

General Assembly

January Session, 2025

Amendment

LCO No. 8785



Offered by: REP. RUTIGLIANO, 123rd Dist. REP. CANDELORA V., 86th Dist. REP. O'DEA, 125th Dist.

To: House Bill No. **7065**

File No. 730 Cal. No. 465

"AN ACT CONCERNING THE DECRIMINALIZATION OF POSSESSION OF SMALL AMOUNTS OF PSILOCYBIN."

Strike everything after the enacting clause and substitute the
 following in lieu thereof:

"Section 1. (NEW) (*Effective October 1, 2025*) No person may possess
any amount of psilocybin, except as authorized in chapter 420 of the
general statutes, unless and until the federal Food and Drug
Administration has reviewed and approved the use of psilocybin.

Sec. 2. (NEW) (*Effective October 1, 2025*) Until October 1, 2028, there
shall be no direct to consumer advertising in this state for any drug not
reviewed and approved by the federal Food and Drug Administration.

10 Sec. 3. (NEW) (*Effective October 1, 2025*) (a) As used in this section:

(1) "Consumer" means an individual who is physically present in thisstate;

_	HB 7065 Amendment		
13	(2) "Legend drug" has the same meaning as provided in section 20-		
14	571 of the general statutes;		
15	(3) "Person" has the same meaning as provided in section 20-571 of		
16	the general statutes;		
17	(4) "Pharmaceutical manufacturer" has the same meaning as		
18	provided in section 21a-70h of the general statutes;		
10	provided in section 214 / on or the general statutes,		
19	(5) "Pharmaceutical marketing firm" has the same meaning as		
20	provided in section 21a-70h of the general statutes;		
01	() "Dharman actival record tativa" has the same meaning of		
21	(6) "Pharmaceutical representative" has the same meaning as		
22	provided in section 21a-70h of the general statutes; and		
23	(7) "Prescribing practitioner" has the same meaning as provided in		
24	section 20-571 of the general statutes.		
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25	(b) No person engaged in trade or commerce in this state shall		
26	directly advertise any legend drug to a consumer. Nothing in this		
27	subsection shall be construed to prohibit any pharmaceutical		
28 20	manufacturer that is registered with the Department of Consumer		
29 30	Protection as a pharmaceutical marketing firm under section 21a-70i of		
30 31	the general statutes, or any individual who is employed or compensated by such pharmaceutical marketing firm to perform the duties of a		
32	pharmaceutical representative in accordance with the provisions of said		
33	section, from promoting or providing information regarding a legend		
34	drug for human use to a prescribing practitioner.		
51	and for numar use to a presenting practitioner.		
35	(c) The Commissioner of Consumer Protection may adopt		
36	regulations, in accordance with the provisions of chapter 54 of the		
37	general statutes, to implement the provisions of this section.		
20	(d) Any violation of the maximizers of subsection (b) of this section		
38 39	(d) Any violation of the provisions of subsection (b) of this section shall be deemed an unfair or deceptive trade practice under subsection		

38 (d) Any violation of the provisions of subsection (b) of this section
39 shall be deemed an unfair or deceptive trade practice under subsection
40 (a) of section 42-110b of the general statutes."

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This act shall take effect as follows and shall amend the following sections:			
Section 1	October 1, 2025	New section	
Sec. 2	October 1, 2025	New section	
Sec. 3	October 1, 2025	New section	