

101ST GENERAL ASSEMBLY State of Illinois 2019 and 2020 SB3545

Introduced 2/14/2020, by Sen. Laura Fine

SYNOPSIS AS INTRODUCED:

215 ILCS 5/356z.43 new 215 ILCS 134/25 225 ILCS 85/26

Amends the Illinois Insurance Code. Provides that an individual or group policy of accident and health insurance amended, delivered, issued, or renewed after the effective date of the amendatory Act shall provide coverage for anti-epileptic drugs. Provides that coverage for anti-epileptic drugs may not impose a waiting period or any deductible, coinsurance, copayment, or other cost-sharing limitation. Defines "anti-epileptic drug", "epilepsy", and "seizure". Amends the Managed Care Reform and Patient Rights Act. Provides that anti-seizure prescription drugs may not be substituted with a generic drug under provisions of the Pharmacy Practice Act under which a pharmacist may substitute a therapeutically equivalent generic drug for a prescription drug. Amends the Pharmacy Practice Act. Provides that a pharmacist may not interchange an anti-epileptic drug or formulation of an anti-epileptic drug for the treatment of epilepsy. Provides that a prescribing physician shall document that such anti-epileptic drug or formulation of an anti-epileptic drug for the treatment of epilepsy is clinically necessary for the patient's optimal care. Removes provisions concerning notification and consent required when a physician substitutes a generic prescription in place of a brand-name anti-epileptic drug.

LRB101 17805 BMS 67235 b

1 AN ACT concerning regulation.

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

- Section 5. The Illinois Insurance Code is amended by adding Section 356z.43 as follows:
- 6 (215 ILCS 5/356z.43 new)
- 7 Sec. 356z.43. Coverage of anti-seizure prescription drugs.
- 8 (a) The General Assembly finds that this Section is
- 9 necessary for the immediate preservation of public peace,
- 10 <u>health</u>, and safety.
- 11 (b) In this Section:
- 12 <u>"Anti-epileptic drug" means (i)</u> a drug prescribed for the
- treatment of epilepsy or (ii) a drug used to treat or prevent
- 14 seizures.
- 15 <u>"Epilepsy" means a neurological condition characterized by</u>
- 16 recurrent seizures.
- "Seizure" means a brief disturbance in the electrical
- 18 activity of the brain.
- 19 (c) An individual or group policy of accident and health
- insurance amended, delivered, issued, or renewed in this State
- 21 after the effective date of this amendatory Act of the 101st
- 22 General Assembly shall provide coverage for anti-epileptic
- drugs.

1	(d) Coverage required under this Section may not impose a
2	waiting period or any deductible, coinsurance, copayment, or
3	other cost-sharing limitation that is greater than that
4	required for other coverage under the policy.

Section 10. The Managed Care Reform and Patient Rights Act is amended by changing Section 25 as follows:

(215 ILCS 134/25)

- Sec. 25. Transition of services.
- 9 (a) A health care plan shall provide for continuity of care for its enrollees as follows:
 - (1) If an enrollee's physician leaves the health care plan's network of health care providers for reasons other than termination of a contract in situations involving imminent harm to a patient or a final disciplinary action by a State licensing board and the physician remains within the health care plan's service area, the health care plan shall permit the enrollee to continue an ongoing course of treatment with that physician during a transitional period:
 - (A) of 90 days from the date of the notice of physician's termination from the health care plan to the enrollee of the physician's disaffiliation from the health care plan if the enrollee has an ongoing course of treatment; or

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plan:

1	(B) if the enrollee has entered the third trimester
2	of pregnancy at the time of the physician's
3	disaffiliation, that includes the provision of
4	post-partum care directly related to the delivery.
5	(2) Notwithstanding the provisions in item (1) of this
6	subsection, such care shall be authorized by the health
7	care plan during the transitional period only if the
8	physician agrees:
9	(A) to continue to accept reimbursement from the
10	health care plan at the rates applicable prior to the
11	start of the transitional period;
12	(B) to adhere to the health care plan's quality
13	assurance requirements and to provide to the health
14	care plan necessary medical information related to
15	such care; and
16	(C) to otherwise adhere to the health care plan's
17	policies and procedures, including but not limited to
18	procedures regarding referrals and obtaining
19	preauthorizations for treatment.
20	(3) During an enrollee's plan year, a health care plan
21	shall not remove a drug from its formulary or negatively
22	change its preferred or cost-tier sharing unless, at least
23	60 days before making the formulary change, the health care

(A) provides general notification of the change in

its formulary to current and prospective enrollees;

(B)	d:	irectly	noti	notifies		ollees	3 (currently			
receivi	ing co	overage f	for the	drug,	inc	luding	ginf	format	tion		
on the	spec	ific dru	gs invo	olved	and t	the st	eps	they	may		
take	to	request	COV	erage	de	termir	natio	ns	and		
exceptions, including a statement that a certification									tion		
of med	ical	necessi	ty by	the e	enrol	lee's	pre	escrik	oing		
provider will result in continuation of coverage at the											
existing level; and											

(C) directly notifies by first class mail and through an electronic transmission, if available, the prescribing provider of all health care plan enrollees currently prescribed the drug affected by the proposed change; the notice shall include a one-page form by which the prescribing provider can notify the health care plan by first class mail that coverage of the drug for the enrollee is medically necessary.

The notification in paragraph (C) may direct the prescribing provider to an electronic portal through which the prescribing provider may electronically file a certification to the health care plan that coverage of the drug for the enrollee is medically necessary. The prescribing provider may make a secure electronic signature beside the words "certification of medical necessity", and this certification shall authorize continuation of coverage for the drug.

If the prescribing provider certifies to the health

care plan either in writing or electronically that the drug is medically necessary for the enrollee as provided in paragraph (C), a health care plan shall authorize coverage for the drug prescribed based solely on the prescribing provider's assertion that coverage is medically necessary, and the health care plan is prohibited from making modifications to the coverage related to the covered drug, including, but not limited to:

- (i) increasing the out-of-pocket costs for the covered drug;
- (ii) moving the covered drug to a more restrictive
 tier; or
- (iii) denying an enrollee coverage of the drug for which the enrollee has been previously approved for coverage by the health care plan.

Nothing in this item (3) prevents a health care plan from removing a drug from its formulary or denying an enrollee coverage if the United States Food and Drug Administration has issued a statement about the drug that calls into question the clinical safety of the drug, the drug manufacturer has notified the United States Food and Drug Administration of a manufacturing discontinuance or potential discontinuance of the drug as required by Section 506C of the Federal Food, Drug, and Cosmetic Act, as codified in 21 U.S.C. 356c, or the drug manufacturer has removed the drug from the market.

Nothing in this item (3) prohibits a health care plan, by contract, written policy or procedure, or any other agreement or course of conduct, from requiring a pharmacist to effect substitutions of prescription drugs consistent with Section 19.5 of the Pharmacy Practice Act, under which a pharmacist may substitute an interchangeable biologic for a prescribed biologic product, and Section 25 of the Pharmacy Practice Act, under which a pharmacist may select a generic drug determined to be therapeutically equivalent by the United States Food and Drug Administration and in accordance with the Illinois Food, Drug and Cosmetic Act, with the exception of anti-seizure prescription drugs, which may not be substituted with a generic drug.

This item (3) applies to a policy or contract that is amended, delivered, issued, or renewed on or after January 1, 2019. This item (3) does not apply to a health plan as defined in the State Employees Group Insurance Act of 1971 or medical assistance under Article V of the Illinois Public Aid Code.

- (b) A health care plan shall provide for continuity of care for new enrollees as follows:
 - (1) If a new enrollee whose physician is not a member of the health care plan's provider network, but is within the health care plan's service area, enrolls in the health care plan, the health care plan shall permit the enrollee to continue an ongoing course of treatment with the

enrollee's current physician during a transitional period:

- (A) of 90 days from the effective date of enrollment if the enrollee has an ongoing course of treatment; or
- (B) if the enrollee has entered the third trimester of pregnancy at the effective date of enrollment, that includes the provision of post-partum care directly related to the delivery.
- (2) If an enrollee elects to continue to receive care from such physician pursuant to item (1) of this subsection, such care shall be authorized by the health care plan for the transitional period only if the physician agrees:
 - (A) to accept reimbursement from the health care plan at rates established by the health care plan; such rates shall be the level of reimbursement applicable to similar physicians within the health care plan for such services;
 - (B) to adhere to the health care plan's quality assurance requirements and to provide to the health care plan necessary medical information related to such care; and
 - (C) to otherwise adhere to the health care plan's policies and procedures including, but not limited to procedures regarding referrals and obtaining preauthorization for treatment.

- 1 (c) In no event shall this Section be construed to require
- 2 a health care plan to provide coverage for benefits not
- 3 otherwise covered or to diminish or impair preexisting
- 4 condition limitations contained in the enrollee's contract. In
- 5 no event shall this Section be construed to prohibit the
- 6 addition of prescription drugs to a health care plan's list of
- 7 covered drugs during the coverage year.
- 8 (Source: P.A. 100-1052, eff. 8-24-18.)
- 9 Section 15. The Pharmacy Practice Act is amended by
- 10 changing Section 26 as follows:
- 11 (225 ILCS 85/26)
- 12 (Section scheduled to be repealed on January 1, 2023)
- 13 Sec. 26. Anti-epileptic drug product selection prohibited.
- 14 (a) The General Assembly finds that this Section is
- 15 necessary for the immediate preservation of the public peace,
- 16 health, and safety.
- 17 (b) In this Section:
- "Anti-epileptic drug" means (i) any drug prescribed for the
- 19 treatment of epilepsy or (ii) a drug used to treat or prevent
- 20 seizures.
- "Epilepsy" means a neurological condition characterized by
- 22 recurrent seizures.
- "Seizure" means a brief disturbance in the electrical
- 24 activity of the brain.

- (c) A When the prescribing physician has indicated on the original prescription "may not substitute", a pharmacist may not interchange an anti-epileptic drug or formulation of an anti-epileptic drug for the treatment of epilepsy. The prescribing physician shall document that such anti-epileptic drug or formulation of an anti-epileptic drug for the treatment of epilepsy is clinically necessary for the patient's optimal care without notification and the documented consent of the prescribing physician and the patient or the patient's parent, legal guardian, or spouse. This Section does not apply to medication orders issued for anti-epileptic drugs for any in-patient care in a licensed hospital.
- (d) (Blank). If a pharmacist substitutes any generic prescription in place of a brand-name anti-epileptic drug, then the pharmacist shall provide written notice to the patient no later than the time the prescription is dispensed.
- 17 (Source: P.A. 97-456, eff. 1-1-12.)