
SENATE BILL 5203

State of Washington

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By Senators Van De Wege, Carlyle, Frockt, Hasegawa, Keiser, Lias, Nguyen, Randall, Robinson, Salomon, Stanford, and Wilson, C.

Read first time 01/13/21. Referred to Committee on Health & Long Term Care.

1 AN ACT Relating to the production, distribution, and purchase of
2 generic prescription drugs; amending RCW 70.14.060; and adding a new
3 section to chapter 70.14 RCW.

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

5 NEW SECTION. **Sec. 1.** A new section is added to chapter 70.14
6 RCW to read as follows:

7 (1)(a) The authority may enter into partnerships with another
8 state, a group of states, a state agency, a nonprofit organization,
9 or any other entity to produce, distribute, or purchase generic
10 prescription drugs.

11 (b) The generic prescription drugs must be produced or
12 distributed by a drug company or generic drug manufacturer that is
13 registered with the United States food and drug administration.

14 (2) The authority shall only enter into partnerships, in
15 consultation with other state agencies as necessary, to produce,
16 distribute, or purchase a generic prescription drug at a price that
17 results in savings to public and private purchasers and consumers.

18 (3) For generic prescription drugs that the authority has entered
19 into a partnership under this section:

20 (a) State purchased health care programs must purchase the
21 generic prescription drugs through the partnership, unless the state

1 purchased health care program can obtain the generic prescription
2 drug at a cost savings through another purchasing mechanism; and

3 (b) Local governments, private entities, health carriers, and
4 others may choose to voluntarily purchase the generic prescription
5 drugs from the authority as available quantities allow.

6 (4) All information and documents obtained or created under this
7 section is exempt from disclosure under chapter 42.56 RCW.

8 (5) For purposes of this section, the following definitions
9 apply:

10 (a) "Authority" means the health care authority.

11 (b) "Generic drug" means a drug that is approved pursuant to
12 section 355(j) of the federal food, drug, and cosmetic act (21 U.S.C.
13 Sec. 301 et seq.), or a biosimilar, as defined under the federal
14 public health service act (42 U.S.C. Sec. 262).

15 (c) "State purchased health care" means medical and health care,
16 pharmaceuticals, and medical equipment purchased with state and
17 federal funds by the department of social and health services,
18 department of health, state health care authority, department of
19 labor and industries, department of corrections, and department of
20 veterans affairs. State purchased health care does not include
21 prescription drugs purchased for medical assistance program clients
22 under chapter 74.09 RCW.

23 **Sec. 2.** RCW 70.14.060 and 2020 c 346 s 4 are each amended to
24 read as follows:

25 (1)(a) The (~~administrator~~[~~director~~]) director of the state
26 health care authority shall, directly or by contract, adopt policies
27 necessary for establishment of a prescription drug purchasing
28 consortium. The consortium's purchasing activities shall be based
29 upon the evidence-based prescription drug program established under
30 RCW 70.14.050. (~~State~~) Except as provided in section 1 of this act
31 or exempted under (b) of this subsection, state purchased health care
32 programs as defined in RCW 41.05.011 shall purchase prescription
33 drugs through the consortium for those prescription drugs that are
34 purchased directly by the state and those that are purchased through
35 reimbursement of pharmacies(~~, unless exempted under (b) of this~~
36 subsection)). The (~~administrator~~[~~director~~]) director shall not
37 require any supplemental rebate offered to the health care authority
38 by a pharmaceutical manufacturer for prescription drugs purchased for
39 medical assistance program clients under chapter 74.09 RCW be

1 extended to any other state purchased health care program, or to any
2 other individuals or entities participating in the consortium. The
3 (~~administrator~~ [director]) director shall explore joint purchasing
4 opportunities with other states.

5 (b) State purchased health care programs are exempt from the
6 requirements of this section if they can demonstrate to the
7 (~~administrator~~ [director]) director of the state health care
8 authority that, as a result of the availability of federal programs
9 or other purchasing arrangements, their other purchasing mechanisms
10 will result in greater discounts and aggregate cost savings than
11 would be realized through participation in the consortium.

12 (2) Participation in the purchasing consortium shall be offered
13 as an option beginning January 1, 2006. Participation in the
14 consortium is purely voluntary for units of local government, private
15 entities, labor organizations, health carriers as provided in RCW
16 48.43.005, state purchased health care services from or through
17 health carriers as provided in RCW 48.43.005, and for individuals who
18 lack or are underinsured for prescription drug coverage. The
19 (~~administrator~~ [director]) director may set reasonable fees,
20 including enrollment fees, to cover administrative costs attributable
21 to participation in the prescription drug consortium.

22 (3) The state health care authority is authorized to adopt rules
23 implementing chapter 129, Laws of 2005.

24 NEW SECTION. **Sec. 3.** If any provision of this act or its
25 application to any person or circumstance is held invalid, the
26 remainder of the act or the application of the provision to other
27 persons or circumstances is not affected.

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