

1 **PHARMACEUTICAL REPORTING AMENDMENTS**

2 2020 SIXTH SPECIAL SESSION

3 STATE OF UTAH

4 **Chief Sponsor: Paul Ray**

5 Senate Sponsor: Evan J. Vickers

7 **LONG TITLE**

8 **General Description:**

9 This bill amends the Prescription Drug Price Transparency Act.

10 **Highlighted Provisions:**

11 This bill:

12 ▶ amends certain reporting requirements in the Prescription Drug Price Transparency
13 Act.

14 **Money Appropriated in this Bill:**

15 None

16 **Other Special Clauses:**

17 This bill provides a special effective date.

18 **Utah Code Sections Affected:**

19 AMENDS:

20 **31A-48-103**, as enacted by Laws of Utah 2020, Chapter 198

22 *Be it enacted by the Legislature of the state of Utah:*

23 Section 1. Section **31A-48-103** is amended to read:

24 **31A-48-103. Manufacturer reports -- Insurer report -- Publication by**
25 **department.**

26 (1) (a) A manufacturer of a drug shall, beginning January 1, 2022, report to the
27 department the information described in Subsection (1)(b) no more than 30 days after the day on



28 which an increase to the wholesale acquisition cost of the drug results in an increase to the
29 wholesale acquisition cost of the drug of:

- 30 (i) greater than 16% over the preceding two calendar years; or
- 31 (ii) greater than 10% over the preceding calendar year.
- 32 (b) The manufacturer shall report:
 - 33 (i) (A) the name of the drug;
 - 34 (B) the dosage form of the drug; and
 - 35 (C) the strength of the drug;
 - 36 (ii) whether the drug is a brand name drug or a generic drug;
 - 37 (iii) the effective date of the increase in the wholesale acquisition cost of the drug;
 - 38 (iv) a written description, suitable for public release, of the factors that led to the increase
39 in the wholesale acquisition cost of the drug and the significance of each factor;
 - 40 (v) the manufacturer's aggregate company-wide research and development costs for the
41 most recent year for which final audit data is available;
 - 42 (vi) the name of each of the manufacturer's drugs approved by the United States Food
43 and Drug Administration during the preceding three calendar years; and
 - 44 (vii) the names of drugs manufactured by the manufacturer that lost patent exclusivity in
45 the United States during the preceding three calendar years.
- 46 (c) Subsection (1)(a) applies only to a drug with a wholesale acquisition cost of at least
47 \$100 for a 30-day supply before the effective date of the increase in the wholesale acquisition
48 cost of the drug.
- 49 (d) A manufacturer's obligations under this Subsection (1) are fully satisfied by
50 submission of information and data that a manufacturer includes in the manufacturer's annual
51 consolidated report on Securities and Exchange Commission Form 10-K or any other public
52 disclosure.
- 53 (e) The department shall consult with representatives of manufacturers to establish a
54 single, standardized format for reporting information under this section that minimizes the
55 administrative burden of reporting for manufacturers and the state.
- 56 (f) Information provided to the department under Subsection (1)(b) may not be released
57 in a manner that:
 - 58 (i) would allow for the identification of an individual drug, therapeutic class of drugs, or

59 manufacturer; or

60 (ii) is likely to compromise the financial, competitive, or proprietary nature of the
61 information.

62 (2) [~~Before August 1 of each year,~~ On or before August 1, 2021, and on or before
63 August 1 of each year thereafter, an insurer shall report to the department in aggregate the
64 following information for the preceding [~~plan~~] calendar year for health benefit plans offered by the
65 insurer:

66 (a) for the 25 drugs for which spending by the insurer was the greatest, after adjusting for
67 rebates:

68 (i) the name of the drug;

69 (ii) the dosage form of the drug; and

70 (iii) the strength of the drug;

71 (b) the percentage increase over the previous year in net spending for all drugs, after
72 adjusting for rebates; and

73 (c) the percentage of the increase in premiums over the previous year attributable to all
74 drugs; and

75 (d) the percentage of the increase in premiums over the previous year attributable to
76 specialty drugs.

77 (3) The department shall publish on the department's website:

78 (a) no later than 60 days after receiving the information, information reported to the
79 department under Subsection (1); and

80 (b) no later than [~~November~~] December 1 of each year, information reported to the
81 department under Subsection (2).

82 (4) The department may not publish information under Subsection (3)(b) in a manner that
83 allows the identity of an insurer to be determined.

84 (5) The department shall make rules, as necessary, in accordance with Title 63G, Chapter
85 3, Utah Administrative Rulemaking Act, to promote comparability of information reported to the
86 department under this chapter.

87 Section 2. **Effective date.**

88 If approved by two-thirds of all the members elected to each house, this bill takes effect
89 upon approval by the governor, or the day following the constitutional time limit of Utah

90 Constitution, Article VII, Section 8, without the governor's signature, or in the case of a veto, the
91 date of veto override.