

1-1 By: Burton S.B. No. 1243
 1-2 (In the Senate - Filed March 11, 2015; March 17, 2015, read
 1-3 first time and referred to Committee on Health and Human Services;
 1-4 April 16, 2015, reported adversely, with favorable Committee
 1-5 Substitute by the following vote: Yeas 9, Nays 0; April 16, 2015,
 1-6 sent to printer.)

1-7 COMMITTEE VOTE

	Yea	Nay	Absent	PNV
1-8				
1-9	X			
1-10	X			
1-11	X			
1-12	X			
1-13	X			
1-14	X			
1-15	X			
1-16	X			
1-17	X			

1-18 COMMITTEE SUBSTITUTE FOR S.B. No. 1243 By: Taylor of Collin

1-19 A BILL TO BE ENTITLED
 1-20 AN ACT

1-21 relating to donation of unused prescription drugs.
 1-22 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF TEXAS:
 1-23 SECTION 1. Subtitle A, Title 6, Health and Safety Code, is
 1-24 amended by adding Chapter 442 to read as follows:
 1-25 CHAPTER 442. DONATION OF PRESCRIPTION DRUGS
 1-26 SUBCHAPTER A. GENERAL PROVISIONS
 1-27 Sec. 442.001. DEFINITIONS. In this chapter:
 1-28 (1) "Department" means the Department of State Health
 1-29 Services.
 1-30 (2) "Donor" means an individual who donates unused
 1-31 prescription drugs under this chapter to a participating provider.
 1-32 (3) "Executive commissioner" means the executive
 1-33 commissioner of the Health and Human Services Commission.
 1-34 (4) "Health care facility" means:
 1-35 (A) a general or special hospital as defined by
 1-36 Chapter 241;
 1-37 (B) an ambulatory surgical center licensed under
 1-38 Chapter 243;
 1-39 (C) an institution licensed under Chapter 242; or
 1-40 (D) any other facility that provides health care
 1-41 services to patients and is authorized to maintain an inventory of
 1-42 prescription drugs for dispensing to the facility's patients or
 1-43 residents.
 1-44 (5) "Health care professional" means an individual
 1-45 licensed, certified, or otherwise authorized to administer health
 1-46 care and prescribe prescription drugs, for profit or otherwise, in
 1-47 the ordinary course of business or professional practice. The term
 1-48 does not include a health care facility.
 1-49 (6) "Participating provider" means a health care
 1-50 facility, pharmacy, or health care professional who elects to
 1-51 participate in the collection and redistribution of donated
 1-52 prescription drugs under this chapter.
 1-53 (7) "Pharmacy" means an entity licensed under Chapter
 1-54 560, Occupations Code.
 1-55 (8) "Prescription drug" has the meaning assigned by
 1-56 Section 551.003, Occupations Code.
 1-57 (9) "Recipient" means an individual who voluntarily
 1-58 receives donated prescription drugs under this chapter.
 1-59 (10) "Tamper-evident" means packaging that allows for
 1-60 detection of unauthorized access to a prescription drug.

2-1 Sec. 442.002. RULEMAKING AUTHORITY. The executive
2-2 commissioner may adopt rules to implement this chapter.

2-3 Sec. 442.003. CONSTRUCTION WITH OTHER LAW. This chapter
2-4 does not limit the authority of this state or a political
2-5 subdivision of this state to regulate or prohibit a prescription
2-6 drug.

2-7 SUBCHAPTER B. DONATION AND REDISTRIBUTION OF UNUSED PRESCRIPTION
2-8 DRUGS

2-9 Sec. 442.051. DONATION AND REDISTRIBUTION OF PRESCRIPTION
2-10 DRUGS. (a) A donor may donate unused prescription drugs to a
2-11 participating provider in accordance with this chapter and rules
2-12 adopted under this chapter.

2-13 (b) A participating provider may dispense donated
2-14 prescription drugs to a recipient in accordance with this chapter
2-15 and rules adopted under this chapter.

2-16 Sec. 442.052. STANDARDS FOR DONATION AND REDISTRIBUTION.
2-17 (a) The executive commissioner by rule shall adopt standards and
2-18 procedures for:

2-19 (1) accepting, storing, labeling, and dispensing
2-20 donated prescription drugs; and

2-21 (2) inspecting donated prescription drugs to
2-22 determine whether the drugs are adulterated and whether the drugs
2-23 are safe and suitable for redistribution.

2-24 (b) In adopting standards and procedures under this
2-25 section, the executive commissioner shall ensure that the donation
2-26 and redistribution process is consistent with public health and
2-27 safety standards.

2-28 Sec. 442.053. REQUIREMENTS FOR DONATED PRESCRIPTION DRUGS.

2-29 (a) A donated prescription drug may be accepted or dispensed under
2-30 this chapter only if the drug is in its original, unopened, sealed,
2-31 and tamper-evident unit-dose packaging. A drug packaged in single
2-32 unit doses may be accepted and dispensed if the outside packaging is
2-33 opened but the single unit-dose packaging is unopened.

2-34 (b) A donated prescription drug may not be accepted or
2-35 dispensed under this chapter if the drug:

2-36 (1) is a controlled substance;

2-37 (2) is adulterated or misbranded; or

2-38 (3) is not stored in compliance with the drug's product
2-39 label.

2-40 (c) A participating provider shall comply with all
2-41 applicable provisions of state and federal law relating to the
2-42 inspection, storage, labeling, and dispensing of prescription
2-43 drugs.

2-44 Sec. 442.054. DONATION PROCESS. (a) Before being
2-45 dispensed to a recipient, a prescription drug donated under this
2-46 chapter must be inspected by a health care professional on behalf of
2-47 the participating provider in accordance with federal law, laws of
2-48 this state, and department rule to determine whether the drug is
2-49 adulterated or misbranded and whether the drug has been stored in
2-50 compliance with the requirements of the product label.

2-51 (b) A donated prescription drug dispensed to a recipient
2-52 under this chapter must be prescribed by a health care professional
2-53 for use by the recipient.

2-54 (c) A participating provider may charge a handling fee not
2-55 to exceed \$20 to a recipient to cover the costs of inspecting,
2-56 storing, labeling, and dispensing the donated prescription drug. A
2-57 participating provider may not resell a prescription drug donated
2-58 under this chapter. A donor may not sell a prescription drug to a
2-59 participating provider.

2-60 (d) A participating provider may not submit a claim or
2-61 otherwise seek reimbursement from any public or private third-party
2-62 payor for donated prescription drugs dispensed to a recipient under
2-63 this chapter. A public or private third-party payor is not required
2-64 to provide reimbursement for donated drugs dispensed to a recipient
2-65 under this chapter.

2-66 Sec. 442.055. DONOR FORM. Before donating a prescription
2-67 drug under this chapter, a donor shall sign a form prescribed by the
2-68 department stating that:

2-69 (1) the donor is the owner of the donated prescription

3-1 drug;
 3-2 (2) the donated prescription drug has been properly
 3-3 stored and the container has not been opened or tampered with;
 3-4 (3) the donated prescription drug has not been
 3-5 adulterated or misbranded; and
 3-6 (4) the donor is voluntarily donating the prescription
 3-7 drug.

3-8 Sec. 442.056. RECIPIENT FORM. Before accepting a donated
 3-9 prescription drug under this chapter, a recipient shall sign a form
 3-10 prescribed by the department stating that:

3-11 (1) the recipient acknowledges that the donor is not a
 3-12 pharmacist and the donor took ordinary care of the prescription
 3-13 drug;

3-14 (2) the recipient acknowledges that the donor is known
 3-15 to the participating provider and that there is no reason to believe
 3-16 that the prescription drug was improperly handled or stored;

3-17 (3) by accepting the prescription drug, the recipient
 3-18 accepts any risk that an accidental mishandling could create; and

3-19 (4) the recipient releases the donor, participating
 3-20 provider, and manufacturer of the drug from liability related to
 3-21 the prescription drug.

3-22 Sec. 442.057. LIMITATION OF LIABILITY. (a) A donor or
 3-23 participating provider who acts in good faith in donating,
 3-24 accepting, storing, labeling, distributing, or dispensing
 3-25 prescription drugs under this chapter:

3-26 (1) is not criminally liable and is not subject to
 3-27 professional disciplinary action for those activities; and

3-28 (2) is not civilly liable for damages for bodily
 3-29 injury, death, or property damage that arises from those activities
 3-30 unless the injury, death, or damage arises from the donor or
 3-31 participating provider's recklessness or intentional conduct.

3-32 (b) A manufacturer of a prescription drug donated under this
 3-33 chapter is not liable for bodily injury, death, or property damage
 3-34 arising from a donor or participating provider's failure to
 3-35 properly handle or store the drug. This subsection does not limit
 3-36 the liability of the manufacturer for a dangerous or defective
 3-37 drug.

3-38 Sec. 442.058. DATABASE OF PARTICIPATING PROVIDERS. The
 3-39 department shall establish and maintain an electronic database that
 3-40 lists each participating provider. The department shall post the
 3-41 database on its Internet website.

3-42 SECTION 2. Not later than December 1, 2015, the executive
 3-43 commissioner of the Health and Human Services Commission shall
 3-44 adopt any necessary rules for the implementation of Chapter 442,
 3-45 Health and Safety Code, as added by this Act.

3-46 SECTION 3. If before implementing any provision of this Act
 3-47 a state agency determines that a waiver or authorization from a
 3-48 federal agency is necessary for implementation of that provision,
 3-49 the agency affected by the provision shall request the waiver or
 3-50 authorization and may delay implementing that provision until the
 3-51 waiver or authorization is granted.

3-52 SECTION 4. This Act takes effect September 1, 2015.

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