

By Senator Campbell

32-136-03

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A bill to be entitled  
An act relating to pharmacy; providing a short  
title; defining the term "pharmaceutical  
adverse incident" and requiring that such  
incidents be reported to the Department of  
Health; providing exceptions; requiring the  
department to review reported incidents to  
determine whether the incidents potentially  
involve conduct by a health care practitioner  
that is subject to disciplinary action;  
specifying that any disciplinary action shall  
be taken by the appropriate board; providing  
for the adoption of rules and forms; providing  
effective dates.

Be It Enacted by the Legislature of the State of Florida:

Section 1. This act may be cited as the "Ernest Belles  
Act."

Section 2. (1) As used in this section, the term  
"pharmaceutical adverse incident" means the dispensing of a  
different medication, a different dose, or the correct  
medication in a container with different instructions than  
those specified in the prescription, which dispensation  
results in actual harm to a patient, but does not include the  
dispensing of a generic equivalent medication with the  
patient's consent.

(2) A pharmacist licensed under chapter 465, Florida  
Statutes, or other health care practitioner as defined in  
section 456.001, Florida Statutes, who becomes aware of a  
patient's allegation that a pharmaceutical adverse incident

1 has occurred which was caused by a health care practitioner  
2 must report such allegation to the Department of Health on  
3 forms provided by the department. This section does not apply  
4 to:

5 (a) Pharmacists employed by pharmacies that  
6 participate in the program provided by Rule 64B16-27.300,  
7 Florida Administrative Code; or

8 (b) Pharmacists employed by pharmacies that have  
9 notified the Board of Pharmacy that they will establish a  
10 continuous quality-improvement program consistent with the  
11 requirements of Rule 64B16-27.300, Florida Administrative  
12 Code.

13 (3) The required notification to the department must  
14 be submitted in writing by certified mail and postmarked  
15 within 15 days after the pharmacist or health care  
16 practitioner became aware of the patient's allegation that a  
17 pharmaceutical adverse incident has occurred.

18 (4) Effective July 1, 2005, subject to subsequent act  
19 of the Legislature and a specific appropriation sufficient to  
20 cover the actual costs, the department shall review each  
21 incident and determine whether it potentially involved conduct  
22 by a pharmacist or health care practitioner who is subject to  
23 disciplinary action, in which case section 465.073, Florida  
24 Statutes, applies. Disciplinary action, if any, shall be  
25 taken by the board under which the pharmacist or health care  
26 practitioner is licensed.

27 (5) The Department of Health shall adopt forms and  
28 rules for administering this section.

29 Section 3. Section 2 of this act shall take effect  
30 only upon the effective date of legislation that makes any  
31 such information provided to the Department of Health

1 confidential and exempt from section 119.07(1), Florida  
2 Statutes, and Section 24(a) of Article I of the State  
3 Constitution, until 10 days after probable cause is found that  
4 a violation of law occurred. Such legislation must also  
5 provide that information may be used by the department or the  
6 Board of Pharmacy only in a disciplinary proceeding brought  
7 against the pharmacist or by the department in any study of  
8 adverse incidents without identifying the patient, pharmacist,  
9 pharmacy, office, or entity by name, location, or other  
10 identifier.

11 Section 4. Except as otherwise expressly provided in  
12 this act, this act shall take effect July 1, 2003.

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15 SENATE SUMMARY

16 Requires health care practitioners to report certain  
17 prescription-filling incidents to the Department of  
18 Health. Provides for the adoption of rules and forms.  
19 Defines the term "pharmaceutical adverse incident".  
20 Specifies that a pharmacist or other health care  
21 practitioner is required to report an allegation of a  
22 "pharmaceutical adverse incident" if he or she is aware  
23 of a patient allegation that such an incident was caused  
24 by the health care practitioner. Makes exceptions for  
25 which certain pharmacists may be excluded from reporting  
26 requirements.

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