## **ACT No. 339**

SENATE BILL NO. 117

BY SENATOR MILLS

1	AN ACT
2	To amend and reenact R.S. 46:153.3(B)(2)(a)(iv), the introductory paragraph of (D)(2),
3	(D)(2)(c), (d), (e), (f), (g), (h), (k), and (p), and (D)(5)(b) and (c) and to repeal R.S
4	46:153.3(B)(1)(b), (c), (d), and (e), (B)(2)(d), (B)(3) and (4), (C)(2) and (3), and
5	(D)(5)(d) and (e), relative to the Medicaid Pharmaceutical and Therapeutics
6	Committee; to remove legislative intent and expired implementation restrictions; to
7	remove references to committees that no longer exist; to remove provisions that have
8	been sunset by subsequent legislation; to change the Medicaid Pharmaceutical and
9	Therapeutics Committee membership selection criteria; to change terminology; to
10	provide for an effective date; and to provide for related matters.
11	Be it enacted by the Legislature of Louisiana:
12	Section 1. R.S. $46:153.3(B)(2)(a)(iv)$ , the introductory paragraph of $(D)(2)$ , $(D)(2)(c)$ ,
13	(d), (e), (f), (g), (h), (k), and (p), and (D)(5)(b) and (c) are hereby amended and reenacted to
14	read as follows:
15	§153.3. Medical vendor reimbursements; allowable restrictions; peer-based
16	prescribing and dispensing practice patterns; Medicaid
17	Pharmaceutical and Therapeutics Committee
18	* * *
19	В.
20	* * *
21	(2)(a)
22	* * *
23	(iv) Involve medical personnel, including but not limited to pharmacists;
24	pharmacy technicians, nurses, and physicians.
25	* * *

1	D.
2	* * *
3	(2) Each nominating organization shall certify by affidavit that the practice
4	of each nominee involves either the care of or the supervision of the care of no less
5	than one hundred fifty Medicaid recipients. The committee shall be comprised of the
6	following persons:
7	* * *
8	(c) One practicing physician who is participating in the Title XIX program
9	as a family practitioner recommended from a list of three names submitted by the
10	Louisiana State Medical Society.
11	(d) One practicing physician who is participating in the Title XIX program
12	as an internal medicine specialist recommended from a list of three names submitted
13	by the Louisiana State Medical Society.
14	(e) One practicing physician who is participating in the Title XIX program
15	as a pediatrician recommended from a list of three names submitted by the Louisiana
16	State Medical Society.
17	(f) One practicing physician who is participating in the Title XIX program
18	as a surgeon recommended from a list of three names submitted by the Louisiana
19	State Medical Society.
20	(g) One practicing physician who is participating in the Title XIX program
21	as an obstetrics/gynecologist recommended from a list of three names submitted by
22	the Louisiana State Medical Society.
23	(h) Two practicing physicians who are participating in the Title XIX program
24	recommended from a list of six names submitted by the Louisiana Medical
25	Association.
26	* * *
27	(k) Two practicing pharmacists who are participating in the Title XIX drug
28	program recommended from a list of six names submitted by the Louisiana
29	Pharmacists Association. One pharmacist shall be an independent pharmacist
30	recommended by the Louisiana Independent Pharmacies Association and one

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1	pharmacist shall be a pharmacist representing a chain pharmacy recommended by
2	the Louisiana Pharmacists Association.
3	* * *
4	(p) One practicing physician who is participating in the Title XIX program
5	as a psychiatrist recommended from a list of three names submitted by the Louisiana
6	Psychiatric Medical Association.
7	* * *

(5)

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- (b) The committee shall be responsible for developing and maintaining a pharmacopoeia preferred drug list established in conjunction with a prior approval process as provided in Subparagraph (B)(2)(a) of this Section. The pharmacopoeia preferred drug list shall comply with all applicable state and federal laws, rules, and regulations. The committee may recommend additions and deletions to the pharmacopoeia preferred drug list and the pharmacopoeia preferred drug list may change in accordance with those recommendations. The committee shall also advise the secretary of the department on policy recommendations related to the prudent administration of the Medicaid drug program. The secretary shall assure that all actions of the committee comply with applicable state and federal laws, rules, and regulations prior to implementation or modification of the pharmacopoeia preferred drug list. The clinical decisions regarding the preferred drug list shall be made transparent through a written report that is publicly available. If the decision of the Medicaid Pharmaceutical and Therapeutics Committee is contrary to the clinical evidence found in labeling, drug compendia, or peer review literature, such decisions shall be justified in writing.
- (c) Any <u>new</u> drug approved by the United States Food and Drug Administration shall <u>may</u> be added to the <u>formulary preferred drug list</u> as soon as <u>when</u> it becomes commercially available <u>and the manufacturer enters into a</u> <u>federal medicaid drug rebate program if the department determines it is in the</u> <u>best interest of the medical assistance program</u>. The Medicaid Pharmaceutical and

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APPROVED:

Therapeutics Committee shall conduct an evidence-based analysis of the drug to
determine if the drug shall be maintained on the formulary preferred drug list. The
analysis shall include but not be limited to the medical evidence of the clinical
effectiveness of the drug as well as evidence of the cost-effectiveness of the drug in
treating illness and disease. The determination by the committee on any new drug
approval by the United States Food and Drug Administration shall be made no later
than ninety days after the drug becomes commercially available. Prior to a drug
being prior authorized, it must have been reviewed by the Medicaid Pharmaceutical
and Therapeutics Committee. When a new drug that is included in the Medicaid
Pharmaceutical and Therapeutics Committee process is approved by the United
States Food and Drug Administration, the drug shall be reviewed at the next
Medicaid Pharmaceutical and Therapeutics Committee meeting.
Section 2. R.S. 46:153.3(B)(1)(b), (c), (d), and (e), (B)(2)(d), (B)(3) and (4), (C)(2)
and (3), and (D)(5)(d) and (e) are hereby repealed in their entirety.
Section 3. This Act shall become effective upon signature by the governor or, if not
signed by the governor, upon expiration of the time for bills to become law without signature
by the governor, as provided by Article III, Section 18 of the Constitution of Louisiana. If
vetoed by the governor and subsequently approved by the legislature, this Act shall become
effective on the day following such approval.
PRESIDENT OF THE SENATE
SPEAKER OF THE HOUSE OF REPRESENTATIVES
GOVERNOR OF THE STATE OF LOUISIANA