



EXECUTIVE CHAMBERS
HONOLULU

NEIL ABERCROMBIE
GOVERNOR

July 1, 2013

GOV. MSG. NO. 1353

The Honorable Donna Mercado Kim,
President
and Members of the Senate
Twenty-Seventh State Legislature
State Capitol, Room 409
Honolulu, Hawaii 96813

The Honorable Joseph M. Souki,
Speaker and Members of the
House of Representatives
Twenty-Seventh State Legislature
State Capitol, Room 431
Honolulu, Hawaii 96813

Dear President Kim, Speaker Souki, and Members of the Legislature:

This is to inform you that on July 1, 2013, the following bill was signed into law:

SB655 SD2 HD2 CD1

RELATING TO HEALTH
ACT 250 (13)

NEIL ABERCROMBIE
Governor, State of Hawaii

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 many patients who
2 have been diagnosed with sexually transmitted diseases,
3 including chlamydia and gonorrhoea, have sexual partners who
4 refuse to seek treatment. To prevent reinfection, adequate
5 treatment of sexually transmitted diseases should include
6 treatment of sexual partners. Expedited partner therapy is a
7 partner treatment approach where partners of patients who test
8 positive for certain sexually transmitted diseases are provided
9 medication without previous medical evaluation.

10 The legislature further finds that because of expedited
11 partner therapy's effectiveness in reducing reinfection rates,
12 the Centers for Disease Control and Prevention has recommended
13 its use since 2006 among heterosexual partners of patients
14 diagnosed with chlamydia or gonorrhoea when it is unlikely the
15 partners will seek timely evaluation and treatment. The
16 legislature additionally finds that Hawaii has high reported
17 rates of chlamydia. The most recent Centers for Disease Control
18 and Prevention data ranks Hawaii twenty-second in the nation for



1 reported chlamydia infection rates, with the disease peaking in
2 the age group between fifteen and twenty-four years.

3 The legislature also finds that primary care providers,
4 including persons licensed under chapter 453, Hawaii Revised
5 Statutes, and advanced practice registered nurses with
6 prescriptive authority under chapter 457, Hawaii Revised
7 Statutes, currently diagnose and treat persons with sexually
8 transmitted diseases. Expedited partner therapy will permit
9 these health professionals to adequately treat sexually
10 transmitted diseases and prevent reinfection through the
11 treatment of sexual partners.

12 The purpose of this Act is to allow health professionals to
13 provide expedited partner therapy, in accordance with Centers
14 for Disease Control and Prevention guidelines and
15 recommendations, to the partners of a patient who has been
16 diagnosed as having a sexually transmitted disease.

17 SECTION 2. Chapter 453, Hawaii Revised Statutes, is
18 amended by adding a new part to be appropriately designated and
19 to read as follows:

20 "PART . EXPEDITED PARTNER THERAPY

21 §453-A Definitions. As used in this part:



1 "Expedited partner therapy" means the indirect treatment of
2 partners of a patient who has been diagnosed as having a
3 sexually transmitted disease through the dispensing or
4 prescribing of antibiotic therapy for the treatment of the
5 partners to the patient without the physical examination of the
6 partners by a health professional.

7 "Health professional" means any of the following:

- 8 (1) A person licensed or otherwise authorized by law to
9 practice medicine or surgery under this chapter and
10 whose scope of practice includes the diagnosis and
11 treatment of sexually transmitted diseases;
- 12 (2) An advanced practice registered nurse with
13 prescriptive authority under chapter 457 and duly
14 licensed in the State; or
- 15 (3) For the purpose of dispensing antibiotic therapy under
16 this section, a pharmacist who is licensed or
17 otherwise authorized to engage in the practice of
18 pharmacy under chapter 461.

19 "Sexual activity" means sexual intercourse, cunnilingus,
20 fellatio, anal intercourse, or any other intrusion, however
21 slight, of any part of a person's body or of any object into the



1 genital or anal openings of another person's body, but emission
2 of semen is not required.

3 "Sexually transmitted disease" means chlamydia, gonorrhea,
4 or other sexually transmitted diseases that are or may be
5 recommended by the Centers for Disease Control and Prevention
6 for expedited partner therapy.

7 **§453-B Expedited partner therapy.** (a) A health
8 professional may, in addition to treating a patient, provide
9 expedited partner therapy to the partners of the patient if all
10 of the following requirements are met:

11 (1) The patient has a laboratory-confirmed or suspected
12 clinical diagnosis of a sexually transmitted disease;

13 (2) The patient indicates that the patient has partners
14 with whom the patient has engaged in sexual activity
15 within the sixty-day period immediately preceding the
16 diagnosis of a sexually transmitted disease; and

17 (3) The patient indicates that the patient's partners are
18 unable or unlikely to seek clinical services in a
19 timely manner.

20 (b) A health professional who provides expedited partner
21 therapy as authorized in this section shall do all of the
22 following:



- 1 (1) Dispense or prescribe antibiotic therapy in the name
2 of the partners, if known, without the physical
3 examination of the partners by the health
4 professional. Notwithstanding any law to the
5 contrary, if the name of the partners are not known,
6 the health professional shall dispense or prescribe
7 the antibiotic therapy in the name of "Expedited
8 Partner Therapy";
- 9 (2) Convey to the patient that it is important to notify
10 the patient's partners of the patient's diagnosis and
11 that it is important for the partners to obtain
12 medical care for a complete evaluation, testing for
13 sexually transmitted diseases, counseling, and
14 treatment;
- 15 (3) Distribute to the patient the information sheet
16 developed pursuant to section 453-C; and
- 17 (4) Follow all Centers for Disease Control and Prevention
18 guidelines related to the practices and
19 recommendations for expedited partner therapy.
- 20 **§453-C Information sheet.** The department of health shall
21 develop and, upon request, distribute to health professionals



1 subject to this part an information sheet that includes all of
2 the following:

- 3 (1) A description of expedited partner therapy and its
4 purpose;
- 5 (2) A notice that an individual who has been treated for a
6 sexually transmitted disease should be retested after
7 treatment to detect possible persistent or recurrent
8 infection, including information on the timing of
9 retesting, as recommended by the Centers for Disease
10 Control and Prevention;
- 11 (3) A warning about the possible dangers of administering
12 antibiotic therapy to a pregnant individual;
- 13 (4) Information about antibiotics dispensed or prescribed
14 and dosages of those antibiotics dispensed or
15 prescribed, as recommended by the Centers for Disease
16 Control and Prevention;
- 17 (5) A warning about the risk of allergies to and drug
18 interactions with the antibiotics described in
19 paragraph (4);
- 20 (6) Information about sexually transmitted diseases, the
21 treatment of sexually transmitted diseases, and the
22 prevention of sexually transmitted diseases;



- 1 (7) A notice that the patient and the patient's partners
2 should abstain from sexual activity for seven days
3 after the patient and the partners have completed the
4 antibiotic therapy;
- 5 (8) A notice that the partners should be tested for
6 sexually transmitted diseases;
- 7 (9) A notice of the risk to the patient, the partners, and
8 others, including the public health, if a sexually
9 transmitted disease is not completely treated;
- 10 (10) A notice of the responsibility of the patient to
11 notify sexual partners of the risk of sexually
12 transmitted diseases and the importance of examination
13 and treatment for sexually transmitted diseases; and
- 14 (11) A statement advising any individual who has any
15 questions regarding anything in the information sheet
16 to contact a health professional or the department of
17 health.

18 **§453-D Limitation of liability.** A health professional who
19 provides expedited partner therapy as authorized under section
20 453-B, a person licensed or otherwise authorized by law to
21 practice medicine or surgery under this chapter, an advanced
22 practice registered nurse with prescriptive authority under



1 chapter 457, or a pharmacist who is licensed or otherwise
2 authorized to engage in the practice of pharmacy under chapter
3 461, who reasonably and in good faith renders the expedited
4 partner therapy in accordance with this section and the rules
5 and regulations adopted by the director of commerce and consumer
6 affairs, shall not be subject to civil or criminal liability or
7 be deemed to have engaged in unprofessional conduct for
8 rendering that therapy."

9 SECTION 3. Chapter 457, Hawaii Revised Statutes, is
10 amended by adding a new section to be appropriately designated
11 and to read as follows:

12 "§457- Advanced practice registered nurses; expedited
13 partner therapy. Advanced practice registered nurses who meet
14 the definition of a health professional as defined in section
15 453-A, shall be authorized to provide expedited partner therapy
16 in accordance with part of chapter 453."

17 SECTION 4. Section 328-16, Hawaii Revised Statutes, is
18 amended as follows:

19 1. By amending subsections (a), (b), and (c) to read:

20 "(a) A prescription drug shall be dispensed only if its
21 label bears the following:



- 1 (1) The name, business address, and telephone number of
- 2 the seller. The business address shall be the
- 3 physical location of the pharmacy or the dispensing
- 4 practitioner's office;
- 5 (2) [The] Except as otherwise authorized for expedited
- 6 partner therapy in section 453-B, the name of the
- 7 person for whom the drug was prescribed or the name of
- 8 the owner of the animal for which the drug was
- 9 prescribed;
- 10 (3) The serial number of the prescription;
- 11 (4) The date the prescription was prepared;
- 12 (5) The name of the practitioner if the seller is not the
- 13 practitioner;
- 14 (6) The name, strength, and quantity of the drug;
- 15 (7) The "use by" date for the drug, which shall be:
- 16 (A) The expiration date on the manufacturer's
- 17 container; or
- 18 (B) One year from the date the drug is dispensed,
- 19 whichever is earlier;
- 20 (8) The number of refills available, if any;
- 21 (9) In the case of the dispensing of an equivalent generic
- 22 drug product, the statement "same as (brand name of



1 the drug product prescribed or the referenced listed
2 drug name)", or words of similar meaning; and
3 (10) Specific directions for the drug's use; provided that
4 if the specific directions for use are too lengthy for
5 inclusion on the label, the notation "take according
6 to written instructions" may be used if separate
7 written instructions for use are actually issued with
8 the drug by the practitioner or the pharmacist, but in
9 no event shall the notation "take as directed",
10 referring to oral instructions, be considered
11 acceptable.

12 If any prescription for a drug does not indicate the number of
13 times it may be refilled, if any, the pharmacist shall not
14 refill that prescription unless subsequently authorized to do so
15 by the practitioner. The act of dispensing a prescription drug
16 other than a professional sample or medical oxygen contrary to
17 this subsection shall be deemed to be an act that results in a
18 drug being misbranded while held for sale.

19 (b) In addition to the requirements enumerated in
20 subsection (a), a prescription drug shall be dispensed only:

21 (1) By a pharmacist pursuant to a valid prescription [~~ex~~],
22 section 461-1[+], or section 453-B;



- 1 (2) By a medical oxygen distributor pursuant to a
- 2 prescription or certificate of medical necessity;
- 3 provided that the drug to be dispensed is medical
- 4 oxygen; or
- 5 (3) By a practitioner to an ultimate user; provided that:
- 6 (A) [The] Except as otherwise authorized for
- 7 expedited partner therapy in section 453-B, the
- 8 practitioner shall inform the patient, prior to
- 9 dispensing any drug other than a professional
- 10 sample, that the patient may have a written,
- 11 orally ordered, or electronically transmitted or
- 12 conveyed prescription directed to a pharmacy or a
- 13 medical oxygen distributor of the patient's own
- 14 choice;
- 15 (B) The practitioner shall promptly record in the
- 16 practitioner's records:
- 17 (i) The prescription in full;
- 18 (ii) The name, strength, and quantity of the
- 19 drug, and specific directions for the drug's
- 20 use;
- 21 (iii) The date the drug was dispensed; [and]



1 (iv) [The] Except as otherwise authorized for
2 expedited partner therapy in section 453-B,
3 the name and address of the person for whom
4 the drug was prescribed or the name of the
5 owner of the animal for which the drug was
6 prescribed; and

7 (v) Prescription drugs dispensed or prescribed
8 for expedited partner therapy as authorized
9 under section 453-B;

10 (C) The records described in subparagraph (B) shall
11 be subject to the inspection of the department or
12 its agents at all times; and

13 (D) No undisclosed rebate, refund, commission,
14 preference, discount, or other consideration,
15 whether in the form of money or otherwise, has
16 been offered to the practitioner as compensation
17 or inducement to dispense or prescribe any
18 specific drug in preference to other drugs that
19 might be used for the identical therapeutic
20 indication.



1 (c) A prescription may be communicated in writing, orally,
2 or by electronic transmission, and shall include the following
3 information:

4 (1) The authorization of the practitioner noted as
5 follows:

6 (A) Written prescriptions shall include the original
7 signature of the practitioner;

8 (B) Oral prescriptions shall be promptly recorded by
9 the pharmacist or medical oxygen distributor and
10 shall include the practitioner's oral code
11 designation; and

12 (C) Electronic prescriptions shall be irrefutably
13 traceable to the prescribing practitioner by a
14 recognizable and unique practitioner identifier
15 such as:

16 (i) A bitmap or graphic image of the
17 prescriber's handwritten signature and the
18 prescriber's oral code designation (or
19 license number or other identifier if the
20 prescriber is an out-of-state practitioner);

21 (ii) An electronic signature;

22 (iii) A digital signature; or



- 1 (iv) By other means as approved by the director;
- 2 (2) The date of issuance;
- 3 (3) The practitioner's name, business telephone number,
- 4 and business address, unless the practitioner is
- 5 otherwise uniquely identified and the pharmacy or
- 6 medical oxygen distributor dispensing the prescription
- 7 has the prescriber's contact information on file
- 8 accessible within the dispensing area;
- 9 (4) The name, strength, and quantity of the drug to be
- 10 dispensed, and specific directions for the drug's use;
- 11 (5) [The] Except as otherwise authorized for expedited
- 12 partner therapy in section 453-B, the name and address
- 13 of the person for whom the prescription was written or
- 14 the name of the owner of the animal for which the drug
- 15 was prescribed, unless the pharmacy or medical oxygen
- 16 distributor dispensing the prescription has the
- 17 address on file accessible within the dispensing area;
- 18 (6) The room number and route of administration, if the
- 19 patient is in an institutional facility; and
- 20 (7) The number of allowable refills, if the prescription
- 21 is refillable. If the number of refills authorized by
- 22 the practitioner is indicated using the terms "as



1 needed" or "prn", the prescription may be refilled up
2 to twelve months from the date the original
3 prescription was written. After the twelve-month
4 period, the "as needed" or "prn" prescription may be
5 refilled for a subsequent three-month period;
6 provided:

7 (A) The prescription is refilled only once during the
8 three-month period;

9 (B) The refill does not exceed a thirty-day supply of
10 the drug;

11 (C) The refill does not provide any amount of the
12 drug fifteen months beyond the date the original
13 prescription was written;

14 (D) In the case of medical oxygen, the duration of
15 therapy indicated on a certificate of medical
16 necessity shall supersede any limitations or
17 restrictions on refilling; and

18 (E) Subparagraphs (A) to (D) shall apply only to
19 pharmacies and medical oxygen distributors
20 practicing in the State."

21 2. By amending subsection (g) to read:



1 "(g) Any drug other than medical oxygen dispensed pursuant
2 to a prescription shall be exempt from the requirements of
3 section 328-15 (except paragraphs (1), (9), (11), and (12), and
4 the packaging requirements of paragraphs (7) and (8)), if the
5 drug bears a label containing:

- 6 (1) The name and address of the pharmacy;
- 7 (2) The serial number and the date of the prescription or
8 of its filling;
- 9 (3) The name of the practitioner;
- 10 (4) ~~[The]~~ Except as otherwise authorized for expedited
11 partner therapy in section 453-B, the name of the
12 patient;
- 13 (5) The directions for use; and
- 14 (6) Any cautionary statements contained in the
15 prescription.

16 This exemption shall not apply to any drug dispensed in the
17 course of the conduct of a business of dispensing drugs pursuant
18 to diagnosis by mail, or to a drug dispensed in violation of
19 subsection (a), (b), (c), or (d)."

20 SECTION 5. Section 328-17.6, Hawaii Revised Statutes, is
21 amended as follows:

22 1. By amending subsections (c) and (d) to read:



1 "(c) Any pharmacist or medical oxygen distributor who
2 fills or refills a prescription from an out-of-state
3 practitioner shall:

4 (1) Note the following on the prescription record: the
5 out-of-state practitioner's full name, address, and
6 telephone number;

7 (2) Be responsible for validating and verifying the
8 practitioner's prescriptive authority by virtue of a
9 valid out-of-state license, a Drug Enforcement
10 Administration registration number, or other measures
11 as appropriate; and

12 (3) ~~[Demand]~~ Except as otherwise authorized for expedited
13 partner therapy in section 453-B, demand proper
14 identification from the person whose name appears on
15 the prescription prior to filling the prescription, in
16 addition to complying with any identification
17 procedures established by the department for filling
18 and refilling an out-of-state prescription.

19 (d) Before refilling a transferred out-of-state
20 prescription, a pharmacist or medical oxygen distributor shall:

21 (1) ~~[Advise]~~ Except as otherwise authorized for expedited
22 partner therapy in section 453-B, advise the person



1 whose name appears on the prescription that the
2 prescription on file at the originating out-of-state
3 pharmacy or medical oxygen distributor may be
4 canceled; and

5 (2) Record all information required to be on a
6 prescription, including:

- 7 (A) The date of issuance of the original
- 8 prescription;
- 9 (B) The number of refills authorized on the original
- 10 prescription;
- 11 (C) The date the original prescription was dispensed;
- 12 (D) The number of valid refills remaining and the
- 13 date of the last refill;
- 14 (E) The out-of-state pharmacy's or out-of-state
- 15 medical oxygen distributor's name, telephone
- 16 number, and address, and the original
- 17 prescription number or control number from which
- 18 the prescription information was transferred; and
- 19 (F) The name of the transferor pharmacist or the
- 20 medical oxygen distributor's agent."

21 2. By amending subsection (f) to read:

1 "(f) An out-of-state prescription record shall state the
2 date of filling or refilling and, except as otherwise authorized
3 for expedited partner therapy in section 453-B, the local
4 address of the person whose name appears on the prescription."

5 SECTION 6. Section 328-17.7, Hawaii Revised Statutes, is
6 amended by amending subsection (a) to read as follows:

7 "(a) Every practitioner, pharmacist, or medical oxygen
8 distributor who compounds, sells, or delivers any prescribed
9 drug to a patient or a patient's agent shall maintain records
10 that identify:

- 11 (1) The specific drug product dispensed, including:
 - 12 (A) The product's national drug code (NDC) number; or
 - 13 (B) The brand name or the established name and the
14 name or commonly accepted abbreviation of the
15 principal labeler of the drug product dispensed,
16 the product strength, and the dosage form;
- 17 (2) The quantity of the drug;
- 18 (3) Directions for use;
- 19 (4) The number of allowable refills;
- 20 (5) The date of initial dispensing and the dates of all
21 refilling;
- 22 (6) The date of any transfer of the prescription;



- 1 (7) The name, business address, and telephone number of
- 2 the recipient pharmacist or medical oxygen distributor
- 3 for any transfer of prescription;
- 4 (8) The prescribing practitioner, including name, business
- 5 address, and telephone number;
- 6 (9) The format (oral, written, or electronic) in which the
- 7 prescription was received;
- 8 (10) [The] Except as otherwise authorized for expedited
- 9 partner therapy in section 453-B, the patient,
- 10 including name, address, and telephone number;
- 11 (11) The date of prescribing; and
- 12 (12) The name of the practitioner, pharmacist, or medical
- 13 oxygen distributor dispensing the drug.

14 Every prescription dispensed shall have the name of the
 15 pharmacist, dispensing practitioner, or medical oxygen
 16 distributor responsible for the dispensing appended to the
 17 prescription record, and every prescription record shall be
 18 preserved and legible for a period of not less than five years.
 19 The prescription records shall be subject at all times to the
 20 inspection of the director of health or the director's agent."

21 SECTION 7. In codifying the new sections added by section
 22 2 of this Act, the revisor of statutes shall substitute

1 appropriate section numbers for the letters used in designating
2 the new sections in this Act.

3 SECTION 8. Statutory material to be repealed is bracketed
4 and stricken. New statutory material is underscored.

5 SECTION 9. This Act shall take effect upon its approval.

APPROVED this 1 day of JUL, 2013



GOVERNOR OF THE STATE OF HAWAII