

Bill No. CS for SB 2220

Amendment No. \_\_\_\_

<u>Senate</u>	CHAMBER ACTION	<u>House</u>
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Senator Silver moved the following amendment to amendment (333964):

**Senate Amendment (with title amendment)**

On page 276, between lines 26 and 27,

insert:

Section 201. Subsection (6) of section 465.025, Florida Statutes, is amended to read:

465.025 Substitution of drugs.--

(6) The Board of Pharmacy and the Board of Medicine shall establish by rule a formulary of generic drug type and brand name drug products which have not been rated as therapeutically equivalent in "Approved Drug Products with Therapeutic Equivalence Evaluations" (Orange Book) published by the Federal Food and Drug Administration, and were not eligible for generic substitution for brand name drug products in at least 40 other states on January 1, 1999 ~~are determined by the boards to demonstrate clinically significant biological or therapeutic inequivalence~~ and which, if substituted, would pose a threat to the health and safety of patients receiving

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1 prescription medication.

2 (a) The formulary may be added to or deleted from as  
3 the Board of Pharmacy and the Board of Medicine deem  
4 appropriate. Any person who requests any inclusion, addition,  
5 or deletion of a generic drug type or brand name drug product  
6 to the formulary shall have the burden of proof to show cause  
7 why such inclusion, addition, or deletion should be made.

8 (b) Upon adoption of the formulary required by this  
9 subsection, and upon each addition, deletion, or modification  
10 to the formulary, the Board of Pharmacy shall mail a copy to  
11 each manager of the prescription department of each community  
12 pharmacy licensed by the state, each nonresident pharmacy  
13 registered in the state, and each board regulating  
14 practitioners licensed by the laws of the state to prescribe  
15 drugs shall incorporate such formulary into its rules. No  
16 pharmacist shall substitute a generically equivalent drug  
17 product for a prescribed brand name drug product if the brand  
18 name drug product or the generic drug type drug product is  
19 included in the said formulary, unless the generically  
20 equivalent drug product has been rated as therapeutically  
21 equivalent in the Orange Book.

22  
23 (Redesignate subsequent sections.)

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26 ===== T I T L E A M E N D M E N T =====

27 And the title is amended as follows:

28 On page 296, line 29, after the semicolon,

29  
30 insert:

31 amending s. 465.025, F.S.; revising

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1 requirements for the Board of Pharmacy and the  
2 Board of Medicine in adopting a formulary of  
3 generic drugs and brand name drugs;  
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