
A BILL FOR AN ACT

RELATING TO PSEUDOEPHEDRINE.

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

1 SECTION 1. Section 329-75, Hawaii Revised Statutes, is
2 amended to read as follows:

3 "**§329-75 Sales of products, mixtures, or preparations**
4 **containing pseudoephedrine; reporting requirement for**
5 **wholesalers.** (a) Notwithstanding any other law to the
6 contrary, a pharmacy or retailer may sell or distribute to a
7 person without a prescription products containing not more than
8 3.6 grams per day~~[,]~~ or not more than nine grams per thirty-day
9 period of pseudoephedrine, without regard to the number of
10 transactions~~[, of any product, mixture, or preparation~~
11 ~~containing any detectable quantity of pseudoephedrine, its~~
12 ~~salts, optical isomers, or salts of optical isomers as the only~~
13 ~~active ingredient or in combination with other active~~
14 ~~ingredients]~~; provided that the pharmacy or retailer shall
15 comply with the following conditions:

16 (1) The product, mixture, or preparation shall be sold or
17 distributed from an area not accessible by customers
18 or the general public, such as behind the counter or



1 in a locked display case and where the [seller]
2 pharmacy or retailer delivers the product directly
3 into the custody of the [~~purchaser,~~] person purchasing
4 or obtaining the substances;

5 (2) Any person purchasing or otherwise [~~acquiring~~]
6 obtaining any product, mixture, or preparation shall
7 produce [~~proper~~] valid, government-issued
8 identification containing the photograph, date of
9 birth, printed name, signature, and address of the
10 [~~individual~~] person purchasing or obtaining the
11 substance;

12 (3) The pharmacy or retailer shall [~~record, in an~~
13 ~~electronic log on software provided by the narcotics~~
14 ~~enforcement division of the department and approved by~~
15 ~~the administrator,~~] maintain a written or electronic
16 log of required information for each sale of a
17 nonprescription product containing pseudoephedrine,
18 including:

19 (A) The date and time of any transaction under
20 paragraph (2);

21 (B) The name, address, and date of birth of the
22 person[+] purchasing or obtaining the substance;



1 (C) The type of identification provided by the
2 ~~[individual]~~ person purchasing or obtaining the
3 substance[+] and identification number;

4 (D) The agency issuing the identification used; and

5 (E) The name of the compound, mixture, or
6 preparation, and the amount; and

7 (4) The pharmacy or retailer shall[+]

8 ~~(A) Record the information required under paragraph~~
9 ~~(3) on an electronic worksheet on software~~
10 ~~provided by the narcotics enforcement division of~~
11 ~~the department; and~~

12 ~~(B) Electronically mail the worksheet record to the~~
13 ~~narcotics enforcement division once a month.]~~
14 require every person purchasing or obtaining the
15 substance to sign a written or electronic log
16 attesting to the validity of the information.

17 The information shall be retained by the pharmacy or
18 retailer for a period of two years. The written or
19 electronic log shall be capable of being checked for
20 compliance against all state and federal laws,
21 including interfacing with other states to ensure
22 comprehensive compliance, and shall be subject to



1 random and warrantless inspection by county or state
2 law enforcement officers.

3 (b) Beginning January 1, 2013, before completing a sale of
4 an over-the-counter product containing pseudoephedrine, a
5 pharmacy or retailer shall electronically submit the information
6 required pursuant to subsection (a) to the National Precursor
7 Log Exchange administered by the National Association of Drug
8 Diversion Investigators; provided that the National Precursor
9 Log Exchange is available to pharmacies or retailers in the
10 State without a charge for accessing the system. The pharmacy
11 or retailer shall not complete the sale if the system generates
12 a stop sale alert. Except in the case of negligence,
13 wantonness, recklessness, or deliberate misconduct, any pharmacy
14 or retailer using the electronic sales tracking system in
15 accordance with this subsection shall not be civilly liable as a
16 result of any act or omission in carrying out the duties
17 required by this subsection and shall be immune from liability
18 to any third party, unless the pharmacy or retailer has violated
19 this subsection, in relation to a claim brought for such
20 violation.

21 (c) If a pharmacy or retailer selling an over-the-counter
22 product containing pseudoephedrine experiences mechanical or



1 electronic failure of the electronic sales tracking system and
2 is unable to comply with the electronic sales tracking
3 requirement under this section, the pharmacy or retailer shall
4 maintain a written log or an alternative electronic
5 recordkeeping mechanism until such time as the pharmacy or
6 retailer is able to comply with the electronic sales tracking
7 requirement.

8 (d) A pharmacy or retailer selling an over-the-counter
9 product containing pseudoephedrine may seek an exemption from
10 submitting transactions to the electronic sales tracking system
11 in writing to the administrator stating the reasons therefore.
12 The administrator may grant an exemption for good cause shown,
13 but in no event shall the exemption exceed one hundred eighty
14 days. Any pharmacy or retailer that receives an exemption shall
15 maintain a hard copy log and shall require the person purchasing
16 or obtaining the substance to provide the information required
17 under this section before completion of any sale. The log shall
18 be maintained as a record of each sale for inspection by any law
19 enforcement officer or inspector of the board of pharmacy during
20 normal business hours.

21 (e) The National Association of Drug Diversion
22 Investigators shall forward Hawaii transaction records in the



1 National Precursor Log Exchange to the narcotics enforcement
2 division of the department of public safety weekly and provide
3 real-time access to National Precursor Log Exchange information
4 through the National Precursor Log Exchange online portal to law
5 enforcement in the State as authorized by the narcotics
6 enforcement division; provided that the narcotics enforcement
7 division executes a memorandum of understanding with the
8 National Association of Drug Diversion Investigators governing
9 access to the information; provided further that the department
10 of public safety narcotics enforcement division shall establish
11 the electronic tracking system in conjunction with the State's
12 existing narcotics tracking system beginning no later than
13 January 1, 2015.

14 (f) This system shall be capable of generating a stop sale
15 alert, which shall be a notification that completion of the sale
16 would result in the pharmacy or retailer, or person purchasing
17 or obtaining the substance, violating the quantity limits set
18 forth in this section. The system shall contain an override
19 function that may be used by a pharmacy or retailer selling
20 pseudoephedrine who has a reasonable fear that imminent bodily
21 harm will result if the sale is not completed. Each instance



1 where the override function is used shall be logged by the
2 system.

3 ~~[(b)]~~ (g) No person shall knowingly purchase, ~~[possess,]~~
4 receive, or otherwise acquire products containing more than 3.6
5 grams per day or more than nine grams ~~[of any product, mixture,~~
6 ~~or preparation containing any detectable quantity of~~
7 ~~pseudoephedrine or its salts, isomers, or salts of optical~~
8 ~~isomers within a thirty day period,]~~ per thirty-day period of
9 pseudoephedrine, except that this limit shall not apply to any
10 quantity of such product, mixture, or preparation dispensed
11 pursuant to a valid prescription.

12 ~~[(e)]~~ (h) Any person who violates ~~[subsection]~~ subsections
13 (b) through (g) is guilty of a class C felony.

14 ~~[(d)]~~ (i) The department, by rule, may exempt other products
15 from this section, if the administrator finds that the products
16 are not used in the illegal manufacture of methamphetamine or
17 other controlled substances. A manufacturer of a drug product
18 may apply for removal of the product from this section if the
19 product is determined by the administrator to have been
20 formulated in such a way as to effectively prevent the
21 conversion of the active ingredient into methamphetamine.



1 [~~(e)~~] (j) Notwithstanding any other provision of this
2 chapter to the contrary, every wholesaler shall report to the
3 administrator all sales made to any retailer, of any product,
4 mixture, or preparation containing any detectable quantity of
5 pseudoephedrine, its salts, optical isomers, or salts of optical
6 isomers, as the only active ingredient or in combination with
7 other active ingredients. The department shall provide a common
8 reporting form that contains at least the following information
9 about the product, mixture, or preparation:

- 10 (1) Generic or other name;
- 11 (2) Quantity sold;
- 12 (3) Date of sale;
- 13 (4) Name and address of the wholesaler; and
- 14 (5) Name and address of the retailer.

15 [~~(f)~~] (k) Intentional or knowing failure of a retailer or
16 pharmacy to transmit any information as required by this section
17 shall be a misdemeanor and shall result in the immediate
18 suspension of that retailer's ability to sell any product,
19 mixture, or preparation containing any detectable quantity of
20 pseudoephedrine, its salts, optical isomers, or salts of optical
21 isomers as the only active ingredient or in combination with
22 other active ingredients until authorized by the administrator."



1 SECTION 2. This Act does not affect rights and duties that
2 matured, penalties that were incurred, and proceedings that were
3 begun before its effective date.

4 SECTION 3. Statutory material to be repealed is bracketed
5 and stricken. New statutory material is underscored.

6 SECTION 4. This Act shall take effect upon its approval.



Report Title:

Pseudoephedrine; Tracking

Description:

Establishes an electronic tracking system for the sale of products containing pseudoephedrine. Limits the amount of pseudoephedrine products that may be sold or distributed to a person without a prescription to products containing not more than 3.6 grams of pseudoephedrine per day or not more than nine grams of pseudoephedrine per thirty-day period. Requires the pharmacy or retailer to maintain a written or electronic log of nonprescription products containing pseudoephedrine sold.
(SB2228 HD2)

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