

WEST VIRGINIA LEGISLATURE

SECOND REGULAR SESSION, 2002

ENROLLED

House Bill No. 4666

(By Delegates Warner, Leach, Boggs, Campbell, Proudfoot and Fletcher)

Passed March 7, 2002

In Effect from Passage

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GATTLE MEST VIRGINIA SECRETARY OF STATE

ENROLLED

H. B. 4666

(By Delegates Warner, Leach, Boggs, Campbell, Proudfoot and Fletcher)

[Passed March 7, 2002; in effect from passage.]

AN ACT to amend and reenact section fifteen, article five, chapter nine of the code of West Virginia, one thousand nine hundred thirty-one, as amended, relating to establishing a process for the secretary of health and human services to enter into negotiations with pharmaceutical companies for rebates that cannot be accessed through Freedom of Information Act requests or through open meetings.

Be it enacted by the Legislature of West Virginia:

That section fifteen, article five, chapter nine of the code of West Virginia, one thousand nine hundred thirty-one, as amended, be amended and reenacted to read as follows:

ARTICLE 5. MISCELLANEOUS PROVISIONS.

§9-5-15. Medicaid program; preferred drug list and drug utilization review.

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1 The Legislature finds that it is a public necessity that trade 2 secrets, rebate amounts, percentage of rebate, manufacturer's 3 pricing and supplemental rebates that are contained in records, 4 as well as any meetings at which this information is negotiated 5 or discussed need confidentiality to insure the most significant 6 rebates available for the state. Information pertaining to similar 7 agreements with the federal government and negotiated by 8 pharmaceutical manufacturers is confidential pursuant to 42 9 U.S.C 1396r-8. A rebate as a percentage of average manufac-10 ture price is confidential under federal law and the federal 11 rebate could be made known if not protected by state law. 12 Because of the protection afforded by federal law, if this 13 information is not protected by state law, manufacturers will 14 not be willing to offer a rebate in West Virginia. Further, the 15 Legislature finds that the number and value of supplemental 16 rebates obtained by the department will increase, to the benefit 17 of Medicaid recipients, if information related to the supplemen-18 tal rebates is protected in the records of the department and in 19 meetings in which this information is disclosed because 20 manufactures will be assured they will not to be placed at a 21 competitive disadvantage by exposure of this information.

The secretary of the department of health and human resources has the authority to develop a preferred drug list, in accordance with federal law, which shall consist of federally approved drugs. The department, through administration of the medicaid program, may reimburse, where applicable and in accordance with federal law, entities providing and dispensing prescription drugs from the preferred drug list.

The secretary of the department is hereby authorized to negotiate and enter into agreements with pharmaceutical manufacturers for supplemental rebates for medicaid reimbursable drugs.

- The provisions of article three, chapter five-a of this code shall not apply to any contract or contracts entered into under this section.
- Trade secrets, rebate amounts, percentage of rebate, manufacturer's pricing and supplemental rebates which are contained in the department's records and those of its agents with respect to supplemental rebate negotiations and which are prepared pursuant to a supplemental rebate agreement are confidential and exempt from all of article one, chapter twentynine-b of this code.
- Those portions of any meetings of the committee at which trade secrets, rebate amounts, percentage of rebate, manufacturer's pricing and supplemental rebates are disclosed for discussion or negotiation of a supplemental rebate agreement are exempt from all of article nine-a, chapter six of this code.
- The secretary of the department will monitor and evaluate the effects of this provision on medicaid recipients, the medicaid program, physicians and pharmacies.
- The commissioner shall implement a drug utilization review program to assure that prescribing and dispensing of drug products result in the most rational cost-effective medication therapy for medicaid patients.
- Any moneys received in supplemental rebates will be deposited in the medical services fund established in section two, article four, chapter nine of this code.

That Joint Committee on Enrolled Bills hereby certifies that the
foregoing bill is correctly enrolled.
Chairman Senate Committee
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Chairman House Committee
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