



102ND GENERAL ASSEMBLY

State of Illinois

2021 and 2022

HB4556

Introduced 1/21/2022, by Rep. Will Guzzardi

SYNOPSIS AS INTRODUCED:

410 ILCS 710/5
410 ILCS 710/10 new
410 ILCS 710/15 new
410 ILCS 710/20 new

Amends the Overdose Prevention and Harm Reduction Act. Provides that a pharmacist or physician may dispense drug adulterant testing supplies, such as reagents, test strips, or quantification instruments, to any person. Provides that no employee or volunteer of or participant in a program established under the Act or any employee or customer of a pharmacy, hospital, clinic, or other health care facility or medical office dispensing drug adulterant testing supplies in accordance with the Act shall be charged with or prosecuted for possession of specified materials. Provides that a law enforcement officer who, acting on good faith, arrests or charges a person who is thereafter determined to be entitled to immunity from prosecution shall not be subject to civil liability for the arrest or filing of charges. Provides that any record of a person that is created or obtained for use by a needle and hypodermic syringe access program or by a pharmacy, hospital, clinic, or other health care facility or medical office in connection with the dispensing of drug adulterant testing supplies must be kept confidential. Contains other provisions. Effective immediately.

LRB102 23790 CPF 33826 b

1 AN ACT concerning health.

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

4 Section 5. The Overdose Prevention and Harm Reduction Act
5 is amended by changing Section 5 and by adding Sections 10, 15,
6 and 20 as follows:

7 (410 ILCS 710/5)

8 Sec. 5. Needle and hypodermic syringe access program.

9 (a) Any governmental or nongovernmental organization,
10 including a local health department, community-based
11 organization, or a person or entity, that promotes
12 scientifically proven ways of mitigating health risks
13 associated with drug use and other high-risk behaviors may
14 establish and operate a needle and hypodermic syringe access
15 program. The objective of the program shall be accomplishing
16 all of the following:

17 (1) reducing the spread of HIV, AIDS, viral hepatitis,
18 and other bloodborne diseases;

19 (2) reducing the potential for needle stick injuries
20 from discarded contaminated equipment; and

21 (3) facilitating connections or linkages to
22 evidence-based treatment.

23 (b) Programs established under this Act shall provide all

1 of the following:

2 (1) Disposal of used needles and hypodermic syringes.

3 (2) Needles, hypodermic syringes, and other safer drug
4 consumption supplies, at no cost and in quantities
5 sufficient to ensure that needles, hypodermic syringes, or
6 other supplies are not shared or reused.

7 (3) Educational materials or training on:

8 (A) overdose prevention and intervention; and

9 (B) the prevention of HIV, AIDS, viral hepatitis,
10 and other common bloodborne diseases resulting from
11 shared drug consumption equipment and supplies.

12 (4) Access to opioid antagonists approved for the
13 reversal of an opioid overdose, or referrals to programs
14 that provide access to opioid antagonists approved for the
15 reversal of an opioid overdose.

16 (5) Linkages to needed services, including mental
17 health treatment, housing programs, substance use disorder
18 treatment, and other relevant community services.

19 (6) Individual consultations from a trained employee
20 tailored to individual needs.

21 (7) If feasible, a hygienic, separate space for
22 individuals who need to administer a prescribed injectable
23 medication that can also be used as a quiet space to gather
24 composure in the event of an adverse on-site incident,
25 such as a nonfatal overdose.

26 (8) If feasible, access to on-site drug adulterant

1 testing supplies such as reagents, test strips, or
2 quantification instruments that provide critical real-time
3 information on the composition of substances obtained for
4 consumption.

5 (c) (Blank). ~~Notwithstanding any provision of the Illinois~~
6 ~~Controlled Substances Act, the Drug Paraphernalia Control Act,~~
7 ~~or any other law, no employee or volunteer of or participant in~~
8 ~~a program established under this Act shall be charged with or~~
9 ~~prosecuted for possession of any of the following:~~

10 ~~(1) Needles, hypodermic syringes, or other drug~~
11 ~~consumption paraphernalia obtained from or returned,~~
12 ~~directly or indirectly, to a program established under~~
13 ~~this Act.~~

14 ~~(2) Residual amounts of a controlled substance~~
15 ~~contained in used needles, used hypodermic syringes, or~~
16 ~~other used drug consumption paraphernalia obtained from or~~
17 ~~returned, directly or indirectly, to a program established~~
18 ~~under this Act.~~

19 ~~(3) Drug adulterant testing supplies such as reagents,~~
20 ~~test strips, or quantification instruments obtained from~~
21 ~~or returned, directly or indirectly, to a program~~
22 ~~established under this Act.~~

23 ~~(4) Any residual amounts of controlled substances used~~
24 ~~in the course of testing the controlled substance to~~
25 ~~determine the chemical composition and potential threat of~~
26 ~~the substances obtained for consumption that are obtained~~

1 ~~from or returned, directly or indirectly, to a program~~
2 ~~established under this Act.~~

3 ~~In addition to any other applicable immunity or limitation~~
4 ~~on civil liability, a law enforcement officer who, acting on~~
5 ~~good faith, arrests or charges a person who is thereafter~~
6 ~~determined to be entitled to immunity from prosecution under~~
7 ~~this subsection (c) shall not be subject to civil liability~~
8 ~~for the arrest or filing of charges.~~

9 (d) Prior to the commencing of operations of a program
10 established under this Act, the governmental or
11 nongovernmental organization shall submit to the Illinois
12 Department of Public Health all of the following information:

13 (1) the name of the organization, agency, group,
14 person, or entity operating the program;

15 (2) the areas and populations to be served by the
16 program; and

17 (3) the methods by which the program will meet the
18 requirements of subsection (b) of this Section.

19 The Department of Public Health may adopt rules to
20 implement this subsection.

21 (Source: P.A. 101-356, eff. 8-9-19.)

22 (410 ILCS 710/10 new)

23 Sec. 10. Dispensing of drug adulterant testing supplies. A
24 pharmacist or physician may dispense drug adulterant testing
25 supplies such as reagents, test strips, or quantification

1 instruments to any person. Drug adulterant testing supplies
2 dispensed under this Section must be stored at a pharmacy,
3 hospital, clinic, or other health care facility licensed under
4 State or federal law or at the medical office of a physician
5 and in a manner that limits access to the drug adulterant
6 testing supplies to pharmacists and physicians employed at the
7 pharmacy, hospital, clinic, or other health care facility,
8 medical office, and any person designated by the pharmacist or
9 physician. Drug adulterant testing supplies dispensed under
10 this Section at a retail store containing a pharmacy may be
11 dispensed only from the pharmacy department of the retail
12 store.

13 (410 ILCS 710/15 new)

14 Sec. 15. Waiver of criminal penalties. Notwithstanding any
15 provision of the Illinois Controlled Substances Act, the Drug
16 Paraphernalia Control Act, or any other law, no employee or
17 volunteer of or participant in a program established under
18 this Act or any employee or customer of a pharmacy, hospital,
19 clinic, or other health care facility or medical office
20 dispensing drug adulterant testing supplies in accordance with
21 Section 10 of this Act shall be charged with or prosecuted for
22 possession of any of the following:

23 (1) Needles, hypodermic syringes, or other drug
24 consumption paraphernalia obtained from or returned,
25 directly or indirectly, to a program established under

1 this Act.

2 (2) Residual amounts of a controlled substance
3 contained in used needles, used hypodermic syringes, or
4 other used drug consumption paraphernalia obtained from or
5 returned, directly or indirectly, to a program established
6 under this Act.

7 (3) Drug adulterant testing supplies such as reagents,
8 test strips, or quantification instruments obtained from
9 or returned, directly or indirectly, to a program
10 established under this Act or a pharmacy, hospital,
11 clinic, or other health care facility or medical office
12 dispensing such drug adulterant testing supplies in
13 accordance with Section 10.

14 (4) Residual amounts of a controlled substance used in
15 the course of testing the controlled substance to
16 determine the chemical composition and potential threat of
17 the substance obtained for consumption that is obtained
18 from or returned, directly or indirectly, to a program
19 established under this Act.

20 In addition to any other applicable immunity or limitation
21 on civil liability, a law enforcement officer who, acting on
22 good faith, arrests or charges a person who is thereafter
23 determined to be entitled to immunity from prosecution under
24 this Section shall not be subject to civil liability for the
25 arrest or filing of charges.

1 (410 ILCS 710/20 new)

2 Sec. 20. Confidentiality.

3 (a) Any record of a person that is created or obtained for
4 use by a needle and hypodermic syringe access program or by a
5 pharmacy, hospital, clinic, or other health care facility or
6 medical office in connection with the dispensing of drug
7 adulterant testing supplies in accordance with Section 10 must
8 be kept confidential and:

9 (1) is not open for public inspection or disclosure;

10 (2) must not be shared with any other person or entity
11 without the consent of the person to whom the record
12 relates; and

13 (3) is not discoverable or admissible during any legal
14 proceeding.

15 (b) A record described in subsection (a) must not be used:

16 (1) to initiate or substantiate any criminal charge
17 against a person who participates in the needle and
18 hypodermic syringe access program or who obtains drug
19 adulterant testing supplies from a pharmacy, hospital,
20 clinic, or other health care facility or medical office in
21 accordance with Section 10; or

22 (2) as grounds for conducting any investigation of a
23 person who participates in the needle and hypodermic
24 syringe access program or who obtains drug adulterant
25 testing supplies from a pharmacy, hospital, clinic, or
26 other health care facility or medical office in accordance

1 with Section 10.

2 (c) The staff and volunteers of a needle and hypodermic
3 syringe access program or a pharmacy, hospital, clinic, or
4 other health care facility or medical office dispensing drug
5 adulterant testing supplies in accordance with Section 10
6 shall not be compelled to provide evidence in any criminal
7 proceeding conducted pursuant to the laws of this State
8 concerning any information that was entrusted to them or
9 became known to them through the program or through the
10 dispensing.

11 (d) The use of any personal information of (i) any person
12 who participates in a needle and hypodermic syringe access
13 program or obtains drug adulterant testing supplies from a
14 pharmacy, hospital, clinic, or other health care facility or
15 medical office in accordance with Section 10, or (ii) the
16 staff or volunteers of the needle and hypodermic syringe
17 access program or the pharmacy, hospital, clinic, or other
18 health care facility or medical office, in research and
19 evaluation, must be done in such a manner as to guarantee the
20 anonymity of the person.

21 (e) Aggregate data from a needle and hypodermic syringe
22 access program, including, without limitation, demographic
23 information, the number of clients contacted, and the types of
24 referrals may be made available to the public.

25 Section 99. Effective date. This Act takes effect upon
26 becoming law.