

Senate File 2204 - Introduced

SENATE FILE 2204
BY COMMITTEE ON COMMERCE

(SUCCESSOR TO SSB 3087)

A BILL FOR

1 An Act relating to the regulation of pharmacy benefits
2 managers.
3 BE IT ENACTED BY THE GENERAL ASSEMBLY OF THE STATE OF IOWA:

1 Section 1. Section 510B.1, Code 2014, is amended by adding
2 the following new subsection:

3 NEW SUBSECTION. 5A. *“Maximum reimbursement amount”* means
4 the maximum reimbursement amount for a therapeutically and
5 pharmaceutically equivalent multiple-source prescription drug
6 that is listed in the most recent edition of the publication
7 entitled approved drug products with therapeutic equivalence
8 evaluations, published by the United States food and drug
9 administration, otherwise known as the orange book.

10 Sec. 2. Section 510B.2, Code 2014, is amended by adding the
11 following new unnumbered paragraph:

12 NEW UNNUMBERED PARAGRAPH. As a condition of complying with
13 this section regarding certification, a pharmacy benefits
14 manager shall enter into and maintain an agreement with
15 the commissioner setting out the terms of disclosure of
16 information, as deemed necessary by the commissioner, relating
17 to rebates received by the pharmacy benefits manager.

18 Sec. 3. NEW SECTION. 510B.8 **Pricing methodology for maximum**
19 **reimbursement amount.**

20 1. The commissioner may require a pharmacy benefits manager
21 to submit information to the commissioner related to the
22 pharmacy benefits manager’s pricing methodology for maximum
23 reimbursement amount.

24 2. For purposes of the disclosure of pricing methodology,
25 maximum reimbursement amounts shall be implemented as follows:

26 a. Established for multiple source prescription drugs
27 prescribed after the expiration of any generic exclusivity
28 period.

29 b. Established for any prescription drug with at least two
30 or more A-rated therapeutically equivalent, multiple source
31 prescription drugs with a significant cost difference.

32 c. Determined using comparable prescription drug prices
33 obtained from multiple nationally recognized comprehensive data
34 sources including wholesalers, prescription drug file vendors,
35 and pharmaceutical manufacturers for prescription drugs that

1 are nationally available and available for purchase locally by
2 multiple pharmacies in the state.

3 3. For those prescription drugs to which maximum
4 reimbursement amount pricing applies, a pharmacy benefits
5 manager shall include in a contract with a pharmacy information
6 regarding which of the national compendia is used to
7 obtain pricing data used in the calculation of the maximum
8 reimbursement amount pricing and shall provide a process to
9 allow a pharmacy to comment on, contest, or appeal the maximum
10 reimbursement amount rates or maximum reimbursement amount
11 list. The right to comment on, contest, or appeal the maximum
12 reimbursement amount rates or maximum reimbursement amount list
13 shall be limited in duration and allow for retroactive payment
14 in the event that it is determined that maximum reimbursement
15 amount pricing has been applied incorrectly.

16 Sec. 4. COMPLIANCE PERIOD FOR DISCLOSURE AGREEMENTS. In
17 order to comply with the certification and disclosure
18 requirements of section 510B.2, as amended in this Act, a
19 pharmacy benefits manager shall have one hundred eighty days
20 after the effective date of this Act within which to enter
21 into an agreement with the commissioner regarding disclosure
22 of rebate information.

23

EXPLANATION

24 The inclusion of this explanation does not constitute agreement with
25 the explanation's substance by the members of the general assembly.

26 This bill relates to the regulation of pharmacy benefits
27 managers. The bill authorizes the commissioner of insurance
28 to require a pharmacy benefits manager to submit information
29 to the commissioner related to the pharmacy benefits manager's
30 pricing methodology for maximum reimbursement amounts for
31 prescription drugs. "Maximum reimbursement amount" is defined
32 as the maximum reimbursement amount for a therapeutically
33 and pharmaceutically equivalent multiple-source prescription
34 drug that is listed in the United States food and drug
35 administration's publication entitled approved drug products

1 with therapeutic equivalence evaluations, otherwise known as
2 the orange book.

3 For those prescription drugs to which maximum reimbursement
4 amount pricing applies, a pharmacy benefits manager must
5 include information in a contract with a pharmacy showing how
6 maximum reimbursement amount pricing is calculated and allowing
7 the pharmacy the opportunity to comment on, contest, or appeal
8 the maximum reimbursement amount rates and list. The contract
9 must also allow for retroactive payment if it is determined
10 that maximum reimbursement amount pricing has been applied
11 incorrectly.

12 The bill also allows the commissioner to implement the
13 information submission requirements in relation to generic
14 exclusivity, therapeutically equivalent drugs, and multiple
15 drug manufacturers, wholesalers, and vendors.

16 The bill provides that pharmacy benefits managers must,
17 within six months of the effective date of the bill, enter into
18 an agreement with the commissioner relating to disclosure of
19 rebates received by the pharmacy benefits managers.