

**EXPEDITED PARTNER THERAPY TREATMENT**

2009 GENERAL SESSION

STATE OF UTAH

**Chief Sponsor: Jennifer M. Seelig**

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**LONG TITLE**

**General Description:**

This bill amends the Pharmacy Practice Act in the Division of Occupational and Professional Licensing Act.

**Highlighted Provisions:**

This bill:

- ▶ defines terms;
- ▶ excludes from the definition of unprofessional conduct and unlawful conduct under the Division of Occupational and Professional Licensing, issuing a prescription for an antibiotic to an unnamed partner of a person who has any one of certain designated sexually transmitted diseases;
  - ▶ does not mandate the use of expedited partner therapy;
  - ▶ provides an option for a practitioner to use expedited partner therapy;
  - ▶ makes conforming changes to the Pharmacy Practices Act;
  - ▶ provides immunity from medical malpractice actions for a practitioner who uses expedited partner therapy; and
- ▶ makes technical changes.

**Monies Appropriated in this Bill:**

None

31 **Other Special Clauses:**

32 None

33 **Utah Code Sections Affected:**

34 AMENDS:

35 **58-17b-602**, as last amended by Laws of Utah 2007, Chapter 279

36 ENACTS:

37 **58-1-501.3**, Utah Code Annotated 1953



39 *Be it enacted by the Legislature of the state of Utah:*

40 Section 1. Section **58-1-501.3** is enacted to read:

41 **58-1-501.3. Health professional prescribing exceptions for expedited partner**  
42 **therapy for sexually transmitted diseases.**

43 (1) For purposes of this section:

44 (a) "Drug to treat a sexually transmitted disease" means a drug:

45 (i) as defined in Section 58-17b-102; and

46 (ii) that is:

47 (A) an antibiotic; and

48 (B) prescribed in accordance with guidelines from the Centers for Disease Control and

49 Prevention for patient delivered expedited partner therapy in the management of sexually  
50 transmitted disease.

51 (b) "Partner" means a person:

52 (i) with whom a practitioner does not have a bonafide practitioner-patient relationship;

53 and

54 (ii) who is identified as, or claims to be a sexual partner of a patient.

55 (c) "Patient" means a person who:

56 (i) has a sexually transmitted disease; and

57 (ii) has a bonafide practitioner-patient relationship with a practitioner.

58 (d) "Sexually transmitted disease" means:

59 (i) gonorrhea; or

60 (ii) chlamydia.

61 (2) This section does not require a practitioner or a licensee under this chapter to  
62 prescribe or dispense a drug to treat a sexually transmitted disease for patient delivered  
63 expedited partner therapy. A practitioner's or licensee's decision to use expedited partner  
64 therapy as allowed by this section is voluntary.

65 (3) Notwithstanding Sections 58-1-501, 58-17b-501, and 58-17b-502, it is not  
66 unlawful conduct or unprofessional conduct, and it does not violate the provisions of this  
67 chapter if:

68 (a) a practitioner, in accordance with this Subsection (3):

69 (i) issues a prescription for a drug to treat a sexually transmitted disease to a partner  
70 by:

71 (A) writing "partner of (patient name)" on the prescription order; and

72 (B) giving the partner's prescription to the patient for subsequent use by the partner; or

73 (ii) notwithstanding Section 58-17b-610, dispenses a drug sample to treat a sexually  
74 transmitted disease to the patient for the subsequent use of the partner; or

75 (b) a pharmacist, in accordance with this Subsection (3), dispenses a prescription drug  
76 for the treatment of a sexually transmitted disease to:

77 (i) a person who:

78 (A) claims to be a partner; and

79 (B) presents a prescription for the drug to the pharmacist which is written for the  
80 unnamed partner of a named patient;

81 (ii) the patient for the subsequent use by the unnamed partner; or

82 (iii) an agent of the patient or partner.

83 (4) (a) For purposes of Subsection (3), and notwithstanding Section 58-17b-602:

84 (i) the partner does not have to be identified on the prescription order by information  
85 that would disclose the identity of the partner; and

86 (ii) when dispensing a drug to treat a sexually transmitted disease directly to the

87 partner, the patient's identifying information may, but does not need to, be included on the  
88 partner's drug label.

89 (b) Information provided by a pharmacist to a patient or the patient's agent for  
90 subsequent use by a partner satisfies the requirements of patient counseling for both the  
91 patient and the partner under Section 58-17b-613.

92 (5) (a) The Legislature finds that the prevention and treatment of sexually transmitted  
93 diseases in the state is a compelling public health issue.

94 (b) A practitioner or licensee under this chapter is not liable for a medical malpractice  
95 action if the use of expedited partner therapy is in compliance with this section, except for  
96 those acts which are grossly negligent or willful and wanton.

97 Section 2. Section **58-17b-602** is amended to read:

98 **58-17b-602. Prescription orders -- Information required -- Alteration -- Labels --**  
99 **Signatures -- Dispensing in pharmacies.**

100 (1) [~~The~~] Except as provided in Section 58-1-501.3, the minimum information that  
101 shall be included in a prescription order, and that may be defined by rule, is:

102 (a) the prescriber's name, address, and telephone number, and, if the order is for a  
103 controlled substance, the patient's age and the prescriber's DEA number;

104 (b) the patient's name and address or, in the case of an animal, the name of the owner  
105 and species of the animal;

106 (c) the date of issuance;

107 (d) the name of the medication or device prescribed and dispensing instructions, if  
108 necessary;

109 (e) the directions, if appropriate, for the use of the prescription by the patient or  
110 animal and any refill, special labeling, or other instructions;

111 (f) the prescriber's signature if the prescription order is written;

112 (g) if the order is an electronically transmitted prescription order, the prescribing  
113 practitioner's electronic signature; and

114 (h) if the order is a hard copy prescription order generated from electronic media, the

115 prescribing practitioner's electronic or manual signature.

116 (2) The requirement of Subsection (1)(a) does not apply to prescription orders  
117 dispensed for inpatients by hospital pharmacies if the prescriber is a current member of the  
118 hospital staff and the prescription order is on file in the patient's medical record.

119 (3) Unless it is for a Schedule II controlled substance, a prescription order may be  
120 dispensed by [~~pharmacists~~] a pharmacist or pharmacy [~~interns~~] intern upon an oral  
121 prescription of a practitioner only if the oral prescription is promptly reduced to writing.

122 (4) (a) Except as provided under Subsection (4)(b), a pharmacist or pharmacy intern  
123 may not dispense or compound any prescription of a practitioner if [~~it~~] the prescription shows  
124 evidence of alteration, erasure, or addition by any person other than the person writing the  
125 prescription.

126 (b) A pharmacist or pharmacy intern dispensing or compounding a prescription may  
127 alter or make additions to the prescription after receiving permission of the prescriber and may  
128 make entries or additions on the prescription required by law or necessitated in the  
129 compounding and dispensing procedures.

130 (5) Each drug dispensed shall have a label securely affixed to the container indicating  
131 the following minimum information:

- 132 (a) the name, address, and telephone number of the pharmacy;
- 133 (b) the serial number of the prescription as assigned by the dispensing pharmacy;
- 134 (c) the filling date of the prescription or its last dispensing date;
- 135 (d) the name of the patient, or in the case of an animal, the name of the owner and  
136 species of the animal;
- 137 (e) the name of the prescriber;
- 138 (f) the directions for use and cautionary statements, if any, which are contained in the  
139 prescription order or are needed;
- 140 (g) except as provided in Subsection (6), the trade, generic, or chemical name, amount  
141 dispensed and the strength of dosage form, but if multiple ingredient products with established  
142 proprietary or nonproprietary names are prescribed, those products' names may be used; and

143 (h) the beyond use date.

144 (6) If the prescriber specifically indicates the name of the prescription product should  
145 not appear on the label, then any of the trade, generic, chemical, established proprietary, and  
146 established nonproprietary names and the strength of dosage form may not be included.

147 (7) Except when it is delivered to the ultimate user via the United States Postal  
148 Service, licensed common carrier, or supportive personnel, a prescription drug may be  
149 dispensed to the ultimate user or his agent only at a licensed pharmacy.