
A BILL FOR AN ACT

RELATING TO HEALTH.

BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF HAWAII:

1 SECTION 1. The legislature finds that the National
2 Institute on Drug Abuse has estimated that eighteen million
3 Americans misused prescription medications at least once in
4 2017. All too often, addiction begins at home, stemming from
5 abused prescriptions or unused medications falling into the
6 wrong hands. Unused medications in households, especially
7 controlled substances, can further expose residents to potential
8 harm due to mistaken ingestion and increase the risk of theft
9 and assault. Requiring pharmacies to educate patients about
10 proper disposal of unused or expired controlled substances at
11 the time they are dispensed and make available to patients
12 certain disposal options and information concerning those
13 options, can help promote public health and safety.

14 Accordingly, the purpose of this Act is to require
15 pharmacies, when dispensing controlled substances, to:



- 1 (1) Provide written notice to patients advising them of
- 2 certain risks associated with not properly disposing
- 3 of unwanted or expired drugs;
- 4 (2) Make available certain drug disposal options; and
- 5 (3) Provide written informational materials concerning
- 6 available drug disposal options.

7 SECTION 2. Section 461-10.2, Hawaii Revised Statutes, is
8 amended to read as follows:

9 "[+]§461-10.2[+] Return for disposal of unused, remaining,
10 or expired drugs; pharmacy [~~options.~~] requirements; written
11 information. (a) A pharmacy that dispenses prescription drugs,
12 other than a long-term care pharmacy, when dispensing to an
13 individual located in this State a prescription drug or
14 medication which is classified as a controlled substance under
15 federal law, and when dispensing any other prescription drug or
16 medication as may be designated under chapter 329, shall:

- 17 (1) Provide the patient with written informational
- 18 materials advising that when unused, unwanted, or
- 19 expired drugs and medications are not properly,
- 20 safely, and promptly disposed of:



- 1 (A) There is a risk that the drug or medication can
2 be stolen, diverted, abused, misused, or
3 accidentally ingested, which can pose a risk to
4 the health and safety of the patient and other
5 members of the patient's household;
- 6 (B) Children are particularly at risk of accidentally
7 ingesting unused, unwanted, and expired
8 medications that have not been properly, safely,
9 and promptly disposed of;
- 10 (C) When drugs or medications are disposed of in the
11 household trash or flushed down the drain, the
12 drugs and medications can leak into the
13 ecosystem, which can have a potentially adverse
14 or harmful effect on the environment; and
- 15 (D) When drugs or medications are disposed of in the
16 household trash without the drug or medication
17 having been rendered deactivated, inaccessible,
18 or otherwise unusable, the drug or medication may
19 be stolen by individuals seeking to divert,
20 abuse, or misuse the drug or medication; and



1 (2) Make available on site, for purchase, or at no cost to
2 the patient, at least one consumer method for
3 individuals to dispose of unwanted or expired
4 prescription drugs, including but not limited to over
5 the counter, at home, or site-of-use compositions or
6 secured medication collection kiosks, boxes, or
7 receptacles, subject to the following requirements:
8 (A) All at home or site-of-use drug disposal products
9 shall alter the characteristics of the
10 prescription drug through chemical, biological,
11 or physical means so as to have a beneficial
12 effect on the environment;
13 (B) Secured medication collection kiosks, boxes, or
14 receptables shall comply with Title 21 Code of
15 Federal Regulations section 1317.75 and be marked
16 and identified by prominent signage;
17 (C) Any manufacturer of a non-toxic at home or site-
18 of-use composition for consumer drug disposal
19 shall provide a method that renders the active
20 ingredients in the prescription medication, as
21 defined in Title 21 Code of Federal Regulations



1 section 210.3(b)(7) or as defined in a successor
2 regulation, unusable so that the active
3 ingredients cannot be transformed to a physical
4 or chemical condition or transformed to the state
5 of a controlled substance or controlled substance
6 analog, as per Title 21 Code of Federal
7 Regulations section 1317.90, or a successor
8 regulation; and

9 (D) The manufacturer of an at home or site-of-use
10 composition or a secured medicine collection
11 kiosk, box, or receptacle made available by a
12 pharmacy pursuant to this paragraph shall
13 represent to the pharmacy that none of the
14 components or methods of disposal, individually
15 or as a blend or as a solution, or the methods of
16 treating or disposing of the medication at any
17 facility, are toxic, and that the composition or
18 medicine collection kiosk, box, or receptacle
19 follows waste regulations outlined by the United
20 States Environmental Protection Agency for
21 municipal household waste disposal; and



1 (3) Provide the patient with written informational
2 materials concerning how to properly, safely, and
3 promptly dispose of unused, unwanted, or expired drugs
4 and medications, which may include but shall not be
5 limited to information concerning drug disposal
6 options pursuant to paragraph (2) of this subsection
7 and any other available medication take back programs
8 in the State. The individual dispensing the
9 prescription drug, or an appropriate designee, shall
10 answer any questions the patient may have upon
11 receiving the written informational materials pursuant
12 to this paragraph.

13 (b) The requirements of subsection (a) of this section
14 shall apply regardless of whether the prescription is an initial
15 prescription or a renewal or refill of an existing prescription,
16 and regardless of whether the patient is a new or returning
17 customer at the pharmacy.

18 (c) Any time a pharmacy that dispenses prescription drugs,
19 other than a long-term care pharmacy, sells or dispenses a
20 sterile hypodermic syringe or needle, regardless of whether the
21 sterile hypodermic syringe or needle is sold or dispensed



1 pursuant to a prescription, the pharmacy shall provide the
2 patient with the written educational materials required under
3 section 325-21 regarding the safe disposal of used syringes at
4 sites where syringes are sold. The individual selling or
5 dispensing the sterile hypodermic syringe or needle, or an
6 appropriate designee, shall answer any questions the patient may
7 have upon receiving the written informational materials pursuant
8 to this subsection.

9 ~~[(a)]~~ (d) No pharmacy shall accept the return of any
10 prescription drug unless:

11 (1) The pharmacy is collecting the prescription drug for
12 disposal only; and

13 (2) The pharmacy is registered with the United States Drug
14 Enforcement Administration as an authorized collector
15 pursuant to title 21 Code of Federal Regulations
16 section 1317.40.

17 ~~[(b)]~~ (e) No prescription drug returned to the pharmacy
18 for disposal shall be redispensed or returned for cash or
19 credit.



1 ~~[(e)]~~ (f) Any pharmacy accepting for disposal any
2 prescription drugs ~~[for disposal]~~ other than those identified in
3 subsection (a) shall use the following methods:

4 (1) Secured collection receptacles in compliance with
5 title 21 Code of Federal Regulations section 1317.75;

6 or

7 (2) Mail-back programs.

8 ~~[(d)]~~ (g) In any pharmacy accepting prescription drugs for
9 disposal under this section, the pharmacist-in-charge shall
10 ensure that only Drug Enforcement Administration approved
11 reverse distributors acquire prescription drugs collected
12 through collection receptacles and mail-back programs."

13 SECTION 3. This Act does not affect rights and duties that
14 matured, penalties that were incurred, and proceedings that were
15 begun before its effective date.

16 SECTION 4. Statutory material to be repealed is bracketed
17 and stricken. New statutory material is underscored.

18 SECTION 5. This Act shall take effect on July 1, 2020.



Report Title:

Pharmacies; Prescription Drugs; Controlled Substances;
Requirements for Proper Disposal; Written Information

Description:

Requires pharmacies to provide written notice to patients advising them of certain risks associated with not properly disposing of unwanted or expired drugs, make available certain drug disposal options, and provide written informational materials concerning available drug disposal options. (Proposed SD1)

The summary description of legislation appearing on this page is for informational purposes only and is not legislation or evidence of legislative intent.

