

# SENATE, No. 2769

## STATE OF NEW JERSEY 217th LEGISLATURE

INTRODUCED NOVEMBER 10, 2016

**Sponsored by:**

**Senator RICHARD J. CODEY**

**District 27 (Essex and Morris)**

**Senator JOSEPH F. VITALE**

**District 19 (Middlesex)**

**SYNOPSIS**

Prohibits use of manufacturer coupons for certain prescription drug and prescription biological products.

**CURRENT VERSION OF TEXT**

As introduced.



**(Sponsorship Updated As Of: 11/15/2016)**

1 AN ACT concerning prescription medications and supplementing  
2 Title 24 of the Revised Statutes.

3

4 **BE IT ENACTED** by the Senate and General Assembly of the State  
5 of New Jersey:

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7 1. a. A manufacturer of a prescription drug product or  
8 prescription biological product shall be prohibited from offering  
9 any discount, rebate, product voucher, or other reduction in an  
10 individual's out-of-pocket expenses, including a co-payment or  
11 deductible, for any prescription drug or biological product if a  
12 lower cost brand name or nonbrand name product is available that is  
13 designated by the United States Food and Drug Administration as  
14 therapeutically equivalent to, or interchangeable with, the  
15 manufacturer's product.

16 b. Subsection a. of this section shall not apply to a discount,  
17 rebate, or other payment by a manufacturer of a prescription drug  
18 product or prescription biological product to a patient or another  
19 person on the patient's behalf, other than the prescriber of the drug  
20 or biological product, for health care items or services related to the  
21 patient's use of a prescription drug or biological product where the  
22 item or service is required under a U.S. Food and Drug  
23 Administration Risk Evaluation and Mitigation Strategy or are for  
24 the purpose of monitoring or facilitating the use of the prescription  
25 drug or biological product in a manner consistent with the product's  
26 approved labeling.

27 c. Nothing in this section shall be deemed to:

28 (1) Restrict a manufacturer of a prescription drug product or  
29 prescription biological product with regard to how it distributes a  
30 drug or biological product;

31 (2) Restrict a carrier, as that term is defined in section 2 of  
32 P.L.1997, c.192 (C.26:2S-2), with regard to how its health benefits  
33 plan design treats such discount, rebate, product voucher, or other  
34 reduction in out-of-pocket expenses; or

35 (3) Affect a pharmacist's duties with regard to substitutions  
36 involving interchangeable drug products pursuant to P.L.1977,  
37 c.240 (C.24:6E-1 et seq.) or interchangeable biological products  
38 pursuant to P.L.2015, c.130 (C.24:6K-1 et al.).

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40 2. This act shall take effect the first day of the fourth month  
41 next following the date of enactment.

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#### STATEMENT

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46 This bill prohibits manufacturers of prescription drug products  
47 and prescription biological products from offering discounts,  
48 rebates, product vouchers, or other reductions in an individual's

1 out-of-pocket expenses, including co-payments and deductibles, for  
2 prescription drug or biological products if a lower-cost alternative  
3 product is available that has been designated by the United States  
4 Food and Drug Administration (FDA) as therapeutically equivalent  
5 to, or interchangeable with, the manufacturer's product. Alternative  
6 products may include both brand name and nonbrand name  
7 products.

8 This restriction will not apply to discounts, rebates, or other  
9 payments made by a manufacturer to a patient for health care items  
10 or services related to the patient's use of a prescription drug or  
11 biological product when the items or services are required under an  
12 FDA Risk Evaluation and Mitigation Strategy or are for the purpose  
13 of monitoring or facilitating an approved use of the product.

14 Nothing in the bill will restrict how a manufacturer may  
15 distribute a drug or product; restrict a health benefits carrier in how  
16 its plan design will treat discounts, rebates, product vouchers, or  
17 other reductions in out-of-pocket expenses; or affect a pharmacist's  
18 duties with regard to substitutions involving interchangeable drug  
19 and biological products under current law.

20 A 2013 article published in the New England Journal of  
21 Medicine discusses how pharmaceutical manufacturers frequently  
22 use coupons and discount offers to encourage consumers to choose  
23 that manufacturer's products. In the short term, the coupons and  
24 discounts reduce the consumer's out-of-pocket costs for the  
25 product, providing the consumer with an incentive to choose the  
26 product even when lower cost alternatives, such as generic versions  
27 of the product, are available. However, the article suggests this  
28 practice raises the cost of providing coverage for health and  
29 prescription benefits providers, which in turn results in higher costs  
30 for all consumers. Additionally, the practice generally results in  
31 higher long-term costs for individual consumers, who frequently  
32 continue to use a manufacturer's product even after the coupon or  
33 discount is no longer available, which is generally not effective for  
34 more than one year.

35 It is the sponsor's belief that restricting the use of coupon and  
36 discount offers when lower cost alternatives are available will help  
37 reduce the overall cost of health care for the citizens of New Jersey.