

House Bill 546

By: Representatives Jasperse of the 11th, Stephens of the 164th, and Parrish of the 158th

A BILL TO BE ENTITLED
AN ACT

1 To amend Code Section 26-4-5 of the Official Code of Georgia Annotated, relating to
2 definitions relative to the "Georgia Pharmacy Practice Act," so as to revise the definition of
3 "pharmacy care"; to provide for related matters; to repeal conflicting laws; and for other
4 purposes.

5 BE IT ENACTED BY THE GENERAL ASSEMBLY OF GEORGIA:

6 style="text-align:center">**SECTION 1.**

7 Code Section 26-4-5 of the Official Code of Georgia Annotated, relating to definitions
8 relative to the "Georgia Pharmacy Practice Act," is amended by revising paragraph (31) as
9 follows:

10 "(31) 'Pharmacy care' means:

11 (A) Those services related to the interpretation, evaluation, or dispensing of
12 prescription drug orders, the participation in drug and device selection, drug
13 administration, and drug regimen reviews, and the provision of patient counseling
14 related thereto; ~~and~~

15 (B) Ordering and administering:

16 (i) Tests that have been cleared or approved ~~for home use~~ under a certificate of
 17 waiver by the federal Food and Drug Administration pursuant to the federal Clinical
 18 Laboratory Improvement Amendments of 1998 and interpreting the results as a means
 19 to screen for or monitor disease, disease risk factors, or drug use and to facilitate
 20 patient education; and

21 (ii) Viral and serology COVID-19 tests, provided that such authority shall expire 12
 22 months after the end of the public health emergency declared by the United States
 23 secretary of health and human services on January 31, 2020; ~~provided, however, that~~
 24 ~~such expiration shall not apply to viral and serology COVID-19 tests cleared or~~
 25 ~~approved pursuant to division (i) of this subparagraph.~~

26 A pharmacist conducting such a test shall do so at a pharmacy or other facility that has
 27 obtained any necessary certification from or that is operating under a certificate of
 28 waiver from the federal Centers for Medicare and Medicaid Services pursuant to the
 29 federal Clinical Laboratory Improvement Amendments of 1998; and

30 (C) Adaptation of a prescription drug order to:

31 (i) Change the quantity of medication prescribed if:

32 (I) The prescribed quantity or package size is not commercially available;

33 (II) The change in quantity is related to a change in dosage form, strength, or
 34 therapeutic interchange;

35 (III) The change is intended to dispense up to a 90 day supply of maintenance
 36 medication in accordance with subsection (q) of Code Section 26-4-80; or

37 (IV) The change extends a maintenance medication for the limited quantity
 38 necessary to coordinate a patient's refills in a medication synchronization program
 39 in accordance with Code Section 33-24-59.22;

40 (ii) Change the dosage form of the prescription if it is in the best interest of patient
 41 care, so long as the prescriber's directions are also modified to equate to an equivalent
 42 amount of drug dispensed as prescribed; or

43 (iii) Complete missing information on a prescription drug order if there is evidence
44 to support the change.

45 A pharmacist who adapts a prescription drug order in accordance with this
46 subparagraph shall: (1) make such adaptation in the exercise of his or her professional
47 judgment; (2) document such adaptation in the patient's record; and (3) obtain the
48 patient's consent prior to making such adaptation. Notwithstanding anything in this
49 chapter to the contrary, a pharmacist shall not make any changes to a prescription drug
50 order pursuant to this subparagraph if the prescriber indicates on such order that
51 adaptation is not permitted."

52 **SECTION 2.**

53 All laws and parts of laws in conflict with this Act are repealed.