

1 STATE OF OKLAHOMA

2 1st Session of the 58th Legislature (2021)

3 SENATE BILL 468

By: Hicks

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6 AS INTRODUCED

7 An Act relating to health insurance; prohibiting a
8 health insurer from modifying coverage of
9 prescription drug in certain circumstance; construing
10 clause; providing for certain civil penalty;
11 providing for codification; and providing an
12 effective date.

13 BE IT ENACTED BY THE PEOPLE OF THE STATE OF OKLAHOMA:

14 SECTION 1. NEW LAW A new section of law to be codified
15 in the Oklahoma Statutes as Section 6850.2 of Title 36, unless there
16 is created a duplication in numbering, reads as follows:

17 A. An insurer, as defined in Section 6054 of Title 36 of the
18 Oklahoma Statutes, shall not modify an insured's coverage of a
19 prescription drug, as defined in Section 367.2 of Title 59 of the
20 Oklahoma Statutes, if the following conditions are met:

21 1. The prescription drug had been previously preauthorized for
22 coverage by the insurer or was listed on the formulary of the
23 insurer at the time the insured was prescribed the drug by his or
24 her practitioner, as defined in Section 6054 of Title 36 of the
25 Oklahoma Statutes;

1 2. The insured has already received the prescription drug; and

2 3. A practitioner continues to prescribe the drug to the
3 insured.

4 Modification prohibited pursuant to this section shall include,
5 but is not limited to, raising the premium, copayment, coinsurance
6 or deductible, denying or otherwise failing to provide continued
7 coverage of the prescription drug, moving the drug to a more
8 restrictive coverage category or tier, or replacing a brand-name
9 drug for a generic drug after the insured has qualified for the
10 brand-name drug pursuant to this section.

11 B. Nothing in this section shall be construed to prohibit an
12 insurer from modifying coverage of a prescription drug if:

13 1. The federal Food and Drug Administration has issued a
14 statement calling into question the clinical safety of the drug; or

15 2. The manufacturer of the drug has notified the federal Food
16 and Drug Administration of a manufacturing discontinuance or
17 potential discontinuance of the drug as required by 21 U.S.C. 356c.

18 C. Any insurer that violates the provisions of this section
19 shall be subject to a civil penalty in an amount to be determined by
20 the Insurance Commissioner.

21 SECTION 2. This act shall become effective November 1, 2021.

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