GENERAL ASSEMBLY OF NORTH CAROLINA SESSION 2021

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HOUSE BILL 178 PROPOSED COMMITTEE SUBSTITUTE H178-PCS10483-BC-9

	Short Title:	Access to Prescription Drug Cost Information. (Public)					
	Sponsors:						
_	Referred to:						
		March 1, 2021					
1 2 3	AN ACT TO R INFORMA	A BILL TO BE ENTITLED EQUIRE ACCESS TO ACCURATE PRESCRIPTION DRUG BENEFIT COST TION.					
4		ssembly of North Carolina enacts:					
5	SEC	CTION 1. Chapter 58 of the General Statutes is amended by adding a new Article					
6	to read:						
7		" <u>Article 56B.</u>					
8		"Access to Prescription Drug Benefit Cost Information.					
9	" <u>§ 58-56B-1. 1</u>						
10		ing definitions apply in this Article:					
11	<u>(1)</u>	<u>Coverage. – The drug formulary information for a health benefit plan that</u>					
12		includes the brand and generic prescription drugs that the payor will cover for					
13	(2)	a specific patient under the patient's health benefit plan.					
14	<u>(2)</u>	<u>Dispenser. – Anyone licensed to dispense prescription drugs under the laws</u>					
15 16	(2)	of this State.					
10	<u>(3)</u>	<u>Health care services. – A health or medical care procedure or service rendered</u> by a health care provider or prescriber that does at least one of the following:					
18		<u>a.</u> <u>Provides testing, diagnosis, or treatment of a human disease or</u>					
19		dysfunction.					
20		b. <u>Dispenses drugs, medical devices, medical appliances, or medical</u>					
21		goods for the treatment of a human disease or dysfunction.					
22	<u>(4)</u>	Intermediary. – Any entity, including real-time networks and translation					
23	<u></u>	services, that accepts an electronic transaction from another organization and					
24		electronically routes the transaction to a receiving entity or facilitates the					
25		routing of prescription drug benefit transactions.					
26	<u>(5)</u>	Patient-specific eligibility information Information on the status of the					
27		health benefit plan and the prescription benefit available under a health benefit					
28		plan provided to a specific patient by a payor, including any exclusions and					
29		limitations under the health benefit plan and the prescription drug benefit					
30		under the health benefit plan.					
31	<u>(6)</u>	Patient-specific prescription drug benefit and cost information The type of					
32		prescription drug coverage offered to a patient by the patient's payor and any					
33		out-of-pocket costs that may be incurred by the patient under the coverage,					
34		including the patient's copayment, coinsurance, and deductible.					
35	<u>(7)</u>	Payor. – Any of the following:					



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	General Assemb	oly Of N	orth Carolina	Session 2021
1		<u>a.</u>	An insurer or nonprofit health service	plan that provides hospital,
2			medical, prescription drug, or surgical	benefits to individuals or
3			groups on an expense-incurred basis und	ler health insurance policies
4			or contracts that are issued or delivered in	<u>the State.</u>
5		<u>b.</u>	A health maintenance organization that p	
6			surgical benefits or prescribes drugs to	
7			contracts that are issued or delivered in th	
8	<u>(8)</u>		acy benefits manager. – As defined in G.S	
9	<u>(9)</u>		ber A licensed health care profession	onal authorized by law to
10		_	be a prescription drug.	
11	<u>(10)</u>		er. – Any person or facility that is licensed	or authorized in this State to
12		-	e health care services.	
13	<u>(11)</u>		me. – Exchange of patient eligibility, pr	-
14			als for a choice product and pharmacy an	
15			ions and alternatives when they exist. The	
16 17			iately after product selection using electro	onic prescribing platforms or
17 18	(12)	system Standa		at door all of the following:
18 19	<u>(12)</u>		rd transaction. – Any electronic process th Facilitates interoperability and data exc	
20		<u>a.</u>	benefit and investigation response inform	
20 21		<u>b.</u>	Is developed by an organization accredite	
21		<u>U.</u>	Standards Institute.	the American National
22	(13)	Switch	. – Has the same meaning as the term "inte	ermediary "
23 24	$\frac{(13)}{(14)}$		eutically equivalent alternative. – Any pr	
25	<u>(11)</u>		following:	esemption and that abos an
26		<u>a.</u>	Has the same clinical effect and safety pr	ofile to another prescription
27		—	drug prescribed for a patient.	<u> </u>
28		<u>b.</u>	Is known to have nearly identical prope	rties to another prescription
29			drug prescribed for a patient.	* *
30		<u>c.</u>	Uses real-time prescription benefit st	
31			organization accredited by the American	National Standards Institute.
32	" <u>§ 58-56B-5. Fi</u> 1			
33			y of North Carolina makes the following f	-
34	<u>(1)</u>		is a need for clear and meaningful	· ·
35			pocket prescription drug costs for pat	
36			riate, data-driven shared decision makin	
37		-	ed and understand the full range of optio	ons to obtain their medically
38			ary medications.	
39 40	<u>(2)</u>		s need to understand the opportunity to de	
40			plan formularies and understand coverage	
41 42			gs on those formularies, including lower-	cost chinear and therapeutic
42 43	(2)	<u>alterna</u>		a hanafit from compatitive
43 44	<u>(3)</u>		<u>s need to understand the opportunity to</u> of prescription drugs outside their healt	-
44 45			prmulary, whether in the form of a lower c	
46			idation programs.	asii price, patient assistance,
40 47	"8 58-56R-10 A		prescription drug benefit and cost info	rmation.
48			plans, pharmacy benefits managers, or an	
49			electronically provide to (i) any point of	
50			spensing of a prescription drug, or (iii) a	
	<u></u>	01 01	representation and and and and and and and and and an	

	General Assembly Of North Carolina Session 202	21			
1	benefit tool the minimum information described in G.S. 58-56B-15(c) to inform pati				
2	prescription price transparency and patients' access to their prescribed medications.				
3	(b) Payors, providers, pharmacies, and other organizations involved in the process of	of			
4	prescribing, dispensing, paying for, and exchanging information relating to prescription drug				
5	including intermediaries, real-time networks, switches, and translation services shall take an	-			
6	actions necessary to facilitate the creation of, access to, and use of the technology described				
7	subsection (a) of this section.	<u> </u>			
8	(c) Patient prescription price transparency technology shall not be prohibited fro	m			
9	displaying patient financial and resource assistance when that information is available for the				
10	prescription drug selected by a provider.				
11	" <u>§ 58-56B-15. Real-time requirements.</u>				
12	(a) Requests for patient-specific drug benefit and cost information through the formation the format	he			
12	technology required under G.S. 58-56B-10 and any responses to those requests using the				
13	technology shall be sent and received in real time.	<u>1at</u>			
15	(b) The real-time exchange of patient-specific eligibility information, including an	nv			
16	information related to a health benefit plan's coverage, benefits, formulary, and cost-sharin				
17	requirements, shall be facilitated using health care industry standards developed by a				
18	organization accredited by the American National Standards Institute.	un			
19	(c) Electronic health records shall display, through real-time integration, the mo	net			
20	up-to-date patient-specific eligibility information, including information on a health benefit				
20	plan's coverage, benefits, formulary, cost-sharing requirements, therapeutically equivale				
22	alternatives, and prior authorization requirements.	<u>/111</u>			
23	(d) <u>Electronic health record vendors, payors, providers, prescribers, pharmacies, and providers, pharmacies, and providers, pharmacies, pharmacies, and pharmacies, pharm</u>	nd			
24	other organizations involved in the process of prescribing, dispensing, paying for, an				
25	exchanging information relating to prescription drugs shall partner with intermediaries to ensu				
26	the delivery of accurate patient-specific prescription price transparency information.	10			
27	(e) Intermediaries shall be capable of supporting and using a standard transaction th	nat			
28	meets the requirements of this section.	iui			
29	(f) Patient-specific information, as described in G.S. 58-56B-15(c), shall be provided	in			
30	real time.				
31	"§ 58-56B-20. Benefit and cost information requirements.				
32	(a) Nothing in this Article shall interfere with patient choice and a health ca	ire			
33	professional's ability to convey the full range of prescription drug cost options to a patient. Heal				
34	benefit plans, pharmacy benefit managers, or any entities acting on behalf of a health benefit pla				
35	shall not restrict a health care professional from communicating prescription cost options to				
36	patient.				
37	(b) A payor shall not prohibit the display of patient-specific prescription drug benefit ar	nd			
38	cost information at the point of prescribing that reflects options available for covering the cost				
39	a prescription drug other than what may be available under the patient's health benefit pla				
40	including cash-pay options, coverage through assistance or support programs, and cost coverage				
41	options at the patient's pharmacy of choice.	<u> </u>			
42	(c) A provider or prescriber shall communicate to a patient the most therapeutical	lly			
43	appropriate treatment for the patient's diagnosis and, when appropriate, prescription drug co				
44	information, including the cash price, therapeutically equivalent alternatives, and delive				
45	options for a prescription drug.	-			
46	(d) In order to protect a patient's privacy and right to choose the means of prescription	on			
47	drug cost coverage, if a patient chooses not to use the prescription drug benefit under the patient				
48	health benefit plan to obtain a prescription drug, a provider does not have an obligation to conve				
49	that fact to the payor who provides the health benefit plan.				
50	(e) A pharmacist filling a prescription for a specific biological product may substitute a	an			
51	interchangeable biological product only if (i) the prescriber has not indicated that the pharmaci				

General Assembly Of North Carolina Session 2021 1 may not substitute an interchangeable biosimilar biological product for the prescribed biological 2 product and (ii) the Food and Drug Administration has determined the biological product to be 3 substituted is interchangeable with the prescribed biological product. 4 "§ 58-56B-25. Construction. 5 Nothing in this Article shall be construed to interfere with a patient's choice of prescription drug cost coverage or to interfere with patient choice and the ability of a health care professional 6 7 to convey the full range of prescription drug cost options to a patient. Health benefit plans, 8 pharmacy benefit managers, or any entities acting on behalf of a health benefit plan shall not 9 restrict a health care professional from communicating prescription cost options to a patient." **SECTION 2.** This act becomes effective January 1, 2023. 10