

94TH GENERAL ASSEMBLY State of Illinois 2005 and 2006 HB0734

Introduced 2/1/2005, by Rep. William Delgado

SYNOPSIS AS INTRODUCED:

305 ILCS 5/5-5.12

from Ch. 23, par. 5-5.12

Amends the Illinois Public Aid Code. In provisions concerning pharmacy payments under the Medicaid program, provides that the Department of Public Aid may not impose prior approval requirements for antidepressants or other drugs used in treating serious mental illnesses (deletes the provision allowing imposition of such requirements after conducting an impact study). Requires continuous access to medications. Makes other changes, Effective immediately.

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FISCAL NOTE ACT MAY APPLY

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1 AN ACT concerning public aid

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

- Section 5. The Illinois Public Aid Code is amended by changing Section 5-5.12 as follows:
- 6 (305 ILCS 5/5-5.12) (from Ch. 23, par. 5-5.12)
- 7 Sec. 5-5.12. Pharmacy payments.
 - (a) Every request submitted by a pharmacy for reimbursement under this Article for prescription drugs provided to a recipient of aid under this Article shall include the name of the prescriber or an acceptable identification number as established by the Department.
 - (b) Pharmacies providing prescription drugs under this Article shall be reimbursed at a rate which shall include a professional dispensing fee as determined by the Illinois current acquisition Department, plus the cost of prescription drug dispensed. The Illinois Department shall update its information on the acquisition costs of all prescription drugs no less frequently than every 30 days. However, the Illinois Department may set the rate reimbursement for the acquisition cost, by rule, at percentage of the current average wholesale acquisition cost.
 - (c) Reimbursement under this Article for prescription drugs shall be limited to reimbursement for 4 brand-name prescription drugs per patient per month. This subsection applies only if (i) the brand-name drug was not prescribed for an acute or urgent condition, (ii) the brand-name drug was not prescribed for Alzheimer's disease, arthritis, diabetes, HIV/AIDS, a mental health condition, or respiratory disease, and (iii) a therapeutically equivalent generic medication has been approved by the federal Food and Drug Administration.
 - (d) The Department shall not impose requirements for prior

- 1 approval based on a preferred drug list for anti-retroviral, 2 anti-hemophilic factor concentrates, or any atypical 3 antipsychotics, conventional antipsychotics, 4 anticonvulsants, or antidepressants used for the treatment of 5 serious mental illnesses until 30 days after it has conducted a 6 study of the impact of such requirements on patient care and submitted a report to the Speaker of the House of 7 Representatives and the President of the Senate. The Department 8 9 shall adopt policies and implement procedures that ensure continuous access to medications 24 hours per day, 7 days per 10 11 week. In an emergency situation, when prior approval or an edit 12 override is not available, a pharmacy may dispense, and the Department shall pay for, up to a 72-hour supply of a drug 13 prescribed to an eligible recipient for a mental illness. 14 Nothing in this Section shall be construed as preventing the 15 16 Department from implementing other restrictions to ensure the 17 appropriate use of medications described in this subsection by persons with a serious mental illness, as supported by 18 19 evidence-based medicine. (Source: P.A. 92-597, eff. 6-28-02; 92-825, eff. 8-21-02; 20 93-106, eff. 7-8-03.) 21
- 22 Section 99. Effective date. This Act takes effect upon 23 becoming law.