

THE GENERAL ASSEMBLY OF PENNSYLVANIA

HOUSE BILL

No. 793 Session of  
2003

INTRODUCED BY J. TAYLOR, BEBKO-JONES, BELARDI, BROWNE, BUNT,  
BUTKOVITZ, CORNELL, COY, DAILEY, DALEY, DALLY, DeLUCA,  
DeWEESE, FRANKEL, GANNON, HARHAI, HENNESSEY, HORSEY, JAMES,  
KELLER, KENNEY, LEDERER, LEWIS, MAITLAND, McGEEHAN, MELIO,  
PAYNE, PETRONE, PISTELLA, SATHER, SAYLOR, SCRIMENTI, SOLOBAY,  
THOMAS, TIGUE, VANCE, WALKO AND YOUNGBLOOD, MARCH 10, 2003

REFERRED TO COMMITTEE ON INSURANCE, MARCH 10, 2003

AN ACT

1 Amending the act of May 17, 1921 (P.L.682, No.284), entitled "An  
2 act relating to insurance; amending, revising, and  
3 consolidating the law providing for the incorporation of  
4 insurance companies, and the regulation, supervision, and  
5 protection of home and foreign insurance companies, Lloyds  
6 associations, reciprocal and inter-insurance exchanges, and  
7 fire insurance rating bureaus, and the regulation and  
8 supervision of insurance carried by such companies,  
9 associations, and exchanges, including insurance carried by  
10 the State Workmen's Insurance Fund; providing penalties; and  
11 repealing existing laws," providing for certain prescription  
12 drug coverage.

13 The General Assembly of the Commonwealth of Pennsylvania  
14 hereby enacts as follows:

15 Section 1. The act of May 17, 1921 (P.L.682, No.284), known  
16 as The Insurance Company Law of 1921, is amended by adding a  
17 section to read:

18 Section 635.2. Prescription Drug Coverage.--(a) No health  
19 insurance policy which covers prescription drug benefits shall  
20 be issued, amended, delivered, or renewed in this Commonwealth  
21 if the plan limits or excludes coverage for a drug on the basis

1 that the drug is prescribed for a use that is different from the  
2 use for which that drug has been approved for marketing by the  
3 FDA, provided that all of the following conditions have been  
4 met:

5 (1) The drug is approved by the FDA for at least one  
6 condition.

7 (2) The drug is prescribed by a participating licensed  
8 health care professional for the treatment of a life-threatening  
9 condition or a chronic and seriously debilitating condition.

10 (3) The drug has been recognized for treatment of that  
11 condition by one of the following:

12 (i) The American Hospital Formulary Service Drug  
13 Information.

14 (ii) The American Medical Association Drug Evaluations.

15 (iii) The United States Pharmacopoeia Drug Information,  
16 Volume 1, "Drug Information for the Health Care Professional."

17 (iv) The drug is supported by clinical research for the  
18 treatment of that condition in scientific, medical or  
19 pharmaceutical publications in which original manuscripts are  
20 published, only after having been critically reviewed for  
21 scientific accuracy, validity and reliability by unbiased  
22 experts.

23 (b) Nothing in this section shall be construed to prohibit  
24 the use of a formulary, copayment, technology assessment panel,  
25 or similar mechanism as a means for appropriately controlling  
26 utilization of a drug prescribed for a use different from the  
27 use for which the drug has been approved for marketing by the  
28 FDA.

29 (c) Nothing in this section shall be construed to require:

30 (1) coverage of a new drug or biological product not

otherwise approved for a use by the FDA;

(2) coverage of a disease or condition that is not a covered condition under the policy, subscriber contract or certificate;

(3) aggregate payments in excess of the amounts required to be paid under the policy, subscriber contract or certificate;

(4) modification of any coinsurance or copayment requirements used to manage a formulary; or

(5) coverage for FDA-approved drugs excluded from an enrollee's formulary coverage, except as such drugs may be available through any prior authorization procedures.

(d) Coverage of a drug required by this section shall include coverage of medically necessary services associated with the administration of the drug when they are covered benefits under the policy.

(e) As used in this section:

(1) "Life-threatening" means either or both of the following:

(i) diseases or conditions where the likelihood of death is high unless the course of the disease is interrupted; or

(ii) diseases or conditions with potentially fatal outcomes, where the end point of clinical intervention is survival.

(2) "Chronic and seriously debilitating" means diseases or conditions that require ongoing treatment to maintain remission or prevent deterioration and cause significant long-term morbidity.

(3) "Health insurance policy" means any individual or group health, sickness or accident insurance policy, subscriber contract or certificate issued by any entity subject to:

(i) this act;

(ii) 40 Pa.C.S. Ch. 61 (relating to hospital plan

1 corporations);

2 (iii) 40 Pa.C.S. Ch. 63 (relating to professional health  
3 services plan corporations); or

4 (iv) the act of December 29, 1972 (P.L.1701, No.364), known  
5 as the "Health Maintenance Organization Act."

6 (4) "FDA" means the Food and Drug Administration of the  
7 United States Department of Health and Human Services.

8 Section 2. This act shall take effect in 60 days.