

By: Senator(s) DeLano

To: Insurance

SENATE BILL NO. 2783

1 AN ACT TO PROHIBIT HEALTH INSURANCE PLANS FROM MODIFYING, ON  
2 RENEWAL, AN INSURED'S CONTRACTED BENEFIT LEVEL FOR ANY  
3 PRESCRIPTION DRUG THAT WAS APPROVED OR COVERED UNDER THE PLAN IN  
4 THE IMMEDIATELY PRECEDING PLAN YEAR AND PRESCRIBED DURING THAT  
5 YEAR FOR A MENTAL ILLNESS; TO LIST MODIFICATIONS PROHIBITED; TO  
6 CLARIFY WHAT IS NOT PROHIBITED; AND FOR RELATED PURPOSES.

7 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MISSISSIPPI:

8 **SECTION 1.** (1) All individual and group health insurance  
9 policies providing coverage on an expense incurred basis,  
10 individual and group service or indemnity type contracts issued by  
11 a nonprofit corporation, individual and group service contracts  
12 issued by a health maintenance organization, all self-insured  
13 group arrangements to the extent not preempted by federal law and  
14 all managed health care delivery entities of any type or  
15 description that are delivered, issued for delivery, continued or  
16 renewed on or after July 1, 2024, and providing coverage to any  
17 resident of this state may not modify, on renewal of the policy,  
18 plan or contract, an insured's contracted benefit level for any  
19 prescription drug that was approved or covered under the plan in  
20 the immediately preceding plan year and prescribed during that



21 year for a mental illness or psychiatric condition if the insured  
22 (a) was covered by the policy, plan or contract on the date  
23 immediately preceding the renewal date, (b) a physician or other  
24 prescribing provider prescribes the drug for the mental illness or  
25 psychiatric condition; and (c) the physician or other prescribing  
26 provider in consultation with the insured determines that the drug  
27 is the most appropriate course of treatment.

28 (2) Modifications prohibited under subsection (1) of this  
29 section include:

30 (a) Removing a drug from a formulary;

31 (b) Adding a requirement that an enrollee receive prior  
32 authorization for a drug;

33 (c) Imposing or altering a quantity limit for a drug;

34 (d) Imposing a step-therapy restriction for a drug;

35 (e) Moving a drug to a higher cost-sharing tier;

36 (f) Increasing a coinsurance, copayment, deductible, or  
37 other out-of-pocket expense that an enrollee must pay for a drug;

38 and

39 (g) Reducing the maximum drug coverage amount.

40 (3) This section shall not be construed to prohibit a  
41 policy, plan or contract issuer from removing a drug from its  
42 formulary or denying an insured's coverage for the drug if:

43 (a) The United States Food and Drug Administration has  
44 issued a statement about the drug that calls into question the  
45 clinical safety of the drug;



46                   (b) The drug manufacturer has notified the United  
47 States Food and Drug Administration of a manufacturing  
48 discontinuance or potential discontinuance of the drug as required  
49 by Section 506C, Federal Food, Drug, and Cosmetic Act (21 USC  
50 Section 356c); or

51                   (c) The drug manufacturer has removed the drug from the  
52 market.

53                   **SECTION 2.** This act shall take effect and be in force from  
54 and after July 1, 2024.

