LC000379

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STATE OF RHODE ISLAND

IN GENERAL ASSEMBLY

JANUARY SESSION, A.D. 2015

AN ACT

RELATING TO INSURANCE - ACCESS TO ABUSE-DETERRENT PAIN MEDICATIONS

<u>Introduced By:</u> Senators Miller, Crowley, Sosnowski, Goldin, and Jabour

Date Introduced: February 05, 2015

Referred To: Senate Health & Human Services

It is enacted by the General Assembly as follows:

considered an abuse-deterrent formulation.

1	SECTION 1. Chapter 27-18 of the General Laws entitled "Accident and Sickness
2	Insurance Policies" is hereby amended by adding thereto the following section:
3	27-18-82. Access to Abuse-Deterrent Pain Medications (a) Every individual or
4	group health insurance contract or every individual or group hospital or medical expense
5	insurance policy, plan, or group policy issued for delivery, or renewed in this state on or after the
6	passage of this act that provides coverage for prescription drugs shall not require, as a condition
7	of coverage:
8	(1) Use of an opioid drug not indicated by the United States Food and Drug
9	Administration (FDA) for the condition being treated prior to use of a non-opioid drug that is
10	approved by the FDA for the condition being treated; or
11	(2) Use of a non-abuse-deterrent formulation prior to use of an abuse-deterrent
12	formulation for the treatment of pain.
13	(b) For the purpose of this section:
14	(1) "Abuse-deterrent formulation" means a drug used to treat pain that is considered to
15	have abuse-deterrent properties if the FDA determines there is sufficient evidence to support
16	abuse-deterrent claims based on published FDA guidance.
17	(2) "Non-abuse-deterrent formulation" means a drug used to treat pain that is not

(c) Health insurance contracts, plans or policies to which this section applies may require

1	an insured to use, prior to using a brand name prescription drug prescribed by a licensed
2	prescriber a therapeutically equivalent generic drug, unless, pursuant to §§ 5-19.1-19, 5-37-18.1
3	and 21-31-15(b), the prescriber indicates "brand name necessary" on the prescription form, or if
4	the prescriber gives oral direction to that effect to the dispensing pharmacist.
5	SECTION 2. Chapter 27-19 of the General Laws entitled "Nonprofit Hospital Service
6	Corporations" is hereby amended by adding thereto the following section:
7	27-19-73. Access to Abuse-Deterrent Pain Medications (a) Every individual or
8	group health insurance contract or every individual or group hospital or medical expense
9	insurance policy, plan, or group policy issued for delivery, or renewed in this state on or after the
10	passage of this act that provides coverage for prescription drugs shall not require, as a condition
11	of coverage:
12	(1) Use of an opioid drug not indicated by the United States Food and Drug
13	Administration (FDA) for the condition being treated prior to use of a non-opioid drug that is
14	approved by the FDA for the condition being treated; or
15	(2) Use of a non-abuse-deterrent formulation prior to use of an abuse-deterrent
16	formulation for the treatment of pain.
17	(b) For the purpose of this section:
18	(1) "Abuse-deterrent formulation" means a drug used to treat pain that is considered to
19	have abuse-deterrent properties if the FDA determines there is sufficient evidence to support
20	abuse-deterrent claims based on published FDA guidance.
21	(2) "Non-abuse-deterrent formulation" means a drug used to treat pain that is not
22	considered an abuse-deterrent formulation.
23	(c) Health insurance contracts, plans or policies to which this section applies may require
24	an insured to use, prior to using a brand name prescription drug prescribed by a licensed
25	prescriber, a therapeutically equivalent generic drug, unless pursuant to §§ 5-19.1-19, 5-37-18.1
26	and 21-31-15(b), the prescriber indicates "brand name necessary" on the prescription form, or if
27	the prescriber gives oral direction to that effect to the dispensing pharmacist.
28	SECTION 3. Chapter 27-20 of the General Laws entitled "Nonprofit Medical Service
29	Corporations" is hereby amended by adding thereto the following section:
30	27-20-69. Access to Abuse-Deterrent Pain Medications (a) Every individual or
31	group health insurance contract, or every individual or group hospital or medical expense
32	insurance policy, plan, or group policy issued for delivery, or renewed in this state on or after the
33	passage of this act that provides coverage for prescription drugs shall not require, as a condition
34	of coverage:

1	(1) Use of an opioid drug not indicated by the Officed States Food and Drug
2	Administration (FDA) for the condition being treated prior to use of a non-opioid drug that is
3	approved by the FDA for the condition being treated; or
4	(2) Use of a non-abuse-deterrent formulation prior to use of an abuse-deterrent
5	formulation for the treatment of pain.
6	(b) For the purpose of this section:
7	(1) "Abuse-deterrent formulation" means a drug used to treat pain that is considered to
8	have abuse-deterrent properties if the FDA determines there is sufficient evidence to support
9	abuse-deterrent claims based on published FDA guidance.
10	(2) "Non-abuse-deterrent formulation" means a drug used to treat pain that is not
11	considered an abuse-deterrent formulation.
12	(c) Health insurance contracts, plans or policies to which this section applies may require
13	an insured to use, prior to using a brand name prescription drug prescribed by a licensed
14	prescriber, a therapeutically equivalent generic drug, unless, pursuant to §§ 5-19.1-19, 5-3 7-18.1
15	and 21-31-15(b), the prescriber indicates "brand name necessary" on the prescription form, or if
16	the prescriber gives oral direction to that effect to the dispensing pharmacist.
17	SECTION 4. Chapter 27-41 of the General Laws entitled "Health Maintenance
18	Organizations" is hereby amended by adding thereto the following section:
19	27-41-86. Access to Abuse-Deterrent Pain Medications (a) Every individual or
20	group health insurance contract, or every individual or group hospital or medical expense
21	insurance policy, plan, or group policy issued for delivery, or renewed in this state on or after the
22	passage of this act that provides coverage for prescription drugs shall not require, as a condition
23	of coverage:
24	(1) Use of an opioid drug not indicated by the United States Food and Drug
25	Administration (FDA) for the condition being treated prior to use of a non-opioid drug that is
26	approved by the FDA for the condition being treated; or
27	(2) Use of a non-abuse-deterrent formulation prior to use of an abuse-deterrent
28	formulation for the treatment of pain.
29	(b) For the purpose of this section:
30	(1) "Abuse-deterrent formulation" means a drug used to treat pain that is considered to
31	have abuse-deterrent properties if the FDA determines there is sufficient evidence to support
32	abuse-deterrent claims based on published FDA guidance.
33	(2) "Non-abuse-deterrent formulation" means a drug used to treat pain that is not
34	considered an abuse-deterrent formulation.

- 1 (c) Health insurance contracts, plans or policies to which this section applies may require
 2 an insured to use, prior to using a brand name prescription drug prescribed by a licensed
 3 prescriber, a therapeutically equivalent generic drug, unless, pursuant to §§ 5-19.1-19, 5-37-18.1
 4 and 21-31-15(b), the prescriber indicates "brand name necessary" on the prescription form, or if
 5 the prescriber gives oral direction to that effect to the dispensing pharmacist.
- 6 SECTION 5. This act shall take effect upon passage.

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EXPLANATION

BY THE LEGISLATIVE COUNCIL

OF

AN ACT

RELATING TO INSURANCE - ACCESS TO ABUSE-DETERRENT PAIN MEDICATIONS

1 This act would prevent health insurance policies, plans or contracts that provide coverage 2 for prescription drugs, from requiring a beneficiary to use an opioid drug not indicated by the 3 FDA for the condition being treated prior to the use of a non-opioid drug that is approved by the 4 FDA for the condition being treated, or to use a non-abuse-deterrent formulation prior to using an abuse-deterrent formulation. 5 6 This act would take effect upon passage.

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