	Cannabinoid Research Act.
148	Section 7. Effective date.
149	If approved by two-thirds of all the members elected to each house, this bill takes effect

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- 150 upon approval by the governor, or the day following the constitutional time limit of Utah
- 151 <u>Constitution, Article VII, Section 8, without the governor's signature, or in the case of a veto,</u>
- 152 <u>the date of veto override.</u>