
A BILL FOR AN ACT

RELATING TO CONTROLLED SUBSTANCES.

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

1 SECTION 1. Section 329-22, Hawaii Revised Statutes, is
2 amended to read as follows:

3 "**§329-22 Schedule V.** (a) The controlled substances
4 listed in this section are included in schedule V.

5 (b) Narcotic drugs containing nonnarcotic active medicinal
6 ingredients. Any compound, mixture, or preparation containing
7 limited quantities of any of the following narcotic drugs, which
8 also contains one or more nonnarcotic active medicinal
9 ingredients in sufficient proportion to confer upon the
10 compound, mixture, or preparation, valuable medicinal qualities
11 other than those possessed by the narcotic drug alone:

12 (1) Not more than 200 milligrams of codeine, or any of its
13 salts, per 100 milliliters or per 100 grams;

14 (2) Not more than 100 milligrams of dihydrocodeine, or any
15 of its salts, per 100 milliliters or per 100 grams;

16 (3) Not more than 100 milligrams of ethylmorphine, or any
17 of its salts, per 100 milliliters or per 100 grams;



1 (4) Not more than 2.5 milligrams of diphenoxylate and not
2 less than 25 micrograms of atropine sulfate per dosage
3 unit;

4 (5) Not more than 100 milligrams of opium per 100
5 milliliters or per 100 grams; and

6 (6) Not more than 0.5 milligram of difenoxin and not less
7 than 25 micrograms of atropine sulfate per dosage
8 unit.

9 (c) Stimulants. Unless specifically exempted or excluded
10 or unless listed in another schedule, any material, compound,
11 mixture, or preparation that contains any quantity of the
12 following substances having a stimulant effect on the central
13 nervous system, including its salts, isomers, and salts of
14 isomers:

15 (1) Ephedrine, its salts, optical isomers, and salts of
16 optical isomers as the only active ingredient, or in
17 combination with other active ingredients;

18 (2) Phenylpropanolamine, its salts, optical isomers, and
19 salts of optical isomers as the only active
20 ingredient, or in combination with other active
21 ingredients;



1 (3) Pseudoephedrine, its salts, optical isomers, and salts
2 of optical isomers as the only active ingredient, or
3 in combination with other active ingredients; and

4 ~~[(1)]~~ (4) Pyrovalerone.

5 (d) The department, by rule, may exempt from schedule V
6 any product containing any of the substances enumerated in
7 subsection (c)(1), (2), or (3) if the administrator finds that
8 the product is not used in the illegal manufacture of
9 methamphetamine or other controlled dangerous substances. A
10 manufacturer of such drug product may apply for removal of the
11 product from the schedule if the product is determined by the
12 administrator to have been formulated in such a way as to
13 effectively prevent the conversion of the active ingredient into
14 methamphetamine."

15 SECTION 2. Section 329-64, Hawaii Revised Statutes, is
16 amended by amending subsection (a) to read as follows:

17 "(a) The requirements imposed by sections 329-62~~[7]~~ and
18 329-63(a)~~[7, and 329-67]~~ of this part shall not apply to any of
19 the following:

20 (1) Any pharmacist or other authorized person who sells or
21 furnishes a substance upon the prescription of a
22 physician, dentist, podiatrist, or veterinarian;



1 (2) Any physician, dentist, podiatrist, or veterinarian
2 who administers or furnishes a substance to patients;

3 (3) Any manufacturer or wholesaler licensed by the State
4 who sells, transfers, or otherwise furnishes a
5 substance to a licensed pharmacy, physician, dentist,
6 podiatrist, or veterinarian; and

7 ~~[(4) Any sale, transfer, furnishing, or receipt of any drug
8 that contains pseudoephedrine or nonpseudoephedrine
9 that is lawfully sold, transferred, or furnished over
10 the counter without a prescription pursuant to the
11 federal Food, Drug, and Cosmetic Act (21 United States
12 Code Sec. 301 et seq.) or regulations adopted
13 thereunder as long as it complies with the
14 requirements of sections 329-73, 329-74, and 329-75;
15 and~~

16 ~~(5)]~~ (4) Any "dietary supplement" as defined by the federal
17 Food, Drug, and Cosmetic Act (21 United States Code
18 Sec. 301) containing ephedrine alkaloids extracted
19 from any species of Ephedra that meets all of the
20 following criteria:

21 (A) It contains, per dosage unit or serving, not more
22 than twenty-five milligrams of ephedrine



1 alkaloids and its labeling does not suggest or
2 recommend a total daily intake of more than one
3 hundred milligrams of ephedrine alkaloids;

4 (B) It contains no hydrochloride or sulfate salts of
5 ephedrine alkaloids; and

6 (C) It is packaged with a prominent label securely
7 affixed to each package that states all of the
8 following:

9 (i) The amount in milligrams of ephedrine
10 alkaloids in a dosage unit or serving;

11 (ii) The amount of the dietary supplement that
12 constitutes a dosage unit or serving; and

13 (iii) The maximum recommended dosage of ephedrine
14 alkaloids for a healthy adult human is not
15 more than one hundred milligrams in a
16 twenty-four-hour period.

17 Any dietary supplement that meets the criteria of this
18 paragraph shall also be exempt from the requirements
19 of section 329-67."

20 SECTION 3. Section 329-73, Hawaii Revised Statutes, is
21 repealed:



1 ~~["§329-73 Pseudoephedrine permit. (a) Beginning~~
2 ~~January 1, 2006, any person transporting by any means more than~~
3 ~~three packages of any product the sale of which is restricted by~~
4 ~~section 329-75 shall obtain a pseudoephedrine permit.~~

5 ~~(b) The requirements imposed by subsection (a) shall not~~
6 ~~apply to persons registered with the department under section~~
7 ~~329-67. A pseudoephedrine permit shall be issued by the~~
8 ~~department in a form and manner as prescribed by the department~~
9 ~~by rule. A pseudoephedrine permit shall be valid for one year~~
10 ~~and renewable annually."]~~

11 SECTION 4. Section 329-74, Hawaii Revised Statutes, is
12 repealed:

13 ~~["§329-74 Unlawful transport of pseudoephedrine. (a) A~~
14 ~~person commits the offense of unlawful transport of~~
15 ~~pseudoephedrine if the person transports more than three packages~~
16 ~~of any product the sale of which is restricted by section~~
17 ~~329-75 without a permit issued from the department.~~

18 ~~(b) For purposes of this section, "transportation" means~~
19 ~~the transfer of a pseudoephedrine product by a person other than~~
20 ~~a wholesaler, distributor, or retailer of such product authorized~~
21 ~~to conduct business as such by the State.~~



1 ~~(c) Unlawful transport of pseudoephedrine is a~~
2 ~~misdemeanor."~~]

3 SECTION 5. Section 329-75, Hawaii Revised Statutes, is
4 repealed:

5 ~~["§329-75 Sales of products, mixtures, or preparations~~
6 ~~containing pseudoephedrine; reporting requirement for~~

7 ~~wholesalers. (a) Notwithstanding any other law to the contrary,~~
8 ~~a pharmacy or retailer may dispense, sell, or distribute to a~~
9 ~~person without a prescription not more than 3.6 grams per day~~
10 ~~without regard to the number of transactions, of any product,~~
11 ~~mixture, or preparation containing any detectable quantity of~~
12 ~~pseudoephedrine, its salts, optical isomers, or salts of optical~~
13 ~~isomers, as the only active ingredient or in combination with~~
14 ~~other active ingredients; provided that the pharmacy or retailer~~
15 ~~complies with the following conditions:~~

16 ~~(1) The product, mixture, or preparation shall be~~
17 ~~dispensed, sold, or distributed from an area not~~
18 ~~accessible by customers or the general public, such as~~
19 ~~behind the counter or in a locked display case and~~
20 ~~where the seller delivers the product directly into~~
21 ~~the custody of the purchaser; and~~



1 ~~(2) Any person purchasing or otherwise acquiring any~~
2 ~~product, mixture, or preparation shall:~~

3 ~~(A) Produce proper identification containing the~~
4 ~~photograph, printed name, and signature of the~~
5 ~~individual obtaining the controlled substance,~~
6 ~~and~~

7 ~~(B) Sign a written log, receipt, or other program or~~
8 ~~mechanism approved by the administrator, showing~~
9 ~~the date of the transaction, name and address of~~
10 ~~the person, and the amount of the compound,~~
11 ~~mixture, or preparation.~~

12 ~~No person shall purchase, receive, or otherwise acquire~~
13 ~~more than nine grams of any product, mixture, or~~
14 ~~preparation containing any detectable quantity of~~
15 ~~pseudoephedrine or its salts, isomers, or salts of optical~~
16 ~~isomers within a thirty day period, except that this limit~~
17 ~~shall not apply to any quantity of such product, mixture,~~
18 ~~or preparation dispensed pursuant to a valid prescription.~~

19 ~~(b) The sales restriction in this section, as it applies~~
20 ~~to products, mixtures, or preparations containing any detectable~~
21 ~~quantity of pseudoephedrine, its salts, optical isomers, or~~
22 ~~salts of optical isomers, shall not apply to any products,~~



1 ~~mixtures, or preparations that are in liquid, liquid capsule, or~~
2 ~~gel capsule form if pseudoephedrine is not the only active~~
3 ~~ingredient.~~

4 ~~(c) The department, by rule, may exempt other products~~
5 ~~from this section, if the administrator finds that the products~~
6 ~~are not used in the illegal manufacture of methamphetamine or~~
7 ~~other controlled substances. A manufacturer of a drug product~~
8 ~~may apply for removal of the product from this section if the~~
9 ~~product is determined by the administrator to have been~~
10 ~~formulated in such a way as to effectively prevent the~~
11 ~~conversion of the active ingredient into methamphetamine.~~

12 ~~(d) Notwithstanding any other provision of this chapter to~~
13 ~~the contrary, every wholesaler shall report to the administrator~~
14 ~~all sales made to any retailer, of any product, mixture, or~~
15 ~~preparation containing any detectable quantity of~~
16 ~~pseudoephedrine, its salts, optical isomers, or salts of optical~~
17 ~~isomers, as the only active ingredient or in combination with~~
18 ~~other active ingredients. The department shall provide a common~~
19 ~~reporting form that contains at least the following information~~
20 ~~about the product, mixture, or preparation:~~

21 ~~(1) Generic or other name;~~

22 ~~(2) Quantity sold;~~



1 ~~(3) Date of sale;~~

2 ~~(4) Name and address of the wholesaler; and~~

3 ~~(5) Name and address of the retailer."]~~

4 SECTION 6. Statutory material to be repealed is bracketed
5 and stricken. New statutory material is underscored.

6 SECTION 7. This Act shall take effect on July 1, 2007.

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JAN 18 2007



Report Title:

Ephedrine; Pseudoephedrine; Phenylpropanolamine; Schedule V

Description:

Adds ephedrine, pseudoephedrine, and phenylpropanolamine to the list of Schedule V controlled substances. Repeals certain exemptions from business permits for regulated chemicals for the manufacture of controlled substances.

