

Bill No. CS for SB 518

Barcode 021838

CHAMBER ACTION

Senate

House

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Comm: RCS  
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The Committee on Governmental Operations (King) recommended  
the following **substitute for amendment** (180174):

**Senate Amendment (with title amendment)**

On page 6, line 13, through  
page 9, line 21, delete those lines

and insert:

Section 3. Section 408.0611, Florida Statutes, is  
created to read:

408.0611 Electronic prescribing clearinghouse.--  
(1) It is the intent of the Legislature to promote the  
implementation of electronic prescribing by health care  
practitioners, health care facilities, and pharmacies in order  
to prevent prescription drug abuse, improve patient safety,  
and reduce unnecessary prescriptions. To that end, it is the  
intent of the Legislature to create a clearinghouse of  
information on electronic prescribing to convey the process  
and advantages of electronic prescribing; to provide  
information regarding the availability of electronic  
prescribing products, including no-cost or low-cost products;

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1 and to regularly convene stakeholders to assess and accelerate  
2 the implementation of electronic prescribing.

3 (2) As used in this section, the term:

4 (a) "Electronic prescribing" means, at a minimum, the  
5 electronic review of the patient's medication history, the  
6 electronic generation of the patient's prescription, and the  
7 electronic transmission of the patient's prescription to a  
8 pharmacy.

9 (b) "Health care practitioner" means an individual  
10 authorized by law to prescribe drugs.

11 (3) The agency shall work in collaboration with  
12 private-sector electronic prescribing initiatives and relevant  
13 stakeholders to create a clearinghouse of information on  
14 electronic prescribing for health care practitioners, health  
15 care facilities, and pharmacies. These stakeholders shall  
16 include organizations that represent health care  
17 practitioners; organizations that represent health care  
18 facilities; organizations that represent pharmacies;  
19 organizations that operate electronic prescribing networks;  
20 organizations that create electronic prescribing products; and  
21 regional health information organizations. Specifically, the  
22 agency shall, by October 1, 2007:

23 (a) Provide on its website:

24 1. Information regarding the process of electronic  
25 prescribing and the availability of electronic prescribing  
26 products, including no-cost or low-cost products;

27 2. Information regarding the advantages of electronic  
28 prescribing, including using medication history data to  
29 prevent drug interactions, prevent allergic reactions, and  
30 deter doctor and pharmacy shopping for controlled substances;

31 3. Links to federal and private-sector websites that

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1 provide guidance on selecting an appropriate electronic  
2 prescribing product; and

3 4. Links to state, federal, and private-sector  
4 incentive programs for the implementation of electronic  
5 prescribing.

6 (b) Convene quarterly meetings of the stakeholders to  
7 assess and accelerate the implementation of electronic  
8 prescribing.

9 (4) Pursuant to s. 408.061, the agency shall monitor  
10 the implementation of electronic prescribing by health care  
11 practitioners, health care facilities, and pharmacies. By  
12 January 31 of each year, the agency shall report on the  
13 progress of implementation of electronic prescribing to the  
14 Governor and the Legislature. Information reported pursuant to  
15 this subsection shall include federal and private-sector  
16 electronic prescribing initiatives and, to the extent that  
17 data is readily available from organizations that operate  
18 electronic prescribing networks, the number of health care  
19 practitioners using electronic prescribing and the number of  
20 prescriptions electronically transmitted.

21 Section 4. Section 893.065, Florida Statutes, is  
22 created to read:

23 893.065 Counterfeit-resistant prescription blanks for  
24 controlled substances listed in Schedule II, Schedule III, or  
25 Schedule IV.--The Department of Health shall develop and adopt  
26 by rule the form and content for a counterfeit-resistant  
27 prescription blank which may be used by practitioners for the  
28 purpose of prescribing a controlled substance listed in  
29 Schedule II, Schedule III, or Schedule IV. The Department of  
30 Health may require the prescription blanks to be printed on  
31 distinctive, watermarked paper and to bear the preprinted

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1 name, address, and category of professional licensure of the  
 2 practitioner and that practitioner's federal registry number  
 3 for controlled substances. The prescription blanks may not be  
 4 transferred.

5 Section 5. The penalties created in s. 831.311(2),  
 6 Florida Statutes, by this act shall be effective only upon the  
 7 adoption of the rules required pursuant to s. 893.065, Florida  
 8 Statutes, as created by this act.

9 Section 6. If a person dies of an apparent drug  
 10 overdose:

11 (1) A law enforcement agency shall prepare a report  
 12 identifying each prescribed controlled substance listed in  
 13 Schedule II, Schedule III, or Schedule IV of s. 893.03,  
 14 Florida Statutes, which is found on or near the deceased or  
 15 among the deceased's possessions. The report must identify the  
 16 person who prescribed the controlled substance, if known or  
 17 ascertainable. Thereafter, the law enforcement agency shall  
 18 submit a copy of the report to the medical examiner.

19 (2) A medical examiner who is preparing a report  
 20 pursuant to s. 406.11, Florida Statutes, shall include in the  
 21 report information identifying each prescribed controlled  
 22 substance listed in Schedule II, Schedule III, or Schedule IV  
 23 of s. 893.03, Florida Statutes, that was found in, on, or near  
 24 the deceased or among the deceased's possessions.

25 Section 7. The sum of \$100,000 in nonrecurring general  
 26 revenue is appropriated to the Agency for Health Care  
 27 Administration to implement this act.

28 Section 8. This act shall take effect July 1, 2007.  
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1 ===== T I T L E    A M E N D M E N T =====

2 And the title is amended as follows:

3            On page 1, line 17, through  
4            page 2, line 6, delete those lines

5

6 and insert:

7            prescription; creating s. 408.0611, F.S.;  
8            providing legislative intent; providing  
9            definitions; requiring the Agency for Health  
10            Care Administration to create a clearinghouse  
11            of information on electronic prescribing;  
12            requiring the agency to monitor and report on  
13            the implementation of electronic prescribing;  
14            creating s. 893.065, F.S.; requiring the  
15            department to develop and adopt by rule the  
16            form and content for a counterfeit-proof  
17            prescription blank for voluntary use by  
18            physicians in prescribing a controlled  
19            substance listed in Schedule II, Schedule III,  
20            or Schedule IV; providing that penalties shall  
21            become effective only upon adoption of rules;  
22            prescribing duties of law enforcement agencies  
23            and medical examiners when a person dies of an  
24            apparent drug overdose; providing an  
25            appropriation; providing an effective date.

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