

AMENDMENTS TO HOUSE BILL NO. 946

Sponsor: SENATOR BROWNE

Printer's No. 3933

1 Amend Bill, page 1, line 3, by striking out "AND" and
2 inserting a comma

3 Amend Bill, page 1, line 3, by inserting after "TRANSPARENCY"
4 and for prescription drugs reimbursed under the PACE and
5 PACENET program; and making related repeals

6 Amend Bill, page 21, by inserting after line 30

CHAPTER 7

PACE AND PACENET PROGRAM PAYMENTS

9 Section 701. Definitions.

10 The following words and phrases when used in this chapter
11 shall have the meanings given to them in this section unless the
12 context clearly indicates otherwise:

13 "A-rated generic therapeutically equivalent drug." A drug
14 product that the Commissioner of Food and Drugs of the United
15 States Food and Drug Administration has approved as safe and
16 effective and has determined to be therapeutically equivalent,
17 as listed in "The Approved Drug Products with Therapeutic
18 Equivalence Evaluations" (Food and Drug Administration "Orange
19 Book"), with a specific "A" code designation only.

20 "Claimant." An eligible person who is enrolled in the
21 program.

22 "Department." The Department of Aging of the Commonwealth.

23 "Less expensive." The lowest net cost to the program. The
24 net cost shall include the amount paid by the Commonwealth to a
25 pharmacy for a drug under a current retail pharmacy
26 reimbursement formula less any discount or rebates, including
27 those paid during the previous calendar quarter and inclusive of
28 all dispensing fees.

29 "NADAC per unit." The current National Average Drug
30 Acquisition Cost per unit.

31 "Prescription drug." All drugs requiring a prescription in
32 this Commonwealth, insulin, insulin syringes and insulin
33 needles. Experimental drugs or drugs prescribed for wrinkle
34 removal or hair growth are prohibited.

35 "Program." The Pharmaceutical Assistance Contract for the

1 Elderly (PACE) and the Pharmaceutical Assistance Contract for
2 the Elderly Needs Enhancement Tier (PACENET) as established by
3 the act of August 26, 1971 (P.L.351, No.91), known as the State
4 Lottery Law.

5 "Provider." A pharmacy, dispensing physician or certified
6 registered nurse practitioner enrolled as a provider in the
7 program.

8 "Wholesale acquisition cost." The cost of a dispensed drug
9 based upon the price published in a national drug pricing system
10 in current use by the Department of Aging as the wholesale
11 acquisition cost of a prescription drug in the most common
12 package size.

13 Section 702. Program payment.

14 In addition to the requirements under section 509 of the act
15 of August 26, 1971 (P.L.351, No.91), known as the State Lottery
16 Law, the department shall administer the program in accordance
17 with the following:

18 (1) If the NADAC per unit is available, the program
19 payment shall be the lower of the following amounts:

20 (i) the NADAC per unit:

21 (A) with the addition of a professional
22 dispensing fee of \$13 per prescription; and

23 (B) the subtraction of the copayment; or

24 (ii) the pharmacy's usual and customary charge for
25 the drug dispensed with the subtraction of the copayment.

26 (2) If the NADAC per unit is unavailable, the program
27 payment shall be the lower of the following amounts:

28 (i) the wholesale acquisition cost plus 3.2%:

29 (A) with the addition of a professional
30 dispensing fee of \$13 per prescription; and

31 (B) the subtraction of the copayment; or

32 (ii) the pharmacy's usual and customary charge for
33 the drug dispensed with the subtraction of the copayment.

34 Section 703. Generic drugs.

35 (a) General rule.--Notwithstanding any other statute or
36 regulation, a brand name product shall be dispensed and not
37 substituted with an A-rated generic therapeutically equivalent
38 drug if it is less expensive to the program. If a less expensive
39 A-rated generic therapeutically equivalent drug is available for
40 dispensing to a claimant, the provider shall dispense the A-
41 rated generic therapeutically equivalent drug to the claimant.
42 The department shall reimburse providers based upon the most
43 current listing of the NADAC per unit plus a professional
44 dispensing fee of \$13 per prescription. The department shall not
45 reimburse providers for brand name products except in the
46 following circumstances:

47 (1) There is no A-rated generic therapeutically
48 equivalent drug available on the market. This paragraph does
49 not apply to the lack of availability of an A-rated generic
50 therapeutically equivalent drug in the providing pharmacy
51 unless it can be shown to the department that the provider

1 made reasonable attempts to obtain the A-rated generic
2 therapeutically equivalent drug or that there was an
3 unforeseeable demand and depletion of the supply of the A-
4 rated generic therapeutically equivalent drug. In either
5 case, the department shall reimburse the provider for the
6 NADAC per unit plus a professional dispensing fee of \$13 per
7 prescription.

8 (2) An A-rated generic therapeutically equivalent drug
9 is deemed by the department, in consultation with a
10 utilization review committee, to have too narrow a
11 therapeutic index for safe and effective dispensing in the
12 community setting. The department shall notify providing
13 pharmacies of A-rated generic therapeutically equivalent
14 drugs that are identified pursuant to this paragraph on a
15 regular basis.

16 (3) The Department of Health has determined that a drug
17 shall not be recognized as an A-rated generic therapeutically
18 equivalent drug for purpose of substitution under section
19 5(b) of the act of November 24, 1976 (P.L.1163, No.259),
20 referred to as the Generic Equivalent Drug Law.

21 (4) At the time of dispensing, the provider has a
22 prescription on which the brand name drug dispensed is billed
23 to the program by the provider at a usual and customary
24 charge which is equal to or less than the least expensive
25 usual and customary charge of any A-rated generic
26 therapeutically equivalent drug reasonably available on the
27 market to the provider.

28 (5) The brand name drug is less expensive to the
29 program.

30 (b) Generic not accepted.--If a claimant chooses not to
31 accept the A-rated generic therapeutically equivalent drug
32 required by subsection (a), the claimant shall be liable for the
33 copayment and the NADAC per unit.

34 Amend Bill, page 22, by inserting between lines 2 and 3
35 Section 1101. Repeals.

36 Repeals are as follows:

37 (1) The General Assembly declares that the repeals under
38 paragraph (2) are necessary to effectuate Chapter 7.

39 (2) Sections 509(6) and 510(a) and (b) of the act of
40 August 26, 1971 (P.L.351, No.91), known as the State Lottery
41 Law, are repealed.

42 Amend Bill, page 22, line 3, by striking out "1101" and
43 inserting
44 1102

45 Amend Bill, page 22, by inserting between lines 4 and 5

46 (1) The following provisions shall take effect

1 immediately:
2 (i) This section.
3 (ii) Chapter 7.

4 Amend Bill, page 22, line 5, by striking out "(1) THE
5 ADDITION OF " and inserting

6 (2)

7 Amend Bill, page 22, line 7, by striking out "(2)" and
8 inserting

9 (3)