1	MEDICATION AMENDMENTS
2	2024 GENERAL SESSION
3	STATE OF UTAH
4	Chief Sponsor: Evan J. Vickers
5 6	House Sponsor:
7	LONG TITLE
8	General Description:
9	This bill amends provisions related to pharmaceutical drugs.
10	Highlighted Provisions:
11	This bill:
12	defines terms;
13	 merges provisions related to federally qualified health centers and 340B entities;
14	 enacts provisions related to how insurers and pharmacy benefit managers interact
15	with 340B entities; and
16	► limits how a pharmaceutical company can interact with a 340B entity, directly or
17	indirectly.
18	Money Appropriated in this Bill:
19	None
20	Other Special Clauses:
21	None
22	Utah Code Sections Affected:
23	AMENDS:
24	31A-46-310, as enacted by Laws of Utah 2021, Chapter 317
25	31A-48-102 , as last amended by Laws of Utah 2022, Chapter 198
26	ENACTS:
27	31A-48-104 , Utah Code Annotated 1953



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28	REPEALS:
29	31A-46-309, as enacted by Laws of Utah 2020, Chapter 275
30	31A-48-101, as enacted by Laws of Utah 2020, Chapter 198
31 32	Be it enacted by the Legislature of the state of Utah:
33	Section 1. Section 31A-46-310 is amended to read:
34	31A-46-310. Prohibited actions.
35	[(1) As used in this section, "federally qualified health center":]
36	[(a) means the same as that term is defined in 42 U.S.C. Sec. 1395x(aa)(4); and]
30 37	[(b) includes the pharmacy or pharmacies that are operated by or contract with a
38	federally qualified health center described in Subsection (1)(a) to dispense drugs purchased
39	through the federally qualified health center.]
40	(1) As used in this section, "insurance entity" means:
41	(a) an insurer or an agent of an insurer; or
42	(b) a pharmacy service entity or an agent of a pharmacy service entity.
43	(2) This section applies to a contract entered into or renewed on or after January 1,
44	[2022, between an insurer and a pharmacy described in Subsection (1)(b)] 2025, between a
45	340B entity and an insurance entity.
46	(3) An [insurer] insurance entity may not vary the amount that the [insurer] insurance
47	entity reimburses to a [federally qualified health center] 340B entity for a drug on the basis of
48	whether:
49	(a) the drug is a 340B drug; or
50	(b) the pharmacy dispensing the drug is a 340B entity.
51	(4) Subsection (3) does not apply to a drug reimbursed, directly or indirectly, by the
52	Medicaid program.
53	(5) An [insurer or an insurer's pharmacy service] insurance entity may not:
54	(a) on the basis that a [federally qualified health center] 340B entity participates,
55	directly or through a contractual arrangement, in the 340B drug discount program:
56	(i) refuse to contract with the 340B entity;
57	[(i)] (ii) assess a fee, charge-back, or other adjustment on a [federally qualified health
58	center] 340B entity;

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59	[(iii)] (iii) restrict access to the [insurer's] insurance entity's pharmacy network;
60	[(iii)] (iv) require the [federally qualified health center] 340B entity to enter into a
61	contract with a specific pharmacy to participate in the [insurer's] insurance entity's pharmacy
62	network;
63	[(iv)] (v) create a restriction or an additional charge on a patient who chooses to
64	receive drugs from a [federally qualified health center] 340B entity; [or]
65	(vi) modify a copayment or other cost-sharing requirement of a patient of the 340B
66	entity; or
67	[(v)] (vii) create any additional requirements or restrictions on the [federally qualified
68	health center] 340B entity; [or]
69	(b) base drug formulary or drug coverage decisions on whether a drug is a 340B drug
70	or whether a dispensing pharmacy is a 340B entity;
71	(c) transfer the benefit of 340B drug discount program savings from a 340B entity to an
72	insurance entity;
73	(d) unilaterally modify the definition of pharmacy in a way that is inconsistent with
74	Utah law through a contract, provider manual, or other means;
75	(e) require the 340B entity to reverse, resubmit, or clarify a claim for a 340B drug after
76	an initial adjudication;
77	(f) charge or hold a 340B entity responsible for a fee related to a claim:
78	(i) that is not apparent at the time of claim processing;
79	(ii) that is not reported on the remittance advice of an adjudicated claim; or
80	(iii) after the initial claim is adjudicated at the point of sale; or
81	[(b)] (g) require a claim for a drug to include a modifier to indicate that the drug is a
82	340B drug unless the claim is for payment, directly or indirectly, by the Medicaid program.
83	Section 2. Section 31A-48-102 is amended to read:
84	31A-48-102. Definitions.
85	As used in this chapter:
86	(1) "340B drug" means the same as that term is defined in Section 31A-46-102.
87	(2) "340B entity" means the same as that term is defined in Section 31A-46-102.
88	$[\underbrace{(1)}]$ (a) "Drug" means a substance that is:
89	(i) (A) intended for use in the diagnosis, cure, mitigation, treatment, or prevention of

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90	disease in humans; and
91	(B) recognized in or in a supplement to the official United States Pharmacopoeia, the
92	Homeopathic Pharmacopoeia of the United States, or the official National Formulary;
93	(ii) required by an applicable federal or state law or rule to be dispensed by prescription
94	only;
95	(iii) restricted to administration by practitioners only;
96	(iv) a substance other than food intended to affect the structure or a function of the
97	human body; or
98	(v) intended for use as a component of a substance described in Subsection [(1)(a)(i),
99	(ii), (iii), or (iv)] (3)(a)(i), (ii), (iii), or (iv).
100	(b) "Drug" does not include a dietary supplement.
101	[(2)] (4) "Insurer" means the same as that term is defined in Section 31A-22-634.
102	[(3)] (5) "Manufacturer" means a person that is engaged in the manufacturing of a drug
103	that is available for purchase by residents of the state.
104	(6) "Pharmaceutical manufacturer" means the same as that term is defined in Section
105	<u>31A-46-102.</u>
106	(7) "Pharmacy" means the same as that term is defined in Section 58-17b-102.
107	[4] (8) "Rebate" means the same as that term is defined in Section 31A-46-102.
108	[(5)] (9) "Wholesale acquisition cost" means the same as that term is defined in 42
109	U.S.C. Sec. 1395w-3a.
110	Section 3. Section 31A-48-104 is enacted to read:
111	31A-48-104. Prohibited conduct.
112	A pharmaceutical manufacturer, or any person involved in the distribution of a
113	pharmaceutical manufacturer's products, may not directly or indirectly:
114	(1) prohibit a pharmacy from contracting with a 340B entity, including by denying the
115	pharmacy access to a drug that is manufactured by the pharmaceutical manufacturer;
116	(2) prohibit a 340B entity from contracting with a pharmacy, including by denying the
117	340B entity access to a drug that is manufactured by the pharmaceutical manufacturer;
118	(3) deny or restrict a 340B entity from:
119	(a) acquiring or dispensing a 340B drug; or
120	(b) receiving 340B drug discount program pricing for a 340B drug, including by

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121	imposing a time limitation on a 340B entity to replenish or submit a claim for a 340B drug;
122	(4) require a 340B entity to purchase a 340B drug from a certain supplier if the
123	pharmaceutical manufacturer, or person involved in the distribution of the pharmaceutical
124	manufacturer's products, would otherwise permit the 340B entity to purchase a drug that is not
125	a 340B drug from the supplier; or
126	(5) otherwise interfere with:
127	(a) a contract between a pharmacy and a 340B entity; or
128	(b) the ability of a pharmacy and a 340B entity to enter into a contract.
129	Section 4. Repealer.
130	This bill repeals:
131	Section 31A-46-309, Reimbursement Prohibitions.
132	Section 31A-48-101, Title.
133	Section 5. Effective date.
134	This bill takes effect on May 1, 2024.