MAINE STATE LEGISLATURE

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120th MAINE LEGISLATURE

FIRST REGULAR SESSION-2001

Legislative Document

No. 1651

S.P. 528

In Senate, March 15, 2001

An Act to Preserve Maine Pharmacies.

Reference to the Committee on Health and Human Services suggested and ordered printed.

JOY J. O'BRIEN Secretary of the Senate

Presented by Senator MITCHELL of Penobscot. Cosponsored by Representative FULLER of Manchester and

Senators: ABROMSON of Cumberland, LaFOUNTAIN of York, TURNER of Cumberland,

Representative: LOVETT of Scarborough.

Be it enacted by the People of the State of Maine as follows:

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Sec. 1. 22 MRSA §254, sub-§8, as amended by PL 1999, c. 786, Pt. B, §1, is further amended to read:

- Drug rebate program. Effective May 1, 1992, payment must be denied for drugs from manufacturers that do not enter into a rebate agreement with the department for prescription drugs included in the list of approved drugs under this Each agreement must provide that the pharmaceutical program. manufacturer make rebate payments for both the basic supplemental components of the program to the department according to the following schedule.
 - B. For the quarters beginning October 1, 1992, the rebate percentage is equal to the percentage recommended by the federal Health Care Financing Administration of the manufacturer's wholesale price for the total number of dosage units of each form and strength of a prescription drug that the department reports as reimbursed to providers of prescription drugs, provided payments are not due until 30 days following the manufacturer's receipt of utilization data supplied by the department, including the number of dosage units reimbursed to providers of prescription drugs during the period for which payments are due.
 - Beginning October 1, 1998, the department shall seek to an aggregate rebate amount from achieve all agreements that is 6 percentage points higher than that required by paragraph B of this subsection, provided such rebates result in a net increase in the rebate revenue available to the elderly low-cost drug program. event the department is not able to achieve the rebate amount required by this paragraph without compromising the best interest of recipients of the elderly low-cost drug program, it shall report to the joint standing committee of the Legislature having jurisdiction over health and human services matters and the joint standing committee of the Legislature having jurisdiction over appropriations financial affairs in the First Regular Session of the 119th Legislature.

Upon receipt of data from the department, the pharmaceutical manufacturer shall calculate the quarterly payment. If a discrepancy is discovered, the department may, at its expense, hire a mutually agreed-upon independent auditor to verify the pharmaceutical manufacturer's calculation. If a discrepancy is still found, the pharmaceutical manufacturer shall justify its calculation or make payment to the department for any additional amount due. The pharmaceutical manufacturer may, at its expense,

hire a mutually agreed-upon independent auditor to verify the accuracy of the utilization data provided by the department. If a discrepancy is discovered, the department shall justify its data or refund any excess payment to the pharmaceutical manufacturer.

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If the dispute over the rebate amount is not resolved, a request for a hearing with supporting documentation must be submitted to the Administrative Hearings Unit. Failure to resolve the dispute may be cause for terminating the drug rebate agreement and denying payment to the pharmaceutical manufacturer for any drugs.

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Any prescription drug of a manufacturer that does not enter into an agreement is not reimbursable unless the department determines the prescription drug is essential.

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All prescription drugs of a pharmaceutical manufacturer that enters into an agreement pursuant to this subsection that appear on the list of approved drugs under this program must be immediately available and the cost of the drugs must be reimbursed and is not subject to any restrictions or prior authorization requirements,——except——as——previded——in——this paragraph. If—the—commissioner—establishes maximum—retail—prices for—prescription—drugs—pursuant—to—section—2693,—the—department shall—adept—rules—for—the—elderly—low—cost—drug—program—requiring the—use—of—a—drug—formulary—and—prior—authorization—for—the dispensing—of—certain—drugs—to—be—listed—en—a—formulary.——Rules adepted—pursuant—to—this—paragraph—are—routine—technical—rules—as defined—in—Title—5,—ehapter—375,—subchapter—II—A.

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. Sec. 2. 22 MRSA §2681, first \P , as enacted by PL 1999, c. 786, Pt. A, \S 3, is amended to read:

The Maine Rx Program, referred to in this subchapter as the "program," is established to reduce prescription drug prices for residents of the State. The program is designed for the State to utilize manufacturer rebates and—pharmacy—discounts to reduce prescription drug prices. In implementing the program, the State shall serve as a pharmacy benefit manager in establishing rebates and-discounts on behalf of qualified residents.

Sec. 3. 22 MRSA §2681, sub-§2, ¶B, as enacted by PL 1999, c. 786, Pt. A, §3, is repealed.

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Sec. 4. 22 MRSA §2681, sub-§2, $\P G$, as enacted by PL 1999, c. 786, Pt. A, §3, is amended to read:

G. "Secondary-discounted <u>Discounted</u> price" means a price that is equal to or less than the initial-discounted retail

price minus the amount of any rebate paid by the State to 2 the participating retail pharmacy. Sec. 5. 22 MRSA §2681, sub-§5, ¶B, as enacted by PL 1999, c. 4 786, Pt. A, §3, is repealed. 6 Sec. 6. 22 MRSA §2681, sub-§5, ¶C, as enacted by PL 1999, c. 786, Pt. A, §3, is amended to read: 8 No later than October 1, 2001, a participating retail 10 pharmacy shall offer the secondary discounted price. 12 Sec. 7. 22 MRSA §2693, as enacted by PL 1999, c. 786, Pt. A, §3, is repealed. 14 Sec. 8. 22 MRSA §3173, 13th ¶, as repealed and replaced by PL 16 1979, c. 127, §144, is amended to read: 18 The Department of Human Services may establish fee schedules governing reimbursement for services provided under this chapter. 20 In establishing the fee schedules, the department shall consult with individual providers and their representative associations. 22 The fee schedules shall-be are subject to annual review. 24 schedule for pharmacies must include, in addition to the reimbursement, payment of a dispensing fee in the amount of \$6.50 26 for each prescription filled. The amount of this dispensing fee must increase by \$1 for each percentage point reduction in the 28 reimbursement for prescription drugs. Sec. 9. 22 MRSA §3174-Y, as enacted by PL 1999, c. 786, Pt. 30 B, §3, is repealed. 32 Sec. 10. 24 MRSA §2502, sub-§1-A, as enacted by PL 1985, c. 34 804, §§3 and 22, is amended to read: 1-A. Health care practitioner. "Health care practitioner" 36 means physicians and all others certified, registered or licensed in the healing arts, including, but not limited to, nurses, 38 podiatrists, optometrists, chiropractors, physical therapists, 40 dentists, psychologists, pharmacists and physicians' assistants. Sec. 11. 24-A MRSA §2703-A is enacted to read: 42 44 §2703-A. Identification card required 46 Every health insurer that issues a policy in this State that includes a prescription drug benefit shall provide an identification card to the policyholder that conforms to uniform 48 content and format requirements determined by rule by the 50 superintendent. Rules adopted pursuant to this section are

2	subchapter II-A.
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_	SUMMARY
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_	This bill makes the following changes to the laws governing
8	pharmacies.
10	1. It amends the Maine Rx Program to eliminate discounts
10	that are borne by pharmacies in this State, leaving the discounts
12	that are funded from rebates paid by drug manufacturers.
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14	2. It requires the Department of Human Services to provide
	a dispensing fee to pharmacies in a designated amount for
16	prescriptions that are filled for patients who participate in the
	Medicaid program.
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	3. It specifically names pharmacists as health care
20	practitioners under the Maine Health Security Act.
22	4. It requires health insurers who provide a pharmacy
	benefit to provide identification cards to their policyholders
24	that conform to a uniform format determined by the Superintendent
	of Insurance.

routine technical rules as defined in Title 5, chapter 375,