93RD GENERAL ASSEMBLY

State of Illinois

2003 and 2004

Introduced 2/5/2004, by Mattie Hunter

SYNOPSIS AS INTRODUCED:

305 ILCS 5/5-5.12

from Ch. 23, par. 5-5.12

Amends the Illinois Public Aid Code. Provides that the Department shall not impose requirements for prior approval based on a preferred drug list for any short-acting beta agonists used as rescue therapy in the treatment of life-threatening conditions due to acute bronchospasm in patients with reversible obstructive airway disease that results from asthma, chronic obstructive pulmonary disease or emphysema.

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SB2788

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AN ACT in relation to public aid.

2 Be it enacted by the People of the State of Illinois, 3 represented in the General Assembly:

4 Section 5. The Illinois Public Aid Code is amended by 5 changing Sections 5-5.12 and 9A-11.5 as follows:

- 6 (305 ILCS 5/5-5.12) (from Ch. 23, par. 5-5.12)
- 7 Sec. 5-5.12. Pharmacy payments.

8 (a) Every request submitted by a pharmacy for reimbursement 9 under this Article for prescription drugs provided to a 10 recipient of aid under this Article shall include the name of 11 the prescriber or an acceptable identification number as 12 established by the Department.

(b) Pharmacies providing prescription drugs under this 13 14 Article shall be reimbursed at a rate which shall include a 15 professional dispensing fee as determined by the Illinois current 16 Department, plus the acquisition cost of the 17 prescription drug dispensed. The Illinois Department shall update its information on the acquisition costs of all 18 19 prescription drugs no less frequently than every 30 days. 20 However, the Illinois Department may set the rate of 21 reimbursement for the acquisition cost, by rule, at a 22 percentage of the current average wholesale acquisition cost.

23 (c) Reimbursement under this Article for prescription drugs shall be limited to reimbursement for 4 brand-name 24 25 prescription drugs per patient per month. This subsection 26 applies only if (i) the brand-name drug was not prescribed for an acute or urgent condition, (ii) the brand-name drug was not 27 28 prescribed for Alzheimer's disease, arthritis, diabetes, 29 HIV/AIDS, a mental health condition, or respiratory disease, 30 and (iii) a therapeutically equivalent generic medication has been approved by the federal Food and Drug Administration. 31

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(d) The Department shall not impose requirements for prior

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1 approval based on a preferred drug list for anti-retroviral, 2 anti-hemophilic factor concentrates, or any atypical 3 antipsychotics, conventional antipsychotics, or 4 anticonvulsants used for the treatment of serious mental 5 illnesses until 30 days after it has conducted a study of the 6 impact of such requirements on patient care and submitted a 7 report to the Speaker of the House of Representatives and the 8 President of the Senate.

9 (e) The Department shall not impose requirements for prior approval based on a preferred drug list for any short-acting 10 11 beta agonists used as rescue therapy in the treatment of 12 life-threatening conditions due to acute bronchospasm in patients with reversible obstructive airway disease that 13 results from asthma, chronic obstructive pulmonary disease or 14 15 emphysema. (Source: P.A. 92-597, eff. 6-28-02; 92-825, eff. 8-21-02; 16

17 93-106, eff. 7-8-03.)

Section 99. Effective date. This Act takes effect upon becoming law.