

Requested by Representative DEXTER

**PROPOSED AMENDMENTS TO  
HOUSE BILL 2397**

1 In line 2 of the printed bill, after “prevention” insert “; creating new  
2 provisions; and amending ORS 414.318, 414.320 and 475.525”.

3 Delete lines 4 through 8 and insert:

4 **“SECTION 1. (1) As used in this section, ‘harm reduction supplies’**  
5 **means opioid reversal medications, hypodermic syringes or needles,**  
6 **single-use drug test strips, drug testing tools or any other item de-**  
7 **signed to prevent or reduce the potential harm associated with the use**  
8 **of opioids and other controlled substances, including but not limited**  
9 **to items that reduce the risk of transmission of infectious disease or**  
10 **prevent injury, infection or overdose.**

11 **“(2) The Harm Reduction Clearinghouse Project is established in the**  
12 **Oregon Health Authority to purchase in bulk harm reduction supplies**  
13 **for use throughout this state by community organizations, first**  
14 **responders and other entities that serve populations who are vulner-**  
15 **able to overdose, infections or injuries due to opioid use and use of**  
16 **other controlled substances. Entities that may participate in the**  
17 **project include but are not limited to:**

18 **“(a) Hospitals and emergency departments;**

19 **“(b) First responders;**

20 **“(c) Law enforcement agencies;**

21 **“(d) Courts and other departments within the criminal justice sys-**

1 **tem;**

2 **“(e) Organizations that provide services to individuals experiencing**  
3 **homelessness;**

4 **“(f) Organizations that provide services to individuals at risk of**  
5 **overdose, infections or injuries related to opioid use or use of other**  
6 **controlled substances;**

7 **“(g) Veterans’ organizations;**

8 **“(h) Religious organizations;**

9 **“(i) Schools and universities;**

10 **“(j) Substance use treatment and recovery facilities, including in-**  
11 **patient facilities, outpatient facilities, residential facilities and sober-**  
12 **ing centers;**

13 **“(k) Public libraries;**

14 **“(L) Community health centers;**

15 **“(m) County public health or behavioral health agencies; and**

16 **“(n) Special districts.**

17 **“(3) To make the bulk purchases of harm reduction supplies under**  
18 **this section, the administrator of the Harm Reduction Clearinghouse**  
19 **Project may use funds from the Opioid Reversal Medication and Harm**  
20 **Reduction Clearinghouse Bulk Purchasing Fund established in section**  
21 **2 of this 2023 Act or from gifts, grants, bequests, endowments or do-**  
22 **nations made to the Prescription Drug Purchasing Fund established**  
23 **in ORS 414.318 for the purchase of harm reduction supplies.**

24 **“SECTION 2. (1) The Opioid Reversal Medication and Harm Re-**  
25 **duction Clearinghouse Bulk Purchasing Fund is established in the**  
26 **State Treasury, separate and distinct from the General Fund. Interest**  
27 **earned by the Opioid Reversal Medication and Harm Reduction Clear-**  
28 **inghouse Bulk Purchasing Fund shall be credited to the fund.**

29 **“(2) The Opioid Reversal Medication and Harm Reduction Clearing-**  
30 **house Bulk Purchasing Fund consists of moneys received by the**

1 **Oregon Health Authority from opioid litigation settlements, grants**  
2 **from the Substance Abuse and Mental Health Services Administration**  
3 **within the United States Department of Health and Human Services**  
4 **for the purpose of addressing the opioid overdose epidemic and moneys**  
5 **appropriated to the fund by the Legislative Assembly.**

6 **“(3) The moneys in the Opioid Reversal Medication and Harm Re-**  
7 **duction Clearinghouse Bulk Purchasing Fund are continuously appro-**  
8 **priated to the Oregon Health Authority for the purpose of carrying out**  
9 **section 1 of this 2023 Act.**

10 **“SECTION 3.** ORS 414.318 is amended to read:

11 **“414.318.** The Prescription Drug Purchasing Fund is established separate  
12 and distinct from the General Fund. The Prescription Drug Purchasing Fund  
13 shall consist of moneys appropriated to the fund by the Legislative Assembly  
14 and moneys received by the Oregon Health Authority for the purposes es-  
15 tablished in this section in the form of gifts, grants, bequests, endowments  
16 or donations. The moneys in the Prescription Drug Purchasing Fund are  
17 continuously appropriated to the authority and shall be used to purchase  
18 prescription drugs, reimburse pharmacies for prescription drugs, **purchase**  
19 **harm reduction supplies under section 2 of this 2023 Act** and reimburse  
20 the authority for the costs of administering the Oregon Prescription Drug  
21 Program, including contracted services costs, computer costs, professional  
22 dispensing fees paid to retail pharmacies and other reasonable program costs.  
23 Interest earned on the fund shall be credited to the fund.

24 **“SECTION 4.** ORS 414.320 is amended to read:

25 **“414.320.** The Oregon Health Authority shall adopt rules to implement and  
26 administer ORS 414.312 to 414.318 **and section 1 of this 2023 Act.** The rules  
27 shall include but are not limited to establishing procedures for:

28 **“(1)** Issuing prescription drug identification cards to individuals and en-  
29 tities that participate in the Oregon Prescription Drug Program; and

30 **“(2)** Enrolling pharmacies in the program.

1       **“SECTION 5.** ORS 475.525 is amended to read:

2       “475.525. (1) It is unlawful for any person to sell or deliver, possess with  
3 intent to sell or deliver or manufacture with intent to sell or deliver drug  
4 paraphernalia, knowing that it will be used to unlawfully plant, propagate,  
5 cultivate, grow, harvest, manufacture, compound, convert, produce, process,  
6 prepare, test, analyze, pack, repack, store, contain, conceal, inject, ingest,  
7 inhale or otherwise introduce into the human body a controlled substance  
8 as defined by ORS 475.005.

9       “(2) For the purposes of this section, ‘drug paraphernalia’ means all  
10 equipment, products and materials of any kind that are marketed for use or  
11 designed for use in planting, propagating, cultivating, growing, harvesting,  
12 manufacturing, compounding, converting, producing, processing, preparing,  
13 testing, analyzing, packaging, repackaging, storing, containing, concealing,  
14 injecting, ingesting, inhaling or otherwise introducing into the human body  
15 a controlled substance in violation of ORS 475.752 to 475.980. Drug par-  
16 aphernalia includes, but is not limited to:

17       “(a) Kits marketed for use or designed for use in unlawfully planting,  
18 propagating, cultivating, growing or harvesting of any species of plant that  
19 is a controlled substance or from which a controlled substance can be de-  
20 rived;

21       “(b) Kits marketed for use or designed for use in manufacturing, com-  
22 pounding, converting, producing, processing or preparing controlled sub-  
23 stances;

24       “(c) Isomerization devices marketed for use or designed for use in in-  
25 creasing the potency of any species of plant that is a controlled substance;

26       “[(d) *Testing equipment marketed for use or designed for use in identifying*  
27 *or in analyzing the strength, effectiveness or purity of controlled substances;*]

28       “[(e)] (d) Scales and balances marketed for use or designed for use in  
29 weighing or measuring controlled substances;

30       “[(f)] (e) Diluents and adulterants, such as quinine hydrochloride,

1 mannilol, mannite, dextrose and lactose, marketed for use or designed for  
2 use in cutting controlled substances;

3 “[g)] (f) Lighting equipment specifically designed for growing controlled  
4 substances;

5 “[h)] (g) Containers and other objects marketed for use or designed for  
6 use in storing or concealing controlled substances; and

7 “[i)] (h) Objects marketed for use or designed specifically for use in  
8 ingesting, inhaling or otherwise introducing a controlled substance into the  
9 human body, such as:

10 “[A) *Metal, wooden, acrylic, glass, stone, plastic or ceramic pipes with or*  
11 *without screens;*]

12 “[B) *Water pipes;*]

13 “[C) *Carburetion tubes and devices;*]

14 “[D)] (A) Smoking and carburetion masks;

15 “[E)] (B) Roach clips, meaning objects used to hold burning material that  
16 has become too small or too short to be held in the hand; **or**

17 “[F)] (C) Miniature cocaine spoons and cocaine vials[;].

18 “[G) *Chamber pipes;*]

19 “[H) *Carburetor pipes;*]

20 “[I) *Electric pipes;*]

21 “[J) *Air-driven pipes;*]

22 “[K) *Chillums;*]

23 “[L) *Bongs; and*]

24 “[M) *Ice pipes or chillers.*]

25 “(3) For purposes of this section, ‘drug paraphernalia’ does not include  
26 hypodermic syringes or needles, **single-use drug test strips, drug testing**  
27 **tools or any other item designed to prevent or reduce the potential**  
28 **harm associated with the use of opioids and other controlled sub-**  
29 **stances, including but not limited to items that reduce the risk of**  
30 **transmission of infectious disease or prevent injury, infection or**

1 **overdose.**

2 “(4) The provisions of ORS 475.525 to 475.565 do not apply to persons  
3 registered under the provisions of ORS 475.125 or to persons specified as  
4 exempt from registration under the provisions of that statute.

5 “(5)(a) The provisions of ORS 475.525 to 475.565 do not apply to a person  
6 who sells or delivers marijuana paraphernalia as defined in ORS 475C.373 to  
7 a person 21 years of age or older.

8 “(b) In determining whether an object is drug paraphernalia under this  
9 section or marijuana paraphernalia under ORS 475C.373, a trier of fact shall  
10 consider, in addition to any other relevant factor, the following:

11 “(A) Any oral or written instruction provided with the object related to  
12 the object’s use;

13 “(B) Any descriptive material packaged with the object that explains or  
14 depicts the object’s use;

15 “(C) Any national or local advertising related to the object’s use;

16 “(D) Any proffered expert testimony related to the object’s use;

17 “(E) The manner in which the object is displayed for sale, if applicable;  
18 and

19 “(F) Any other proffered evidence substantiating the object’s intended use.

20 “(6) **A person acting in good faith is immune from civil liability for**  
21 **any act or omission of an action committed during the course of dis-**  
22 **tributing an item described in subsection (3) of this section.**

23 “**SECTION 6. The amendments to ORS 475.525 by section 5 of this**  
24 **2023 Act apply to conduct occurring on or after the effective date of**  
25 **this 2023 Act.”.**

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