

**House File 626 - Reprinted**

HOUSE FILE 626  
BY COMMITTEE ON COMMERCE

(SUCCESSOR TO HF 96)

(As Amended and Passed by the House March 9, 2023)

**A BILL FOR**

1 An Act relating to continuity of care and nonmedical switching  
2 by health carriers, health benefit plans, and utilization  
3 review organizations, and including applicability  
4 provisions.

5 BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF IOWA:

1 Section 1. NEW SECTION. 514F.9 Continuity of care —  
2 nonmedical switching.

3 1. *Definitions.* For the purpose of this section:

4 a. "*Commissioner*" means the commissioner of insurance.

5 b. "*Cost sharing*" means any coverage limit, copayment,  
6 coinsurance, deductible, or other out-of-pocket expense  
7 requirement.

8 c. "*Covered person*" means the same as defined in section  
9 514J.102.

10 d. "*Demonstrated bioavailability*" means the same as defined  
11 in section 155A.3.

12 e. "*Formulary*" means a complete list of prescription drugs  
13 eligible for coverage under a health benefit plan.

14 f. "*Generic name*" means the same as defined in section  
15 155A.3.

16 g. "*Health benefit plan*" means the same as defined in  
17 section 514J.102.

18 h. "*Health care professional*" means the same as defined in  
19 section 514J.102.

20 i. "*Health care services*" means the same as defined in  
21 section 514J.102.

22 j. "*Health carrier*" means an entity subject to the  
23 insurance laws and regulations of this state, or subject  
24 to the jurisdiction of the commissioner, including an  
25 insurance company offering sickness and accident plans, a  
26 health maintenance organization, a nonprofit health service  
27 corporation, a plan established pursuant to chapter 509A  
28 for public employees, or any other entity providing a plan  
29 of health insurance, health care benefits, or health care  
30 services. "*Health carrier*" does not include the department  
31 of human services, or a managed care organization acting  
32 pursuant to a contract with the department of human services to  
33 administer the medical assistance program under chapter 249A  
34 or the healthy and well kids in Iowa (hawk-i) program under  
35 chapter 514I.

1     *k. "Interchangeable biological product"* means the same as  
2 defined in section 155A.3.

3     1. *"Utilization review organization"* means the same as  
4 defined in section 514F.7.

5     2. *Nonmedical switching.* With respect to a health carrier  
6 that has entered into a health benefit plan with a covered  
7 person that covers prescription drug benefits, all of the  
8 following apply:

9     *a.* A health carrier, health benefit plan, or utilization  
10 review organization shall not limit or exclude coverage of  
11 a prescription drug for any covered person who is medically  
12 stable on such drug as determined by the prescribing health  
13 care professional, if all of the following apply:

14     (1) The prescription drug was previously approved by the  
15 health carrier for coverage for the covered person.

16     (2) The covered person's prescribing health care  
17 professional has prescribed the drug for the covered person's  
18 medical condition within the previous six months.

19     (3) The covered person continues to be an enrollee of the  
20 health benefit plan.

21     *b.* Coverage of a covered person's prescription drug, as  
22 described in paragraph "a", shall continue through the last day  
23 of the covered person's eligibility under the health benefit  
24 plan, or through the last day of the health benefit plan year,  
25 whichever is earlier.

26     *c.* Prohibited limitations and exclusions referred to in  
27 paragraph "a" include but are not limited to the following:

28     (1) Limiting or reducing the maximum coverage of  
29 prescription drug benefits.

30     (2) Increasing cost sharing for a covered prescription  
31 drug.

32     (3) Moving a prescription drug to a more restrictive tier if  
33 the health carrier uses a formulary with tiers.

34     (4) Removing a prescription drug from a formulary, unless  
35 the United States food and drug administration has issued a

1 statement about the drug that calls into question the clinical  
2 safety of the drug, or the manufacturer of the drug has  
3 notified the United States food and drug administration of a  
4 manufacturing discontinuance or potential discontinuance of the  
5 drug as required by section 506C of the Federal Food, Drug, and  
6 Cosmetic Act, as codified in 21 U.S.C. §356c.

7 *d.* This subsection shall not be construed to prohibit  
8 a substitution, a formulary change, or a preference by a  
9 health carrier for a prescribed drug product that has the same  
10 generic name and demonstrated bioavailability, or that is an  
11 interchangeable biological product.

12 3. *Limitations.* This section shall not be construed to do  
13 any of the following:

14 *a.* Prevent a health care professional from prescribing  
15 another drug covered by the health carrier that the health care  
16 professional deems medically necessary for the covered person.

17 *b.* Prevent a health carrier from doing any of the following:

18 (1) Adding a prescription drug to its formulary.

19 (2) Removing a prescription drug from its formulary if the  
20 drug manufacturer has removed the drug for sale in the United  
21 States.

22 4. *Enforcement.* The commissioner may take any enforcement  
23 action under the commissioner's authority to enforce compliance  
24 with this section.

25 Sec. 2. APPLICABILITY. This Act applies to a health benefit  
26 plan that is delivered, issued for delivery, continued, or  
27 renewed in this state on or after January 1, 2024.