## SENATE BILL 5697

| Sta | ate of | Wa  | shington | 65        | 5th | Legisla | ature   | 2   | 2017 | Regu | lar | Session |
|-----|--------|-----|----------|-----------|-----|---------|---------|-----|------|------|-----|---------|
| By  | Senat  | ors | Rivers,  | Cleveland | ł,  | Conway, | Keiser, | and | Bai  | ley  |     |         |

AN ACT Relating to developing a standardized prescription drug benefit package for individual and small group market offerings; amending RCW 48.43.700 and 48.43.705; adding a new section to chapter 4 8.43 RCW; and creating a new section.

5 BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF WASHINGTON:

6 NEW SECTION. Sec. 1. The patient out-of-pocket cost task force 7 held several extended discussions on the costs patients with chronic medical conditions face associated with prescription drugs, and the 8 task force explored several benefit design strategies that might 9 10 reduce the impact of the out-of-pocket costs. Several states have 11 enacted laws requiring various levels of standardization in health benefit design, primarily focused on reducing the out-of-pocket 12 obligations for prescription drugs, and providing information that 13 14 allows consumers to compare plans and select a plan that best fits their needs. It is the intent of the legislature to establish a 15 16 process for the development and maintenance of a standardized 17 prescription drug benefit that shall be available in the individual 18 and small group markets.

19 <u>NEW SECTION.</u> Sec. 2. A new section is added to chapter 48.43 20 RCW to read as follows: 1 (1) The commissioner, in collaboration with the health benefit exchange, shall convene a committee to develop a recommendation on a 2 standardized prescription benefit design. Applications to participate 3 in the committee shall be submitted to the commissioner, and the 4 commissioner shall ensure that participants on the committee include 5 representatives from the following groups: Insurance carriers, 6 7 providers, patient groups, labor, small employers, large employers, and drug manufacturers. The commissioner shall retain a neutral 8 consultant or facilitator to assist with meetings of the committee. 9

(2) The committee shall be convened no later than October 1, 10 11 2017, and shall examine the options for designing a standardized prescription drug benefit package for use in the small group and 12 individual markets. Standardized benefit design components must 13 consider limiting or eliminating coinsurance as a cost-sharing 14 method, fixing the copayment amounts for medications, limiting 15 16 deductibles for medications, and defining any necessary parameters 17 for the tiers of coverage. The committee shall submit recommendations to the commissioner and the exchange for a standardized design by 18 October 1, 2018. The commissioner shall adopt the recommendations in 19 rule, ensuring that each carrier offering coverage in the individual 20 21 and small group markets offer at least one health plan that includes the standardized benefit design, beginning with coverage offered for 22 23 January 1, 2020.

(3) The committee shall be retained for an annual review of the standardized benefit design and any recommendations for modifications. The commissioner shall update the rule as needed to reflect recommendations from the committee.

28 **Sec. 3.** RCW 48.43.700 and 2014 c 31 s 1 are each amended to read 29 as follows:

(1) For plan or policy years beginning January 1, 2014, a carrier offering a health benefit plan that meets the definition of bronze level in section 1302 of P.L. 111-148 of 2010, as amended, in the individual market outside of the exchange must also offer plans that meet the definition of silver and gold level plans in section 1302 of P.L. 111-148 of 2010, as amended, in the individual market outside of the exchange.

37 (2) For plan or policy years beginning January 1, 2014, a carrier
38 offering a health benefit plan that meets the definition of bronze
39 level in section 1302 of P.L. 111-148 of 2010, as amended, in the

p. 2

1 small group market outside of the exchange must also offer plans that 2 meet the definition of silver and gold level plans in section 1302 of 3 P.L. 111-148 of 2010, as amended, in the small group market outside 4 of the exchange.

5 (3) A health benefit plan meeting the definition of a 6 catastrophic plan in RCW 48.43.005(8)(c)(i) may only be sold through 7 the exchange.

8 (4) By December 1, 2016, the exchange board, in consultation with 9 the commissioner, must complete a review of the impact of this 10 section on the health and viability of the markets inside and outside 11 the exchange and submit the recommendations to the legislature on 12 whether to maintain the market rules or let them expire.

13 (5) The commissioner shall evaluate plans offered at each 14 actuarial value defined in section 1302 of P.L. 111-148 of 2010, as 15 amended, and determine whether variation in prescription drug benefit 16 cost-sharing, both inside and outside the exchange in both the 17 individual and small group markets results in adverse selection. If 18 so, the commissioner may adopt rules to assure substantial 19 equivalence of prescription drug cost-sharing.

20 (6) For plan or policy years beginning January 1, 2020, a carrier 21 offering a health benefit plan in the individual or small group 22 markets must include at least one health plan, in each market, that 23 includes the standardized prescription drug benefit design developed 24 under section 2 of this act.

25 **Sec. 4.** RCW 48.43.705 and 2014 c 31 s 2 are each amended to read 26 as follows:

27 (1) All nongrandfathered individual and small group health plans, 28 other than catastrophic health plans, offered outside of the exchange 29 must conform with the actuarial value tiers specified in section 1302 30 of P.L. 111-148 of 2010, as amended, as bronze, silver, gold, or 31 platinum.

32 (2) For plan or policy years beginning January 1, 2020, a carrier 33 offering a health benefit plan in the individual or small group 34 market must include at least one health plan, in each market, that 35 includes the standardized prescription drug benefit design developed 36 under section 2 of this act.

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