Sixty-second Legislative Assembly of North Dakota

## HOUSE BILL NO. 1422

Introduced by

Representatives Weisz, Devlin, Kilichowski

Senators Dever, Uglem, Heckaman

1 A BILL for an Act to create and enact chapter 43-15.4 of the North Dakota Century Code,

2 relating to electronic prescription transmission.

## 3 BE IT ENACTED BY THE LEGISLATIVE ASSEMBLY OF NORTH DAKOTA:

SECTION 1. Chapter 43-15.4 of the North Dakota Century Code is created and enacted as
follows:

## 6 <u>43-15.4-01. Application.</u>

7 <u>This chapter applies to all electronic prescribing devices used within this state and to all</u>

8 software and hardware vendors and content managers with respect to such electronic

9 prescribing devices regardless of location. Relevant sections of this chapter also apply to all

10 requirements for prior authorization requests used within this state, and to all software and

11 hardware vendors and content managers with respect to electronic prior authorization requests,

12 <u>regardless of location.</u>

## 13 43-15.4-02. Electronic prescribing transmission standards.

14 <u>All prescription drug orders communicated by way of electronic transmission to a pharmacy</u>

15 or pharmacist must identify the transmitter's telephone number or any other suitable means to

16 contact the transmitter for verbal confirmation, written confirmation, or verbal and written

17 confirmation; the time and date of transmission; the identity of the pharmacy intended to receive

18 the transmission; and any other information required by federal or state law. An electronic

19 transmission is deemed the original prescription drug order, if the electronic transmission meets

- 20 the requirements of this section.
- 21 <u>43-15.4-03. Electronic transmission devices.</u>

22 <u>Electronic transmission devices used to communicate a prescription to a pharmacist must:</u>

23 <u>1.</u> <u>Allow any legal prescription to be written and entered into the device without</u>

24 interference or limitations before submission to a pharmacist.

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1	<u>2.</u>	Allow the prescription to be written through a neutral and open platform that does not			
2		use any means, program, or device, including advertising, instant messaging, and			
3		popup messaging, to influence or attempt to influence, through economic incentives or			
4		otherwise, the prescribing decision of an authorized prescriber at the point of care.			
5		This subsection applies if such means, program, or device is triggered by, initiated by,			
6		or is in specific response to the input, selection, or act or any combination of these of a			
7		prescribing health care professional or that prescribing health care professional's			
8		agent prescribing a covered outpatient drug or selecting a pharmacy for a patient.			
9	<u>3.</u>	Make available information regarding a plan's specific formulary according to the			
10		following conditions:			
11		a. All available covered outpatient drugs shall be readily disclosed to the authorized			
12		prescriber:			
13		b. All available pharmacies, both in and out of network, must be readily disclosed to			
14		the authorized prescriber;			
15		c. Nothing is designed to preclude or make more difficult the authorized prescriber's			
16		or patient's selection of any particular pharmacy or covered outpatient drug;			
17		d. Copay and cost-sharing data, specific to the patient's relevant formulary and			
18		entitled benefits, are electronically accessible to the physician for reference; and			
19		e. An electronic prior authorization process for allowing approval of an exception to			
20		the plan formulary or other restriction is available on the device as required under			
21		section 43-15.4-04, providing real-time adjudication.			
22	<u>4.</u>	As provided under subsection 2, alerts and messages to the prescriber and the			
23		prescriber's staff which are related to the formulary must support better clinical			
24		decisionmaking, including alerts to adverse events and access to formulary			
25		information. These messages and alerts must be consistently supported by scientific			
26		evidence. This information must be:			
27		a. Consistent with the federal food and drug administration regulations for			
28		advertising pharmaceutical products and be categorized or prioritized based on			
29		their clinical importance, including severity and likelihood of any adverse events;			
30		b. Individually suppressible by the prescriber;			

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1		<u>C.</u>	Able to be overridden by the prescriber so that the prescriber can prescribe the			
2			prescriber's medication of choice for the patient; and			
3		<u>d.</u>	Provide access to the decision support rules underlying each alert or message to			
4			include the date of the last update and the source of any financial support			
5			received in connection with the development of those rules.			
6	6 <u>43-15.4-04. Electronic prior authorization.</u>					
7	An electronic prior authorization process for allowing approval of an exception to the plan					
8	formulary or other restriction must:					
9	<u>1.</u>	<u>Be i</u>	required as a part of all electronic medical record systems that facilitate electronic			
10		<u>sub</u>	mission of prescriptions;			
11	<u>2.</u>	<u>Utili</u>	ze a universal format for a prior authorization request;			
12	<u>3.</u>	Pro	vide specific feedback to the provider on acceptable and approvable reasons for			
13		<u>app</u>	roval of a prior authorization request for a medication prescribed for a patient; and			
14	<u>4.</u>	Pro	vide real-time adjudication of the prior authorization request which facilitates an			
15		<u>exp</u>	lanation of benefits for the patient with information on how to appeal the denial of			
16		<u>the</u>	requested medication.			