## **Representative Eric K. Hutchings** proposes the following substitute bill:

1	DRUG PRODUCT EQUIVALENT AMENDMENTS
2	2007 GENERAL SESSION
3	STATE OF UTAH
4	Chief Sponsor: Eric K. Hutchings
5	Senate Sponsor: Sheldon L. Killpack
6 7	LONG TITLE
8	General Description:
9	This bill amends Pharmacy Practice Act provisions governing the substitution of a drug
10	product equivalent for a drug specified in a prescription order.
11	Highlighted Provisions:
12	This bill:
13	<ul><li>amends the Pharmacy Practice Act;</li></ul>
14	<ul> <li>except as specified, prohibits substitution of a drug product equivalent for a drug</li> </ul>
15	prescribed to treat or prevent seizures without the prescribing practitioner's
16	authorization; and
17	<ul> <li>clarifies that the preferred drug list and the generic requirements of the state</li> </ul>
18	Medicaid program supercede provisions related to substitutions for seizure
19	medications.
20	Monies Appropriated in this Bill:
21	None
22	Other Special Clauses:
23	None
24	<b>Utah Code Sections Affected:</b>
25	AMENDS:



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<b>58-17b-605</b> , as enacted by Chapter 280, Laws of Utah 2004
Be it enacted by the Legislature of the state of Utah:
Section 1. Section <b>58-17b-605</b> is amended to read:
58-17b-605. Drug product equivalents.
(1) A pharmacist or pharmacy intern dispensing a prescription order for a specific drug
by brand or proprietary name may substitute another drug product equivalent as defined in
Section 58-17b-102 for the prescribed drug only if:
(a) the purchaser specifically requests or consents to the substitution of a drug product
equivalent;
(b) the substituted drug product equivalent is of the same generic type and is
designated the therapeutic equivalent in the approved drug products with therapeutic
equivalence evaluations prepared by the Center for Drug Evaluation and Research of the
Federal Food and Drug Administration;
(c) the substituted drug product is permitted to move in interstate commerce;
(d) the pharmacist or pharmacy intern counsels the patient on the use and the expected
response to the prescribed drug, whether a substitute or not, and the substitution is not
otherwise prohibited by this chapter;
(e) the prescribing practitioner has not indicated that [an equivalent] a drug product [is]
equivalent may not [to] be substituted for the drug, as provided in Subsection (5); [and]
(f) except when the prescription drug is paid for in whole or in part by Medicaid, if the
prescribed drug has been prescribed in this instance to treat or prevent seizures, the pharmacist
or pharmacy intern obtains the authorization of the prescribing practitioner; and
[(f)] (g) the substitution is not otherwise prohibited by law, including any restrictions
imposed by the Medicaid program.
(2) (a) Each out-of-state mail service pharmacy dispensing a substituted drug product
equivalent as a substitute for another drug into this state shall notify the patient of the
substitution either by telephone or in writing.
(b) Each out-of-state mail service pharmacy shall comply with the requirements of this
chapter with respect to [drugs which may be] a drug product equivalent substituted for another

<u>drug</u>, including labeling [and], record keeping[, when dispensing substituted drug products],

and in accordance with Subsection (1)(f), prescribing practitioner authorization.

- (3) Pharmacists or pharmacy interns may not substitute without the prescriber's authorization on trade name drug product prescriptions unless the product is currently categorized in the approved drug products with therapeutic equivalence evaluations prepared by the Center for Drug Evaluation and Research of the Federal Food and Drug Administration as a drug product considered to be therapeutically equivalent to another drug product.
- (4) A pharmacist or pharmacy intern who dispenses a prescription with a drug product equivalent under this section assumes no greater liability than would be incurred had the pharmacist or pharmacy intern dispensed the prescription with the drug product prescribed.
- (5) (a) If, in the opinion of the <u>prescribing</u> practitioner, it is in the best interest of the patient that [an equivalent] a drug product <u>equivalent</u> not be substituted <u>for another drug</u>, the practitioner may indicate a prohibition on substitution either by writing "dispense as written" or may sign in the appropriate space where two lines have been preprinted on a prescription order and captioned "dispense as written" or "substitution permitted".
- (b) If the prescription is communicated orally by the <u>prescribing</u> practitioner to the pharmacist or pharmacy intern, the practitioner shall indicate the prohibition on substitution and that indication shall be noted in writing by the pharmacist or pharmacy intern with the name of the practitioner and the words "orally by" and the initials of the <u>pharmacist or pharmacy [practitioner] intern</u> written after it.
- (6) The substitution, if any, shall be communicated to the purchaser. The container shall be labeled with the name of the drug <u>product equivalent</u> dispensed and the pharmacist, pharmacy intern, or pharmacy technician shall indicate on the file copy of the prescription both the name of the prescribed drug and the name of the drug <u>product equivalent</u> dispensed in its place.
- (7) Failure of a licensed medical practitioner to specify that no substitution is authorized does not constitute evidence of negligence.

### H.B. 440 1st Sub. (Buff) - Drug Product Equivalent Amendments

# **Fiscal Note**

2007 General Session State of Utah

### **State Impact**

Enactment of this bill will not require additional appropriations.

### Individual, Business and/or Local Impact

Enactment of this bill likely will not result in direct, measurable costs and/or benefits for individuals, businesses, or local governments.

2/16/2007, 8:55:29 AM, Lead Analyst: Greer, W.

Office of the Legislative Fiscal Analyst