- 1 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 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.
- 8 (a) Every request submitted by a pharmacy for
- 9 reimbursement under this Article for prescription drugs
- 10 provided to a recipient of aid under this Article shall
- 11 include the name of the prescriber or an acceptable
- identification number as established by the Department.
- 13 (b) Pharmacies providing prescription drugs under this
- 14 Article shall be reimbursed at a rate which shall include a
- 15 professional dispensing fee as determined by the Illinois
- 16 Department, plus the current acquisition cost of the
- 17 prescription drug dispensed.
- 18 <u>In the fiscal year beginning July 1, 2003 and every 2</u>
- 19 years thereafter, the Illinois Department shall conduct a
- 20 <u>survey of pharmacies providing prescription drugs under this</u>
- 21 Article to assess the appropriate level of dispensing fees to
- be paid under this Section.
- The Illinois Department shall update its information on
- 24 the acquisition costs of all prescription drugs no less
- 25 frequently than every 30 days. However, the Illinois
- 26 Department may set the rate of reimbursement for the
- 27 acquisition cost, by rule, at a percentage of the current
- 28 average wholesale acquisition cost.
- 29 (c) Reimbursement under this Article for prescription
- 30 drugs shall be limited to reimbursement for 4 brand-name
- 31 prescription drugs per patient per month. This subsection

- 1 applies only if (i) the brand-name drug was not prescribed
- 2 for an acute or urgent condition, (ii) the brand-name drug
- 3 was not prescribed for Alzheimer's disease, arthritis,
- 4 diabetes, HIV/AIDS, a mental health condition, or respiratory
- 5 disease, and (iii) a therapeutically equivalent generic
- 6 medication has been approved by the federal Food and Drug
- 7 Administration.
- 8 (d) The Department shall not impose requirements for
- 9 prior approval based on a preferred drug list for
- 10 anti-retroviral or any atypical antipsychotics, conventional
- 11 antipsychotics, or anticonvulsants used for the treatment of
- serious mental illnesses until 30 days after it has conducted
- 13 a study of the impact of such requirements on patient care
- 14 and submitted a report to the Speaker of the House of
- 15 Representatives and the President of the Senate.
- 16 (Source: P.A. 92-597, eff. 6-28-02; 92-825, eff. 8-21-02;
- 17 revised 9-19-02.)