STATE OF RHODE ISLAND

IN GENERAL ASSEMBLY

JANUARY SESSION, A.D. 2020

AN ACT

RELATING TO INSURANCE

Introduced By: Senators Crowley, Lombardo, Conley, and Ruggerio

<u>Date Introduced:</u> February 05, 2020

Referred To: Senate Health & Human Services

It is enacted by the General Assembly as follows:

SECTION 1. Section 27-18-50 of the General Laws in Chapter 27-18 entitled "Accident

and Sickness Insurance Policies" is hereby amended to read as follows:

27-18-50. Drug coverage.

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- (a) Any accident and sickness insurer that utilizes a formulary of medications for which coverage is provided under an individual or group-plan, master contract shall require any physician or other person authorized by the department of health to prescribe medication to prescribe from the formulary. A physician or other person authorized by the department of health to prescribe medication shall be allowed to prescribe medications previously on, or not on, the accident and sickness insurer's formulary if he or she believes that the prescription of the non-formulary medication is medically necessary. An accident and sickness insurer shall be required to provide coverage for a non-formulary medication only when the non-formulary medication meets the accident and sickness insurer's medical-exception criteria for the coverage of that medication.
- (b) An accident and sickness insurer's medical exception criteria for the coverage of non-formulary medications shall be developed in accordance with § 23-17.13-3(e)(3) 27-18.8-3(b)(5).
- (c) Any subscriber who is aggrieved by a denial of benefits to be provided under this section may appeal the denial in accordance with the rules and regulations promulgated by the department of health commissioner pursuant to chapter 17.12 of title 23 chapter 18.9 of title 27.
 - (d) Prior to removing a prescription drug from its plan's formulary or making any change

1	in the preferred of flered, cost sharing status of a covered prescription drug, an accident and
2	sickness insurer must provide at least thirty (30) days' notice to authorized prescribers by
3	established communication methods of policy and program updates and by updating available
4	references on web-based publications. All adversely affected members must be provided at least
5	thirty (30) days' notice prior to the date such change becomes effective by a direct notification:
6	(i) The written or electronic notice must contain the following information:
7	(A) The name of the affected prescription drug;
8	(B) Whether the plan is removing the prescription drug from the formulary, or changing
9	its preferred or tiered, cost-sharing status; and
.0	(C) The means by which subscribers may obtain a coverage determination or medical
1	exception, in the case of drugs that will require prior authorization or are formulary exclusions
2	respectively.
.3	(d) A health benefit plan issuer may modify drug coverage provided under a health
4	benefit plan if:
.5	(1) The modification occurs at the time of coverage renewal;
6	(2) The modification is effective uniformly among all group health benefit plan sponsors
7	covered by identical or substantially identical health benefit plans or all individuals covered by
8	identical or substantially identical individual health benefit plans, as applicable; and
9	(3) Not later than the sixtieth day before the date the modification is effective, the issuer
20	provides written notice of the modification to the commissioner, each affected group health
21	benefit plan sponsor, each affected enrollee in an affected group health benefit plan, and each
22	affected individual health benefit plan holder.
23	(e) Modifications affecting drug coverage that require written or electronic notice under
24	subsection (d) of this section, include:
25	(1) Removing a drug from a formulary;
26	(2) Adding a requirement that an enrollee receive prior authorization for a drug;
27	(3) Imposing or altering a quantity limit for a drug;
28	(4) Imposing a step-therapy restriction for a drug; and
29	(5) Moving a drug to a higher cost-sharing tier unless a generic drug alternative to the
80	drug is available.
31	(ii)(f) An accident and sickness insurer may immediately remove from its plan
32	formularies covered prescription drugs deemed unsafe by the accident and sickness insurer or the
33	Food and Drug Administration, or removed from the market by their manufacturer, without
34	meeting the requirements of this section.

1	(e)(g) This section shall not apply to insurance coverage providing benefits for: (1)
2	Hospital confinement indemnity; (2) Disability income; (3) Accident only; (4) Long-term care;
3	(5) Medicare supplement; (6) Limited-benefit health; (7) Specified-disease indemnity; (8)
4	Sickness or bodily injury or death by accident or both; or (9) Other limited-benefit policies.
5	SECTION 2. Section 27-19-42 of the General Laws in Chapter 27-19 entitled "Nonprofit
6	Hospital Service Corporations" is hereby amended to read as follows:
7	27-19-42. Drug coverage.
8	(a) Any nonprofit, hospital-service corporation that utilizes a formulary of medications
9	for which coverage is provided under an individual or group-plan, master contract shall require
10	any physician or other person authorized by the department of health to prescribe medication to
11	prescribe from the formulary. A physician or other person authorized by the department of health
12	to prescribe medication shall be allowed to prescribe medications previously on, or not on, the
13	nonprofit, hospital-service corporation's formulary if he or she believes that the prescription of
14	the non-formulary medication is medically necessary. A nonprofit, hospital-service corporation
15	shall be required to provide coverage for a non-formulary medication only when the non-
16	formulary medication meets the nonprofit, hospital-service corporation's medical-exception
17	criteria for the coverage of that medication.
18	(b) A nonprofit, hospital-service corporation's medical-exception criteria for the coverage
19	of non-formulary medications shall be developed in accordance with § 23-17.13-3(e)(3) 27-18.8-
20	<u>3(b)(5)</u> .
21	(c) Any subscriber who is aggrieved by a denial of benefits to be provided under this
22	section may appeal the denial in accordance with the rules and regulations promulgated by the
23	department of health commissioner pursuant to chapter 17.12 of title 23 chapter 18.9 of title 27.
24	(d) Prior to removing a prescription drug from its plan's formulary or making any change
25	in the preferred or tiered cost sharing status of a covered prescription drug, a nonprofit, hospital-
26	service corporation must provide at least thirty (30) days' notice to authorized prescribers by
27	established communication methods of policy and program updates and by updating available
28	references on web-based publications. All adversely affected members must be provided at least
29	thirty (30) days' notice prior to the date such change becomes effective by a direct notification:
30	(i) The written or electronic notice must contain the following information:
31	(A) The name of the affected prescription drug;
32	(B) Whether the plan is removing the prescription drug from the formulary, or changing
33	its preferred or tiered, cost-sharing status; and
34	(C) The means by which subscribers may obtain a coverage determination or medical

1	exception, in the case of drugs that will require prior additionzation of the formulary exclusions
2	respectively.
3	(d) A health benefit plan issuer may modify drug coverage provided under a health
4	benefit plan if:
5	(1) The modification occurs at the time of coverage renewal;
6	(2) The modification is effective uniformly among all group health benefit plan sponsors
7	covered by identical or substantially identical health benefit plans or all individuals covered by
8	identical or substantially identical individual health benefit plans, as applicable; and
9	(3) Not later than the sixtieth day before the date the modification is effective, the issuer
10	provides written notice of the modification to the commissioner, each affected group health
11	benefit plan sponsor, each affected enrollee in an affected group health benefit plan, and each
12	affected individual health benefit plan holder.
13	(e) Modifications affecting drug coverage that require written or electronic notice under
14	subsection (d) of this section, include:
15	(1) Removing a drug from a formulary;
16	(2) Adding a requirement that an enrollee receive prior authorization for a drug;
17	(3) Imposing or altering a quantity limit for a drug;
18	(4) Imposing a step-therapy restriction for a drug; and
19	(5) Moving a drug to a higher cost-sharing tier unless a generic drug alternative to the
20	drug is available.
21	(ii)(f) A nonprofit, hospital-service corporation may immediately remove from its plan
22	formularies covered prescription drugs deemed unsafe by the nonprofit, hospital-service
23	corporation or the Food and Drug Administration, or removed from the market by their
24	manufacturer, without meeting the requirements of this section.
25	SECTION 3. Section 27-20-37 of the General Laws in Chapter 27-20 entitled "Nonprofit
26	Medical Service Corporations" is hereby amended to read as follows:
27	27-20-37. Drug coverage.
28	(a) Any nonprofit, medical-service corporation that utilizes a formulary of medications
29	for which coverage is provided under an individual or group-plan, master contract shall require
30	any physician or other person authorized by the department of health to prescribe medication to
31	prescribe from the formulary. A physician or other person authorized by the department of health
32	to prescribe medication shall be allowed to prescribe medications previously on, or not on, the
33	nonprofit, medical-service corporation's formulary if he or she believes that the prescription of
34	the non-formulary medication is medically necessary. A nonprofit, medical-service corporation

2	formulary medication meets the nonprofit, medical-service corporation's medical-exception
3	criteria for the coverage of that medication.
4	(b) A nonprofit, medical-service corporation's medical-exception criteria for the coverage
5	of non-formulary medications shall be developed in accordance with § 23-17.13-3(e)(3) 27-18.8-
6	<u>3(b)(5)</u> .
7	(c) Any subscriber who is aggrieved by a denial of benefits to be provided under this
8	section may appeal the denial in accordance with the rules and regulations promulgated by the
9	department of health commissioner pursuant to chapter 17.12 of title 23 chapter 18.9 of title 27.
10	(d) Prior to removing a prescription drug from its plan's formulary or making any change
11	in the preferred or tiered, cost-sharing status of a covered prescription drug, a nonprofit, medical-
12	service corporation must provide at least thirty (30) days' notice to authorized prescribers by
13	established communication methods of policy and program updates and by updating available
14	references on web-based publications. All adversely affected members must be provided at least
15	thirty (30) days' notice prior to the date such change becomes effective by a direct notification:
16	(i) The written or electronic notice must contain the following information:
17	(A) The name of the affected prescription drug;
18	(B) Whether the plan is removing the prescription drug from the formulary, or changing
19	its preferred or tiered, cost-sharing status; and
20	(C) The means by which subscribers may obtain a coverage determination or medical
21	exception, in the case of drugs that will require prior authorization or are formulary exclusions
22	respectively.
23	(d) A health benefit plan issuer may modify drug coverage provided under a health
24	benefit plan if:
25	(1) The modification occurs at the time of coverage renewal;
26	(2) The modification is effective uniformly among all group health benefit plan sponsors
27	covered by identical or substantially identical health benefit plans or all individuals covered by
28	identical or substantially identical individual health benefit plans, as applicable; and
29	(3) Not later than the sixtieth day before the date the modification is effective, the issuer
30	provides written notice of the modification to the commissioner, each affected group health
31	benefit plan sponsor, each affected enrollee in an affected group health benefit plan, and each
32	affected individual health benefit plan holder.
33	(e) Modifications affecting drug coverage that require written or electronic notice under
34	subsection (d) of this section, include:

shall be required to provide coverage for a non-formulary medication only when the non-

1	(1) Removing a drug from a formulary;
2	(2) Adding a requirement that an enrollee receive prior authorization for a drug;
3	(3) Imposing or altering a quantity limit for a drug;
4	(4) Imposing a step-therapy restriction for a drug; and
5	(5) Moving a drug to a higher cost-sharing tier unless a generic drug alternative to the
6	drug is available.
7	(ii)(f) A nonprofit, medical-service corporation may immediately remove from its plan
8	formularies covered prescription drugs deemed unsafe by the nonprofit, medical-service
9	corporation or the Food and Drug Administration, or removed from the market by their
10	manufacturer, without meeting the requirements of this section.
11	SECTION 4. Section 27-20.1-15 of the General Laws in Chapter 27-20.1 entitled
12	"Nonprofit Dental Service Corporations" is hereby amended to read as follows:
13	27-20.1-15. Drug coverage.
14	(a) Any nonprofit, dental-service corporation that utilizes a formulary of medications for
15	which coverage is provided under an individual or group-plan, master contract shall require any
16	physician or other person authorized by the department of health to prescribe medication to
17	prescribe from the formulary. A physician or other person authorized by the department of health
18	to prescribe medication shall be allowed to prescribe medications previously on, or not on, the
19	nonprofit, dental-service corporation's formulary if he or she believes that the prescription of the
20	non-formulary medication is medically necessary. A nonprofit, dental-service corporation shall be
21	required to provide coverage for a non-formulary medication only when the non-formulary
22	medication meets the nonprofit, dental-service corporation's medical-exception criteria for the
23	coverage of that medication.
24	(b) A nonprofit, dental-service corporation's medical-exception criteria for the coverage
25	of non-formulary medications shall be developed in accordance with § 23-17.13-3(e)(3) 27-18.8-
26	<u>3(b)(5)</u> .
27	(c) Any subscriber who is aggrieved by a denial of benefits to be provided under this
28	section may appeal the denial in accordance with the rules and regulations promulgated by the
29	department of health commissioner pursuant to chapter 17.12 of title 23 chapter 18.9 of title 27.
30	(d) Prior to removing a prescription drug from its plan's formulary or making any change
31	in the preferred or tiered, cost sharing status of a covered prescription drug, a nonprofit, dental
32	service corporation must provide at least thirty (30) days' notice to authorized prescribers by
33	established communication methods of policy and program updates and by updating available
34	references on web based publications. All adversely affected members must be provided at least

1	thirty (30) days' notice prior to the date such change becomes effective by a direct notification:
2	(i) The written or electronic notice must contain the following information:
3	(A) The name of the affected prescription drug;
4	(B) Whether the plan is removing the prescription drug from the formulary, or changing
5	its preferred or tiered, cost-sharing status; and
6	(C) The means by which subscribers may obtain a coverage determination or medical
7	exception, in the case of drugs that will require prior authorization or are formulary exclusions
8	respectively.
9	(d) A health benefit plan issuer may modify drug coverage provided under a health
10	benefit plan if:
11	(1) The modification occurs at the time of coverage renewal;
12	(2) The modification is effective uniformly among all group health benefit plan sponsors
13	covered by identical or substantially identical health benefit plans or all individuals covered by
14	identical or substantially identical individual health benefit plans, as applicable; and
15	(3) Not later than the sixtieth day before the date the modification is effective, the issuer
16	provides written notice of the modification to the commissioner, each affected group health
17	benefit plan sponsor, each affected enrollee in an affected group health benefit plan, and each
18	affected individual health benefit plan holder.
19	(e) Modifications affecting drug coverage that require written or electronic notice under
20	subsection (d) of this section, include:
21	(1) Removing a drug from a formulary;
22	(2) Adding a requirement that an enrollee receive prior authorization for a drug;
23	(3) Imposing or altering a quantity limit for a drug;
24	(4) Imposing a step-therapy restriction for a drug; and
25	(5) Moving a drug to a higher cost-sharing tier unless a generic drug alternative to the
26	drug is available.
27	(ii)(f) A nonprofit, dental-service corporation may immediately remove from its plan
28	formularies covered prescription drugs deemed unsafe by the nonprofit, dental-service
29	corporation or the Food and Drug Administration, or removed from the market by their
30	manufacturer, without meeting the requirements of this section.
31	SECTION 5. Section 27-41-51 of the General Laws in Chapter 27-41 entitled "Health
32	Maintenance Organizations" is hereby amended to read as follows:
33	<u>27-41-51. Drug coverage.</u>
34	(a) Any health-maintenance organization that utilizes a formulary of medications for

1	which coverage is provided under an individual or group-plan, master contract shall require any
2	physician or other person authorized by the department of health to prescribe medication to
3	prescribe from the formulary. A physician or other person authorized by the department of health
4	to prescribe medication shall be allowed to prescribe medications previously on, or not on, the
5	health-maintenance organization's formulary if he or she believes that the prescription of non-
6	formulary medication is medically necessary. A health-maintenance organization shall be
7	required to provide coverage for a non-formulary medication only when the non-formulary
8	medication meets the health-maintenance organization's medical-exception criteria for the
9	coverage of that medication.
10	(b) A health-maintenance organization's medical-exception criteria for the coverage of
11	non-formulary medications shall be developed in accordance with § 23-17.13-3(e)(3) 27-18.8-
12	<u>3(b)(5)</u> .
13	(c) Any subscriber who is aggrieved by a denial of benefits to be provided under this
14	section may appeal the denial in accordance with the rules and regulations promulgated by the
15	department of health commissioner pursuant to chapter 17.12 of title 23 chapter 18.9 of title 27.
16	(d) Prior to removing a prescription drug from its plan's formulary or making any change
17	in the preferred or tiered, cost sharing status of a covered prescription drug, a health-maintenance
18	organization must provide at least thirty (30) days' notice to authorized prescribers by established
19	communication methods of policy and program updates and by updating available references on
20	web based publications. All adversely affected members must be provided at least thirty (30)
21	days' notice prior to the date such change becomes effective by a direct notification:
22	(i) The written or electronic notice must contain the following information:
23	(A) The name of the affected prescription drug;
24	(B) Whether the plan is removing the prescription drug from the formulary, or changing
25	its preferred or tiered, cost-sharing status; and
26	(C) The means by which subscribers may obtain a coverage determination or medical
27	exception, in the case of drugs that will require prior authorization or are formulary exclusions
28	respectively.
29	(d) A health benefit plan issuer may modify drug coverage provided under a health
30	benefit plan if:
31	(1) The modification occurs at the time of coverage renewal:
32	(2) The modification is effective uniformly among all group health benefit plan sponsors
33	covered by identical or substantially identical health benefit plans or all individuals covered by
34	identical or substantially identical individual health benefit plans, as applicable; and

1	(3) Not later than the sixtieth day before the date the modification is effective, the issuer
2	provides written notice of the modification to the commissioner, each affected group health
3	benefit plan sponsor, each affected enrollee in an affected group health benefit plan, and each
4	affected individual health benefit plan holder.
5	(e) Modifications affecting drug coverage that require written or electronic notice under
6	subsection (d) of this section, include:
7	(1) Removing a drug from a formulary;
8	(2) Adding a requirement that an enrollee receive prior authorization for a drug;
9	(3) Imposing or altering a quantity limit for a drug;
10	(4) Imposing a step-therapy restriction for a drug; and
11	(5) Moving a drug to a higher cost-sharing tier unless a generic drug alternative to the
12	drug is available.
13	(ii)(f) A health-maintenance organization may immediately remove from its plan
14	formularies covered prescription drugs deemed unsafe by the health-maintenance organization or
15	the Food and Drug Administration, or removed from the market by their manufacturer, without
16	meeting the requirements of this section.
17	SECTION 6. This act shall take effect upon passage.
	====== LC004122
	ECOVITEE .

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EXPLANATION

BY THE LEGISLATIVE COUNCIL

OF

AN ACT

RELATING TO INSURANCE

1	This act would allow an issuer of a health benefit plan to modify drug coverage pursuant
2	to a health benefit plan if: (1) the modification occurs are the time of coverage renewal; (2) the
3	modification is effective among all identical or substantially identical health benefit plans; and (3)
4	written notice is provided not later than sixty (60) days before the date the modification becomes
5	effective.
6	This act would take effect upon passage.
	====== LC004122

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