## THE GENERAL ASSEMBLY OF PENNSYLVANIA

## SENATE BILL

No. 7

Session of 2003

INTRODUCED BY ORIE, MOWERY, HELFRICK, LEMMOND, GREENLEAF, O'PAKE, TARTAGLIONE, MUSTO, WOZNIAK, BOSCOLA, RAFFERTY AND TOMLINSON, MAY 5, 2003

REFERRED TO AGING AND YOUTH, MAY 5, 2003

## AN ACT

- Amending the act of August 26, 1971 (P.L.351, No.91), entitled "An act providing for a State Lottery and administration 3 thereof; authorizing the creation of a State Lottery 4 Commission; prescribing its powers and duties; disposition of funds; violations and penalties therefor; exemption of prizes 6 from State and local taxation and making an appropriation," 7 defining "best price"; and further providing for prescription 8 drug programs generally, for reimbursement, for the Pharmaceutical Assistance Contract Needs Enhancement Tier, 9 for amount of rebate and for excessive pharmaceutical price 10 inflation discount. 11 12 The General Assembly of the Commonwealth of Pennsylvania 13 hereby enacts as follows: 14 Section 1. Section 502 of the act of August 26, 1971 15 (P.L.351, No.91), known as the State Lottery Law, is amended by 16 adding a definition to read: Section 502. Definitions. 17
- 18 The following words and phrases when used in this chapter
- 19 shall have the meanings given to them in this section unless the
- 20 context clearly indicates otherwise:
- 21 \* \* \*
- 22 "Best price." As defined in section 1927(c)(1)(C) of the

- 1 Social Security Act (49 Stat. 620, 42 U.S.C. § 1396r-
- $2 \frac{8(c)(1)(C)}{.}$
- 3 \* \* \*
- 4 Section 2. Sections 509, 515, 519, 705 and 706 of the act,
- 5 added November 21, 1996 (P.L.741, No.134), are amended to read:
- 6 Section 509. Program generally.
- 7 The program shall include the following:
- 8 (1) Participating pharmacies shall be paid within 21
- 9 days of the contracting firm receiving the appropriate
- 10 substantiation of the transaction. Pharmacies shall be
- 11 entitled to interest for payment not made within the 21-day
- 12 period at a rate approved by the board.
- 13 (2) Collection of the copayment by pharmacies shall be
- 14 mandatory.
- 15 (3) Senior citizens participating in the program are not
- required to maintain records of each transaction.
- 17 (4) A system of rebates or reimbursements to eligible
- 18 claimants for pharmaceutical expenses shall be prohibited.
- 19 (5) PACE shall include a participant copayment schedule
- 20 for each prescription. The copayment may increase or decrease
- on an annual basis by the average percent change of
- 22 ingredient costs for all prescription drugs, plus a
- differential to raise the copayment to the next highest 25ç
- increment. In addition, the department may approve a request
- for increase or decrease in the level of copayment based upon
- 26 the financial experience and projections of PACE and after
- 27 consultation with the board. The department is prohibited
- from approving adjustments to the copayment on more than an
- 29 annual basis.
- 30 (6) The program shall consist of payments to pharmacies

- on behalf of eligible claimants [for]:
- 2 <u>(i) For single-source drugs,</u> 90% of the average
- 3 wholesale costs of prescription drugs which exceed the
- 4 copayment, plus a dispensing fee of at least \$3.50 or the
- 5 dispensing fee established by the department by
- 6 regulation, whichever is greater.
- 7 (ii) For multiple-source drugs, the department shall
- 8 adopt the Federal upper reimbursement limit for each
- 9 <u>multiple-source drug for which the FDA has rated three or</u>
- 10 more products therapeutically and pharmaceutically
- 11 <u>equivalent, regardless of whether all such additional</u>
- formulations are rated as such and shall use only such
- formulations when determining any such upper limit.
- 14 (7) In no case shall the Commonwealth or any person
- enrolled in the program be charged more than the price of the
- drug at the particular pharmacy on the date of the sale.
- 17 Section 515. Reimbursement.
- For-profit third-party insurers, health maintenance
- 19 organizations and not-for-profit prescription plans shall be
- 20 responsible for any payments made to a providing pharmacy on
- 21 behalf of a claimant covered by such a third party.
- 22 Section 519. The Pharmaceutical Assistance Contract for the
- 23 Elderly Needs Enhancement Tier.
- 24 (a) Establishment.--There is hereby established within the
- 25 department a program to be known as the Pharmaceutical
- 26 Assistance Contract for the Elderly Needs Enhancement Tier
- 27 (PACENET).
- 28 (b) PACENET eligibility. -- A claimant with an annual income
- 29 of not less than \$14,000 and not more than [\$16,000] <u>\$17,000</u> in
- 30 the case of a single person and of not less than \$17,200 and not

- 1 more than [\$19,200] \$23,200 in the case of the combined income
- 2 of persons married to each other shall be eligible for enhanced
- 3 pharmaceutical assistance under this section. A person may, in
- 4 reporting income to the department, round the amount of each
- 5 source of income and the income total to the nearest whole
- 6 dollar, whereby any amount which is less than 50ç is eliminated.
- 7 (c) Deductible.--Upon enrollment in PACENET, eligible
- 8 claimants in the income ranges set forth in subsection (b) shall
- 9 be required to meet an annual deductible in unreimbursed
- 10 prescription drug expenses of [\$500] \$480 per person. To qualify
- 11 for the deductible set forth in this subsection the prescription
- 12 drug must be purchased for the use of the eligible claimant from
- 13 a provider as defined in this chapter. The department, after
- 14 consultation with the board, may approve an adjustment in the
- 15 deductible on an annual basis. The annual deductible may be met
- 16 through unreimbursed drug expenses of \$40 per person per
- 17 calendar month.
- 18 (d) Copayment. -- For eligible claimants under this section,
- 19 the copayment schedule, which may be adjusted by the department
- 20 on an annual basis after consultation with the board, shall be:
- 21 (i) eight dollars for noninnovator multiple source
- drugs as defined in section 702; or
- 23 (ii) fifteen dollars for single-source drugs and
- innovator multiple-source drugs as defined in section
- 25 702.
- 26 Section 705. Amount of rebate.
- 27 (a) Single-source drugs and innovator multiple-source
- 28 drugs. -- With respect to single-source drugs and innovator
- 29 multiple-source drugs, each manufacturer shall remit a rebate to
- 30 the Commonwealth. Except as otherwise provided in this section,

- 1 the amount of the rebate to the Commonwealth per calendar
- 2 quarter with respect to each dosage form and strength of single-
- 3 source drugs and innovator multiple-source drugs shall be as
- 4 follows:
- 5 (1) For quarters beginning after September 30, 1992, and
- 6 ending before January 1, 1997, the product of the total
- 7 number of units of each dosage form and strength reimbursed
- 8 by PACE and General Assistance in the quarter and the
- 9 difference between the average manufacturer price and 85% of
- 10 that price, after deducting customary prompt payment
- 11 discounts, for the quarter.
- 12 (2) For quarters beginning after December 31, 1996, the
- 13 product of the total number of units of each dosage form and
- strength reimbursed by [PACE, PACENET and] designated
- pharmaceutical programs other than PACE and PACENET in the
- quarter and the difference between the average manufacturer
- 17 price and 83% of that price, after deducting customary prompt
- 18 payment discounts.
- 19 (3) For quarters beginning after December 31, 2003, the
- 20 product of the total number of units of each dosage form and
- 21 <u>strength reimbursed by PACE and PACENET in the quarter and</u>
- 22 the difference between the average manufacturer price and the
- 23 best price, after deducting customary prompt payment
- 24 <u>discounts.</u>
- 25 (b) Rebate for other drugs.--
- 26 (1) The amount of the rebate to the Commonwealth for a
- 27 calendar quarter with respect to covered prescription drugs
- which are noninnovator multiple-source drugs shall be equal
- 29 to the product of:
- 30 (i) the applicable percentage of the average

- 1 manufacturer price, after deducting customary prompt
- 2 payment discounts, for each dosage form and strength of
- 3 such drugs for the quarter; and
- 4 (ii) the number of units of such form and dosage
- 5 reimbursed by PACE and General Assistance in the quarter.
- 6 (2) For the purposes of paragraph (1), the applicable
- 7 percentage for calendar quarters beginning after September
- 8 30, 1992, and ending before January 1, 1997, is 11%.
- 9 (c) Revised rebate for other drugs.--Beginning after
- 10 December 31, 1996:
- 11 (1) The amount of the rebate to the Commonwealth for a
- 12 calendar quarter with respect to covered prescription drugs
- which are noninnovator multiple-source drugs shall be the
- 14 greater of the product of:
- 15 (i) the applicable percentage of the average
- 16 manufacturer price, after deducting customary prompt
- 17 payment discounts, for each dosage form and strength of
- such drugs for the quarter; and
- 19 (ii) the number of units of such form and dosage
- 20 reimbursed by PACE, PACENET and designated pharmaceutical
- 21 programs in the quarter.
- 22 (2) For purposes of paragraph (1), the applicable
- percentage is 17%.
- 24 (d) Drugs approved after act takes effect.--In the case of a
- 25 covered outpatient drug approved for marketing after the
- 26 effective date of the act of August 14, 1991 (P.L.342, No.36),
- 27 known as the Lottery Fund Preservation Act, any reference to
- 28 January 1, 1991, shall be a reference to the first day of the
- 29 first month during which the drug was marketed.
- 30 Section 706. Excessive pharmaceutical price inflation discount.

- 1 (a) General rule. -- A discount shall be provided to the
- 2 department for all covered prescription drugs. The discount
- 3 shall be calculated as follows:
- 4 (1) For each quarter for which a rebate under section
- 5 705(a) and (b) is to be paid after December 31, 1991, and
- 6 before January 1, 1997, the average manufacturer price for
- 7 each dosage form and strength of a covered prescription drug
- 8 shall be compared to the average manufacturer price for the
- 9 same form and strength in the previous calendar year, and a
- 10 percentage increase shall be calculated.
- 11 (2) For each quarter under paragraph (1), the average
- 12 percentage increase in the Producer Price Index for
- 13 Pharmaceuticals over the same quarter in the previous
- 14 calendar year shall be calculated.
- 15 (3) If the calculation under paragraph (1) is greater
- than the calculation under paragraph (2), the discount amount
- for each quarter shall be equal to the product of:
- 18 (i) the difference between the calculations under
- paragraphs (1) and (2); and
- 20 (ii) the total number of units of each dosage form
- 21 and strength reimbursed by PACE and General Assistance
- and the average manufacturer price reported by the
- manufacturer under section 704(c)(1).
- 24 (b) Revised general rule. -- A discount shall be provided to
- 25 the department for all covered prescription drugs. The discount
- 26 shall be calculated as follows:
- 27 (1) [For] Except as provided in subsection (b.1), for
- each quarter for which a rebate under section 705(a) and (c)
- is to be paid after December 31, 1996, the average
- 30 manufacturer price for each dosage form and strength of a

- 1 covered prescription drug shall be compared to the average
- 2 manufacturer price for the same form and strength in the
- 3 previous calendar year and a percentage increase shall be
- 4 calculated.
- 5 (2) For each quarter under paragraph (1), the average
- 6 percentage increase in the Consumer Price Index-Urban over
- 7 the same quarter in the previous calendar year shall be
- 8 calculated.
- 9 (3) If the calculation under paragraph (1) is greater
- than the calculation under paragraph (2), the discount amount
- for each quarter shall be equal to the product of:
- 12 (i) the difference between the calculations under
- paragraphs (1) and (2); and
- 14 (ii) the total number of units of each dosage form
- and strength reimbursed by PACE, PACENET and designated
- 16 pharmaceutical programs and the average manufacturer
- 17 price reported by the manufacturer under section
- 18 704(c)(1).
- 19 (b.1) Discount after December 31, 2003, for PACE and
- 20 PACENET. -- A discount in lieu of the discount provided under
- 21 subsection (b) shall be provided to the department for all the
- 22 covered prescription drugs under this subsection. The discount
- 23 shall be calculated as provided under this subsection. For each
- 24 quarter for which a discount specified in this subsection for a
- 25 rebate under section 705(a) and (c) is paid after December 31,
- 26 2003, with respect to each dosage form and strength reimbursed
- 27 by PACE and PACENET:
- 28 (1) the amount shall be increased by an amount equal to
- 29 <u>the product of:</u>
- (i) the total number of units of each dosage form

1 and strength reimbursed by PACE and PACENET, for which payment was made under section 705(a) and (c) for the 2. 3 rebate period; and 4 (ii) the amount, if any, by which the average 5 manufacturer price for the dosage form and strength of the drug for the period, exceeds the average manufacturer 6 7 price for such dosage form and strength for the calendar quarter beginning July 1, 2003, without regard to whether 8 9 or not the drug has been sold or transferred to an 10 entity, including a division or subsidiary of the 11 manufacturer, after the first day of such quarter, increased by the percentage by which the Consumer Price 12 13 Index-Urban for the quarter before the quarter in which the rebate period begins exceeds such index for September 14 15 2003; or 16 (2) in the case of covered prescription drugs that have not been marketed before October 1, 2003, the amount shall be 17 18 increased by an amount equal to the product of: (i) the total number of units of each dosage form 19 20 and strength reimbursed by PACE and PACENET, for which payment was made under section 705(a) and (c) for the 21 22 rebate period; and 23 (ii) the amount, if any, by which the average manufacturer price for the dosage form and strength of 24 the drug for the period, exceeds the average manufacturer 25 26 price for such dosage form and strength for the first 27 full calendar quarter after the day on which the drug was 28 first marketed, without regard to whether or not the drug has been sold or transferred to an entity, including a 29 division or subsidiary of the manufacturer, after the 30

- first day of such quarter, increased by the percentage by
  which the Consumer Price Index-Urban for the quarter

  before the quarter in which the rebate period begins

  exceeds such index for the month prior to the first month

  of the first full calendar quarter after the day on which

  the drug was first marketed.
- 7 (c) New bimarketed drugs.--For covered prescription drugs
  8 that have not been marketed for a full calendar year, subsection
  9 (a) shall apply after the covered prescription drug has been on
  10 the market for four consecutive quarters. The drug's initial
  11 average manufacturer price shall be based on the first day of
  12 the first quarter that the drug was marketed.
- 13 Section 3. This act shall take effect in 60 days.