Bill No. <u>CS for SB 270</u>

	CHAMBER ACTION <u>Senate</u> <u>House</u>							
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11	Senator Saunders moved the following amendment:							
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13	Senate Amendment (with title amendment)							
14	On page 2, line 2,							
15								
16	insert:							
17	Section 1. Section 831.311, Florida Statutes, is							
18	created to read:							
19	831.311 Unlawful sale, manufacture, alteration,							
20	delivery, uttering, or possession of counterfeit-resistant							
21	prescription blanks for controlled substances							
22	(1) It is unlawful for any person having the intent to							
23	injure or defraud any person or to facilitate any violation of							
24	s. 893.13 to sell, manufacture, alter, deliver, utter, or							
25	possess with intent to injure or defraud any person, or to							
26	facilitate any violation of s. 893.13, any							
27	counterfeit-resistant prescription blanks for controlled							
28	substances, the form and content of which are adopted by rule							
29	of the Department of Health pursuant to s. 893.065.							
30	(2) Any person who violates this section commits a							
31	felony of the third degree, punishable as provided in s. 1							
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1 775.082, s. 775.083, or s. 775.084. Section 2. Section 893.04, Florida Statutes, is 2 amended to read: 3 4 893.04 Pharmacist and practitioner.--(1) A pharmacist, in good faith and in the course of 5 б professional practice only, may dispense controlled substances 7 upon a written or oral prescription of a practitioner, under the following conditions: 8 9 (a) Oral prescriptions must be promptly reduced to writing by the pharmacist or recorded electronically if 10 11 permitted by federal law. (b) The written prescription must be dated and signed 12 13 by the prescribing practitioner on the day when issued. (c) There shall appear on the face of the prescription 14 15 or written record thereof for the controlled substance the 16 following information: 1. The full name and address of the person for whom, 17 or the owner of the animal for which, the controlled substance 18 19 is dispensed. 20 2. The full name and address of the prescribing practitioner and the practitioner's federal controlled 21 22 substance registry number shall be printed thereon. 3. If the prescription is for an animal, the species 23 24 of animal for which the controlled substance is prescribed. 4. The name of the controlled substance prescribed and 25 the strength, quantity, and directions for use thereof. 26 5. The number of the prescription, as recorded in the 27 prescription files of the pharmacy in which it is filled. 28 29 6. The initials of the pharmacist filling the prescription and the date filled. 30 31 (d) The prescription shall be retained on file by the 2 4:17 PM 04/30/07 s0270c1c-37-r8g

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1 proprietor of the pharmacy in which it is filled for a period of 2 years. 2 (e) Affixed to the original container in which a 3 4 controlled substance is delivered upon a prescription or authorized refill thereof, as hereinafter provided, there 5 shall be a label bearing the following information: 6 7 1. The name and address of the pharmacy from which such controlled substance was dispensed. 8 9 2. The date on which the prescription for such 10 controlled substance was filled. 3. The number of such prescription, as recorded in the 11 prescription files of the pharmacy in which it is filled. 12 13 4. The name of the prescribing practitioner. 5. The name of the patient for whom, or of the owner 14 15 and species of the animal for which, the controlled substance 16 is prescribed. 6. The directions for the use of the controlled 17 18 substance prescribed in the prescription. 19 7. A clear, concise warning that it is a crime to transfer the controlled substance to any person other than the 20 21 patient for whom prescribed. 22 (f) A prescription for a controlled substance listed in Schedule II may be dispensed only upon a written 23 24 prescription of a practitioner, except that in an emergency situation, as defined by regulation of the Department of 25 Health, such controlled substance may be dispensed upon oral 26 prescription but is limited to a 72-hour supply. A No 27 prescription for a controlled substance listed in Schedule II 28 may not be refilled. 29 (g) \underline{A} No prescription for a controlled substance 30 31 listed in <u>Schedule</u> Schedules III, <u>Schedule</u> IV, or <u>Schedule</u> V 3 4:17 PM 04/30/07 s0270c1c-37-r8g

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1	may <u>not</u> be filled or refilled more than five times within a					
2	period of 6 months after the date on which the prescription					
3	was written unless the prescription is renewed by a					
4	practitioner.					
5	(2)(a) A pharmacist may not dispense a controlled					
6	substance listed in Schedule II, Schedule III, or Schedule IV					
7	to any patient or patient's agent without first determining,					
8	in the exercise of her or his professional judgment, that the					
9	order is valid. The pharmacist may dispense the controlled					
10	substance, in the exercise of her or his professional					
11	judgment, when the pharmacist or pharmacist's agent has					
12	obtained satisfactory patient information from the patient or					
13	the patient's agent.					
14	(b) Any pharmacist who dispenses by mail a controlled					
15	substance listed in Schedule II, Schedule III, or Schedule IV					
16	is exempt from the requirement to obtain suitable					
17	identification for the prescription dispensed by mail if the					
18	pharmacist has obtained the patient's identification through					
19	the patient's prescription benefit plan.					
20	(c) Any controlled substance listed in Schedule III or					
21	Schedule IV may be dispensed by a pharmacist upon an oral					
22	prescription if, before filling the prescription, the					
23	pharmacist reduces it to writing or records the prescription					
24	electronically if permitted by federal law. Such prescriptions					
25	must contain the date of the oral authorization.					
26	(d) Each written prescription prescribed by a					
27	practitioner in this state for a controlled substance listed					
28	in Schedule II, Schedule III, or Schedule IV must include both					
29	a written and a numerical notation of the quantity on the face					
30	of the prescription and a notation of the date, with the					
31	abbreviated month written out on the face of the prescription.					
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1 A pharmacist may, upon verification by the prescriber, document any information required by this paragraph. 2 (e) A pharmacist may not dispense more than a 30-day 3 4 supply of a controlled substance listed in Schedule III upon an oral prescription issued in this state. 5 б (f) A pharmacist may not knowingly fill a prescription 7 that has been forged for a controlled substance listed in Schedule II, Schedule III, or Schedule IV. 8 (3)(2) Notwithstanding the provisions of subsection 9 10 (1), a pharmacist may dispense a one-time emergency refill of 11 up to a 72-hour supply of the prescribed medication for any medicinal drug other than a medicinal drug listed in Schedule 12 II, in compliance with the provisions of s. 465.0275. 13 (4)(3) The legal owner of any stock of controlled 14 15 substances in a pharmacy, upon discontinuance of dealing in 16 controlled substances, may sell said stock to a manufacturer, wholesaler, or pharmacy. Such controlled substances may be 17 sold only upon an order form, when such an order form is 18 19 required for sale by the drug abuse laws of the United States 20 or this state, or regulations pursuant thereto. 21 Section 3. Section 408.0611, Florida Statutes, is 22 created to read: 408.0611 Electronic prescribing clearinghouse.--23 2.4 (1) It is the intent of the Legislature to promote the implementation of electronic prescribing by health care 25 practitioners, health care facilities, and pharmacies in order 2.6 to prevent prescription drug abuse, improve patient safety, 27 and reduce unnecessary prescriptions. To that end, it is the 28 29 intent of the Legislature to create a clearinghouse of information on electronic prescribing to convey the process 30 31 and advantages of electronic prescribing; to provide 5 s0270c1c-37-r8g 4:17 PM 04/30/07

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1	information regarding the availability of electronic					
2	prescribing products, including no-cost or low-cost products;					
3	and to regularly convene stakeholders to assess and accelerate					
4	the implementation of electronic prescribing.					
5	(2) As used in this section, the term:					
б	(a) "Electronic prescribing" means, at a minimum, the					
7	electronic review of the patient's medication history, the					
8	electronic generation of the patient's prescription, and the					
9	electronic transmission of the patient's prescription to a					
10	pharmacy.					
11	(b) "Health care practitioner" means an individual					
12	authorized by law to prescribe drugs.					
13	(3) The agency shall work in collaboration with					
14	private-sector electronic prescribing initiatives and relevant					
15	stakeholders to create a clearinghouse of information on					
16	electronic prescribing for health care practitioners, health					
17	care facilities, and pharmacies. These stakeholders shall					
18	include organizations that represent health care					
19	practitioners; organizations that represent health care					
20	facilities; organizations that represent pharmacies;					
21	organizations that operate electronic prescribing networks;					
22	organizations that create electronic prescribing products; and					
23	regional health information organizations. Specifically, the					
24	agency shall, by October 1, 2007:					
25	(a) Provide on its website:					
26	1. Information regarding the process of electronic					
27	prescribing and the availability of electronic prescribing					
28	products, including no-cost or low-cost products;					
29	2. Information regarding the advantages of electronic					
30	prescribing, including using medication history data to					
31	prevent drug interactions, prevent allergic reactions, and					
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1	deter doctor and pharmacy shopping for controlled substances;					
2	3. Links to federal and private-sector websites that					
3	provide guidance on selecting an appropriate electronic					
4	prescribing product; and					
5	4. Links to state, federal, and private-sector					
6	incentive programs for the implementation of electronic					
7	prescribing.					
8	(b) Convene quarterly meetings of the stakeholders to					
9	assess and accelerate the implementation of electronic					
10	prescribing.					
11	(4) Pursuant to s. 408.061, the agency shall monitor					
12	the implementation of electronic prescribing by health care					
13	practitioners, health care facilities, and pharmacies. By					
14	January 31 of each year, the agency shall report on the					
15	progress of implementation of electronic prescribing to the					
16	Governor and the Legislature. Information reported pursuant to					
17	this subsection shall include federal and private-sector					
18	electronic prescribing initiatives and, to the extent that					
19	data is readily available from organizations that operate					
20	electronic prescribing networks, the number of health care					
21	practitioners using electronic prescribing and the number of					
22	prescriptions electronically transmitted.					
23	Section 4. Section 893.065, Florida Statutes, is					
24	created to read:					
25	893.065 Counterfeit-resistant prescription blanks for					
26	controlled substances listed in Schedule II, Schedule III, or					
27	Schedule IVThe Department of Health shall develop and adopt					
28	by rule the form and content for a counterfeit-resistant					
29	prescription blank which may be used by practitioners for the					
30	purpose of prescribing a controlled substance listed in					
31	Schedule II, Schedule III, or Schedule IV. The Department of					
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1	Health may require the prescription blanks to be printed on					
2	distinctive, watermarked paper and to bear the preprinted					
3	name, address, and category of professional licensure of the					
4	practitioner and that practitioner's federal registry number					
5	for controlled substances. The prescription blanks may not be					
б	transferred.					
7	Section 5. The penalties created in s. 831.311(2),					
8	Florida Statutes, by this act shall be effective only upon the					
9	adoption of the rules required pursuant to s. 893.065, Florida					
10	Statutes, as created by this act.					
11	Section 6. If a person dies of an apparent drug					
12	<u>overdose:</u>					
13	(1) A law enforcement agency shall prepare a report					
14	identifying each prescribed controlled substance listed in					
15	Schedule II, Schedule III, or Schedule IV of s. 893.03,					
16	Florida Statutes, which is found on or near the deceased or					
17	among the deceased's possessions. The report must identify the					
18	person who prescribed the controlled substance, if known or					
19	ascertainable. Thereafter, the law enforcement agency shall					
20	submit a copy of the report to the medical examiner.					
21	(2) A medical examiner who is preparing a report					
22	pursuant to s. 406.11, Florida Statutes, shall include in the					
23	report information identifying each prescribed controlled					
24	substance listed in Schedule II, Schedule III, or Schedule IV					
25	of s. 893.03, Florida Statutes, that was found in, on, or near					
26	the deceased or among the deceased's possessions.					
27						
28	(Redesignate subsequent sections.)					
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31						
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Florida Senate - 2007 SENATOR AMENDMENT Bill No. CS for SB 270 Barcode 371292 1 ========== T I T L E And the title is amended as follows: 2 On page 1, line 2, after the semicolon, 3 4 5 insert: б creating s. 831.311, F.S.; prohibiting the 7 sale, manufacture, alteration, delivery, uttering, or possession of 8 9 counterfeit-resistant prescription blanks for 10 controlled substances with the intent to injure 11 or defraud; providing penalties; amending s. 893.04, F.S.; providing additional requirements 12 13 for the dispensing of a controlled substance listed in Schedule II, Schedule III, or 14 15 Schedule IV; specifying circumstances under 16 which a pharmacist who dispenses controlled substances by mail is exempt from certain 17 requirements governing patient identification; 18 19 providing requirements and limitations for 20 dispensing controlled substances upon an oral 21 prescription; creating s. 408.0611, F.S.; 22 providing legislative intent; providing definitions; requiring the Agency for Health 23 2.4 Care Administration to create a clearinghouse of information on electronic prescribing; 25 requiring the agency to monitor and report on 26 the implementation of electronic prescribing; 27 creating s. 893.065, F.S.; requiring the 28 29 department to develop and adopt by rule the form and content for a counterfeit-proof 30 31 prescription blank for voluntary use by 4:17 PM 04/30/07 s0270c1c-37-r8g

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1		physicians in prescrib	ing a controlled	
2		substance listed in Sc	hedule II, Schedu	le III,
3		or Schedule IV; provid	ing that penaltie	s shall
4		become effective only	upon adoption of	rules;
5		prescribing duties of	law enforcement a	gencies
6		and medical examiners	when a person die	s of an
7		apparent drug overdose	;	
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