

**SB2353**



**103RD GENERAL ASSEMBLY**

**State of Illinois**

**2023 and 2024**

**SB2353**

Introduced 2/10/2023, by Sen. Rachel Ventura

**SYNOPSIS AS INTRODUCED:**

720 ILCS 570/302

from Ch. 56 1/2, par. 1302

Amends the Illinois Controlled Substances Act. Provides that notwithstanding any other provision of the Act to the contrary, including the scheduling of psilocybin as a Schedule I controlled substance, the Department of Financial and Professional Regulation shall authorize the distribution of, and make publicly available, psilocybin for medical, psychological, and scientific studies, research, and other information relating to the safety and efficacy of psilocybin and other entheogens to treat mental health conditions, including, but not limited to, addiction, depression, anxiety disorders, headache disorders, and end-of-life psychological distress. Provides that the Department of Financial and Professional Regulation shall begin receiving applications for the registration of persons to perform the following: (1) manufacturing psilocybin products; (2) operating a service center; (3) facilitating psilocybin services; and (4) testing psilocybin products.

LRB103 28977 RLC 55363 b

**A BILL FOR**

1 AN ACT concerning criminal law.

2 **Be it enacted by the People of the State of Illinois,**  
3 **represented in the General Assembly:**

4 Section 5. The Illinois Controlled Substances Act is  
5 amended by changing Section 302 as follows:

6 (720 ILCS 570/302) (from Ch. 56 1/2, par. 1302)

7 Sec. 302. (a) Every person who manufactures, distributes,  
8 or dispenses any controlled substances; engages in chemical  
9 analysis, research, or instructional activities which utilize  
10 controlled substances; purchases, stores, or administers  
11 euthanasia drugs, within this State; provides canine odor  
12 detection services; proposes to engage in the manufacture,  
13 distribution, or dispensing of any controlled substance;  
14 proposes to engage in chemical analysis, research, or  
15 instructional activities which utilize controlled substances;  
16 proposes to engage in purchasing, storing, or administering  
17 euthanasia drugs; or proposes to provide canine odor detection  
18 services within this State, must obtain a registration issued  
19 by the Department of Financial and Professional Regulation in  
20 accordance with its rules. The rules shall include, but not be  
21 limited to, setting the expiration date and renewal period for  
22 each registration under this Act. The Department, any facility  
23 or service licensed by the Department, and any veterinary

1 hospital or clinic operated by a veterinarian or veterinarians  
2 licensed under the Veterinary Medicine and Surgery Practice  
3 Act of 2004 or maintained by a State-supported or publicly  
4 funded university or college shall be exempt from the  
5 regulation requirements of this Section; however, such  
6 exemption shall not operate to bar the University of Illinois  
7 from requesting, nor the Department of Financial and  
8 Professional Regulation from issuing, a registration to the  
9 University of Illinois Veterinary Teaching Hospital under this  
10 Act. Neither a request for such registration nor the issuance  
11 of such registration to the University of Illinois shall  
12 operate to otherwise waive or modify the exemption provided in  
13 this subsection (a).

14 (b) Persons registered by the Department of Financial and  
15 Professional Regulation under this Act to manufacture,  
16 distribute, or dispense controlled substances, engage in  
17 chemical analysis, research, or instructional activities which  
18 utilize controlled substances, purchase, store, or administer  
19 euthanasia drugs, or provide canine odor detection services,  
20 may possess, manufacture, distribute, engage in chemical  
21 analysis, research, or instructional activities which utilize  
22 controlled substances, dispense those substances, or purchase,  
23 store, or administer euthanasia drugs, or provide canine odor  
24 detection services to the extent authorized by their  
25 registration and in conformity with the other provisions of  
26 this Article.

1       (b-5) Notwithstanding any other provision of this Act to  
2 the contrary, including the scheduling of psilocybin as a  
3 Schedule I controlled substance, the Department of Financial  
4 and Professional Regulation shall authorize the distribution  
5 of, and make publicly available, psilocybin for medical,  
6 psychological, and scientific studies, research, and other  
7 information relating to the safety and efficacy of psilocybin  
8 and other entheogens to treat mental health conditions,  
9 including, but not limited to, addiction, depression, anxiety  
10 disorders, headache disorders, and end-of-life psychological  
11 distress. The Department of Financial and Professional  
12 Regulation shall begin receiving applications for the  
13 registration of persons to perform the following:

- 14           (1) manufacturing psilocybin products;  
15           (2) operating a service center;  
16           (3) facilitating psilocybin services; and  
17           (4) testing psilocybin products.

18       (c) The following persons need not register and may  
19 lawfully possess controlled substances under this Act:

20           (1) an agent or employee of any registered  
21 manufacturer, distributor, or dispenser of any controlled  
22 substance if he or she is acting in the usual course of his  
23 or her employer's lawful business or employment;

24           (2) a common or contract carrier or warehouseman, or  
25 an agent or employee thereof, whose possession of any  
26 controlled substance is in the usual lawful course of such

1 business or employment;

2 (3) an ultimate user or a person in possession of a  
3 controlled substance prescribed for the ultimate user  
4 under a lawful prescription of a practitioner, including  
5 an advanced practice registered nurse, practical nurse, or  
6 registered nurse licensed under the Nurse Practice Act, or  
7 a physician assistant licensed under the Physician  
8 Assistant Practice Act of 1987, who provides hospice  
9 services to a hospice patient or who provides home health  
10 services to a person, or a person in possession of any  
11 controlled substance pursuant to a lawful prescription of  
12 a practitioner or in lawful possession of a Schedule V  
13 substance. In this Section, "home health services" has the  
14 meaning ascribed to it in the Home Health, Home Services,  
15 and Home Nursing Agency Licensing Act; and "hospice  
16 patient" and "hospice services" have the meanings ascribed  
17 to them in the Hospice Program Licensing Act;

18 (4) officers and employees of this State or of the  
19 United States while acting in the lawful course of their  
20 official duties which requires possession of controlled  
21 substances;

22 (5) a registered pharmacist who is employed in, or the  
23 owner of, a pharmacy licensed under this Act and the  
24 Federal Controlled Substances Act, at the licensed  
25 location, or if he or she is acting in the usual course of  
26 his or her lawful profession, business, or employment;

1           (6) a holder of a temporary license issued under  
2           Section 17 of the Medical Practice Act of 1987 practicing  
3           within the scope of that license and in compliance with  
4           the rules adopted under this Act. In addition to  
5           possessing controlled substances, a temporary license  
6           holder may order, administer, and prescribe controlled  
7           substances when acting within the scope of his or her  
8           license and in compliance with the rules adopted under  
9           this Act.

10          (d) A separate registration is required at each place of  
11          business or professional practice where the applicant  
12          manufactures, distributes, or dispenses controlled substances,  
13          or purchases, stores, or administers euthanasia drugs. Persons  
14          are required to obtain a separate registration for each place  
15          of business or professional practice where controlled  
16          substances are located or stored. A separate registration is  
17          not required for every location at which a controlled  
18          substance may be prescribed.

19          (e) The Department of Financial and Professional  
20          Regulation or the Illinois State Police may inspect the  
21          controlled premises, as defined in Section 502 of this Act, of  
22          a registrant or applicant for registration in accordance with  
23          this Act and the rules promulgated hereunder and with regard  
24          to persons licensed by the Department, in accordance with  
25          subsection (bb) of Section 30-5 of the Substance Use Disorder  
26          Act and the rules and regulations promulgated thereunder.

1 (Source: P.A. 99-163, eff. 1-1-16; 99-247, eff. 8-3-15;  
2 99-642, eff. 7-28-16; 100-513, eff. 1-1-18; 100-759, eff.  
3 1-1-19.)