

1 SB113
2 125792-3
3 By Senator Orr
4 RFD: Finance and Taxation General Fund
5 First Read: 01-MAR-11

1 SB113

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4 ENROLLED, An Act,

5 To authorize the Alabama Department of Corrections
6 to accept and redispense unused prescription medications.

7 BE IT ENACTED BY THE LEGISLATURE OF ALABAMA:

8 Section 1. As used in this act, the following terms
9 shall have the following meanings:

10 (1) CORRECTIONS FACILITY. Any facility or program
11 controlled or operated by the state Department of Corrections
12 or any of its agencies or departments and supported wholly or
13 in part by state funds for the correctional care of persons.

14 (2) CUSTOMIZED PATIENT MEDICATION PACKAGE. A package
15 that is prepared by a pharmacist for a specific patient and
16 that contains two or more prescribed solid oral dosage forms.

17 (3) REPACKAGING. The process by which the pharmacy
18 prepares a prescription it accepts pursuant to this act in a
19 unit-dose package, unit-of-issue package or customized patient
20 medication package for immediate dispensing in accordance with
21 a current prescription.

22 (4) UNIT-DOSE PACKAGE. A package that contains a
23 single-dose drug with the name, strength, control number, and
24 expiration date of that drug on the label.

1 (5) UNIT-OF-ISSUE PACKAGE. A package that provides
2 multiple doses of the same drug, but each drug is individually
3 separated and includes the name, lot number, and expiration
4 date of the drug.

5 Section 2. (a) A pharmacy operated by the Alabama
6 Department of Corrections (ADOC) or operated by a company
7 under contract with the ADOC, shall accept for the purpose of
8 redispensing a prescription drug that has been dispensed and
9 has left the control of the pharmacy or pharmacist if the
10 prescription drug is being returned by a corrections facility
11 that has met the requirements of routine on-site inspections
12 by the pharmacy or pharmacist and has a registered
13 professional nurse or a licensed practical nurse who is
14 responsible for the security, handling, and administration of
15 prescription drugs within that corrections facility and if all
16 of the following conditions are met:

17 (1) The pharmacy or pharmacist is satisfied that the
18 conditions under which the prescription drug has been
19 delivered, stored, and handled before and during its return
20 were such as to prevent damage, deterioration, or
21 contamination that would adversely affect the identity,
22 strength, quality, purity, stability, integrity, or
23 effectiveness of the prescription drug.

24 (2) The pharmacist is satisfied that the
25 prescription drug did not leave the control of the registered

1 professional nurse or licensed practical nurse responsible for
2 the security, handling, and administration of that
3 prescription drug and that the prescription drug did not come
4 into the physical possession of the individual for whom it was
5 prescribed.

6 (3) The pharmacist is satisfied that the labeling
7 and packaging of the prescription drug are accurate, have not
8 been altered, defaced, or tampered with and include the
9 identity, strength, expiration date, and lot number of the
10 prescription drug.

11 (4) The prescription drug was dispensed in a
12 unit-dose package or unit-of-issue package.

13 (b) A pharmacy operated by the ADOC or operated by a
14 company under contract with the ADOC shall not accept for
15 return prescription drugs as provided pursuant to this section
16 until the pharmacist in charge develops a written set of
17 protocols for accepting, returning to stock, repackaging,
18 labeling, and redispensing prescription drugs. The written
19 protocols shall be maintained on the premises of any pharmacy
20 dispensing prescriptions for the ADOC and shall be readily
21 accessible to each pharmacist on duty. The written protocols
22 shall include, at a minimum, each of the following:

23 (1) Methods for ensuring that damage, deterioration,
24 or contamination has not occurred during the delivery,
25 handling, storage, or return of the prescription drugs such

1 that it would adversely affect the identity, strength,
2 quality, purity, stability, integrity, or effectiveness of the
3 prescription drugs or otherwise render the drugs unfit for
4 distribution.

5 (2) Methods for accepting, returning to stock,
6 repackaging, labeling, and redispensing the prescription drugs
7 returned pursuant to this section.

8 (3) A uniform system of recording and tracking
9 prescription drugs that are returned to stock, repackaged,
10 labeled, and redistributed pursuant to this section.

11 (c) If the condition of a prescription drug and its
12 package meets the standards set forth in subsection (b), a
13 prescription drug shall be returned to stock and redistributed
14 as follows:

15 (1) A prescription drug that was originally
16 dispensed in the manufacturer's unit-dose package or
17 unit-of-issue package that is returned in that same package
18 may be returned to stock, repackaged, and redispensed as
19 needed.

20 (2) A prescription drug that is repackaged into a
21 unit-dose package or a unit-of-issue package by the pharmacy,
22 dispensed and returned to that pharmacy in that unit-dose
23 package or unit-of-issue package may be returned to stock, but
24 it shall not be repackaged. A unit-dose package or
25 unit-of-issue package prepared by the pharmacist and returned

1 to stock shall only be redispensed in that same unit-dose
2 package or unit-of-issue package and shall only be redispensed
3 once. A pharmacist shall not add unit-dose package drugs to a
4 partially used unit-of-issue package.

5 (d) This section does not apply to any of the
6 following:

7 (1) A controlled substance.

8 (2) A prescription drug that is dispensed as part of
9 a customized patient medication package.

10 (3) A prescription drug that is not dispensed as a
11 unit-dose package or a unit-of-issue package.

12 (4) A prescription drug that is not properly labeled
13 with the identity, strength, lot number, and expiration date.

14 Section 3. This act shall become effective
15 immediately following its passage and approval by the
16 Governor, or its otherwise becoming law.

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President and Presiding Officer of the Senate

Speaker of the House of Representatives

SB113
Senate 24-MAR-11
I hereby certify that the within Act originated in and passed
the Senate, as amended.

Patrick Harris
Secretary

House of Representatives
Passed: 09-JUN-11

By: Senator Orr