

STATE OF NEW YORK

1428

2025-2026 Regular Sessions

IN ASSEMBLY

January 9, 2025

Introduced by M. of A. FORREST -- read once and referred to the Committee on Insurance

AN ACT to amend the insurance law, in relation to substituting brand name prescription drugs in the case of a drug shortage

The People of the State of New York, represented in Senate and Assembly, do enact as follows:

1 Section 1. Section 3242 of the insurance law is amended by adding a
2 new subsection (d) to read as follows:

3 (d) (1) As used in this subsection:

4 (A) "Eligible prescription drug" means a prescription drug approved
5 under 21 U.S.C. 355(c) that is not under patent.

6 (B) "Generic drug" means a drug that is approved pursuant to an appli-
7 cation referencing an eligible prescription drug that is submitted under
8 subsection (j) of Section 505 of the Federal Food, Drug, and Cosmetic
9 Act, 21 U.S.C. 355.

10 (C) "Supply issue" means a drug shortage or meaningful disruption as
11 defined in 21 U.S.C. 356(c).

12 (2) In the event an AB-rated generic equivalent or interchangeable
13 biological product for an eligible prescription drug is covered in an
14 insurer's formulary and such generic drug equivalent is unavailable due
15 to a supply issue which has been recognized by the federal food and drug
16 administration pursuant to 21 U.S.C. 356e and the dosage cannot be
17 adjusted, an insurer that delivers or issues for delivery in this state
18 a policy that provides coverage for prescription drugs shall provide
19 coverage for a brand name eligible prescription drug to an insured who
20 is already receiving such prescription drug as a generic equivalent or
21 has been diagnosed with or presented with a condition on or prior to the
22 start of the plan year that is treated by such eligible prescription
23 drug or is an eligible prescription drug that is or would be part of the
24 insured's treatment regimen for such condition. Such brand name eligible
25 prescription drug shall be covered at the same level of coverage as the

EXPLANATION--Matter in italics (underscored) is new; matter in brackets
[-] is old law to be omitted.

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1 generic drug in the insurer's formulary until such time as the supply
2 issue is resolved and the drug has been removed from the federal food
3 and drug administration's shortage list.

4 § 2. Section 4329 of the insurance law is amended by adding a new
5 subsection (d) to read as follows:

6 (d) (1) As used in this subsection:

7 (A) "Eligible prescription drug" means a prescription drug approved
8 under 21 U.S.C. 355(c) that is not under patent.

9 (B) "Generic drug" means a drug that is approved pursuant to an appli-
10 cation referencing an eligible prescription drug that is submitted under
11 subsection (j) of Section 505 of the Federal Food, Drug, and Cosmetic
12 Act, 21 U.S.C. 355.

13 (C) "Supply issue" means a drug shortage or meaningful disruption as
14 defined in 21 U.S.C. 356(c).

15 (2) In the event an AB-rated generic equivalent or interchangeable
16 biological product for an eligible prescription drug is covered in an
17 insurer's formulary and such generic drug equivalent is unavailable due
18 to a supply issue which has been recognized by the federal food and drug
19 administration pursuant to 21 U.S.C. 356e and the dosage cannot be
20 adjusted, a corporation subject to the provisions of this article that
21 issues a contract that provides coverage for prescription drugs shall
22 provide coverage for a brand name eligible prescription drug to an
23 insured who is already receiving such prescription drug as a generic
24 equivalent or has been diagnosed with or presented with a condition on
25 or prior to the start of the plan year that is treated by such eligible
26 prescription drug or is an eligible prescription drug that is or would
27 be part of the insured's treatment regimen for such condition. Such
28 brand name eligible prescription drug shall be covered at the same level
29 of coverage as the generic drug in the insurer's formulary until such
30 time as the supply issue is resolved and the drug has been removed from
31 the federal food and drug administration's shortage list.

32 § 3. This act shall take effect immediately.