LC005138

2014 -- S 2803

STATE OF RHODE ISLAND

IN GENERAL ASSEMBLY

JANUARY SESSION, A.D. 2014

AN ACT

RELATING TO INSURANCE - ACCIDENT AND SICKNESS INSURANCE POLICIES

Introduced By: Senator Christopher S.Ottiano Date Introduced: March 25, 2014 Referred To: Senate Health & Human Services (by request)

It is enacted by the General Assembly as follows:

1	SECTION	1.	Chapter	27-18	of	the	General	Laws	entitled	"Accident	and	Sickness
2	Insurance Policies"	is ł	hereby an	nended	by a	addi	ng thereto	the fo	llowing s	ection:		

3	27-18-82. Cancer patient safety and environmental protection (a) Purpose. It is the
4	policy of the state of Rhode Island not to permit the introduction of pollutants into the
5	groundwaters and water systems of the state, or otherwise to be discharged in concentrations
6	which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same are defined in
7	the Rhode Island department of environmental managements groundwater quality rules and the
8	rules and regulations for hazardous waste management.
9	(b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients
10	undergoing chemotherapy treatment may contain levels of chemicals that are toxic,
11	carcinogenic, mutagenic or teratogenic for a certain period of time, to such an extent that the
12	World Health Organization defines genotoxic waste as chemotherapy drug waste including urine,
13	feces and vomit from patients, which may contain potentially hazardous amounts of the
14	administered cytostatic drugs or of their metabolites, and which should be considered genotoxic
15	for at least forty-eight (48) hours and sometimes up to one week after drug administration.
16	(2) The World Health Organization further states that any discharge of genotoxic waste
17	into the environment could have disastrous ecological consequences. The World Health
18	Organization core principles require that all personnel associated with financing and supporting
19	healthcare activities should provide for the costs of managing healthcare waste. This is the duty of

care. The World Health Organization places the responsibility for genotoxic waste on the chief 2 pharmacist and further states that the chief pharmacist also has the special responsibility of 3 ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely. 4 (3) The federal Occupational Safety and Health Administration ("OSHA") is the main 5 federal agency charged with the enforcement of safety and health legislation. OSHA, in concert with the National Institute for Occupational Safety and Health ("NIOSH") and the joint 6 7 commission on healthcare, an independent, not-for-profit organization that accredits and certifies 8 more than twenty thousand (20,000) healthcare organizations and programs in the United States, 9 stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been 10 known to cause cancer, reproductive and developmental problems, allergic reactions, and other 11 adverse effects that can be irreversible even after low-level exposures"; and 12 (4) The American Cancer Society has published a comprehensive list of safety 13 precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy 14 and their families. Therefore, for the protection of both the public health and the environment, the 15 general assembly shall require that standards as set forth pursuant to this section be observed to 16 address this serious safety issue. 17 (c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other 18 health care professionals licensed in the state of Rhode Island authorized to prescribe and/or 19 administer chemotherapy treatment shall: 20 (1) Provide written notice from the prescribing pharmacist to each patient undergoing 21 such treatment as to the hazards posed to patients and their families of extremely hazardous 22 excretions, including, but not limited to, urine, vomit, and feces for a period following treatment 23 as generally determined by the food and drug administration label accompanying said 24 chemotherapy drug or drugs; 25 (2) Provide a sufficient collection method so that providers and patients can safely collect and contain extremely hazardous excretions for a period of time as determined by the United 26 27 States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription 28 insert(s); and 29 (3) Provide for safe and proper disposal of said collected extremely hazardous excretions. 30 (d) All expenses incurred as a result of this section shall be paid by Medicare, Medicaid 31 or any private insurance company providing health care insurance and licensed pursuant to this 32 chapter. 33 (e) Receipt of notice from the party administering chemotherapy drugs or their agent is 34 responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief

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1 pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder.

2 (f) For the purposes of this section, "extremely hazardous excretions" shall mean any

3 excretion from a patient on a regimen of chemotherapy agents that are antineoplastic or cytotoxic,

4 during the period of administration and the time period referenced in subsection (c) of this

5 section, including, but not limited to, drugs listed in the NIOSH list of antineoplastic and other

- 6 <u>hazardous drugs, as the same may be updated or amended from time to time.</u>
 - SECTION 2. Chapter 27-18.5 of the General Laws entitled "Individual Health Insurance

8 Coverage" is hereby amended by adding thereto the following section:

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27-18.5-11. Cancer patient safety and environmental protection. --

(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
 pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in
 concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same
 are defined in the Rhode Island department of environmental management groundwater quality
 rules and the rules and regulations for hazardous waste management.
 (b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients

undergoing chemotherapy treatment may contain levels of chemicals that are toxic, carcinogenic, mutagenic or teratogenic for a certain period of time, to such an extent that the World Health Organization defines genotoxic waste as chemotherapy drug waste including urine, feces and vomit from patients, which may contain potentially hazardous amounts of the administered cytostatic drugs or of their metabolites, and which should be considered genotoxic for at least forty-eight (48) hours and sometimes up to one week after drug administration.

(2) The World Health Organization further states that any discharge of genotoxic waste into the environment could have disastrous ecological consequences. The World Health Organization core principles require that all personnel associated with financing and supporting healthcare activities should provide for the costs of managing healthcare waste. This is the duty of care. The World Health Organization places the responsibility for genotoxic waste on the chief pharmacist and further states that the chief pharmacist also has the special responsibility of ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely.

(3) The federal Occupational Safety and Health Administration ("OSHA") is the main federal agency charged with the enforcement of safety and health legislation. OSHA, in concert with the National Institute for Occupational Safety and Health ("NIOSH") and the joint commission on healthcare, an independent, not-for-profit organization that accredits and certifies more than twenty thousand (20,000) healthcare organizations and programs in the United States, stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been

1 known to cause cancer, reproductive and developmental problems, allergic reactions, and other

2 adverse effects that can be irreversible even after low-level exposures"; and

3 (4) The American Cancer Society has published a comprehensive list of safety 4 precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy 5 and their families. Therefore, for the protection of both the public health and the environment, the general assembly shall require that standards are set forth pursuant to this section to address this 6 7 serious safety issue. 8 (c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other 9 healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or 10 administer chemotherapy treatment shall: 11 (1) Provide written notice from the prescribing pharmacist to each patient undergoing 12 such treatment as to the hazards posed to patients and their families of extremely hazardous 13 excretions, including, but not limited to, urine, vomit, and feces, for a period following treatment 14 as generally determined by the food and drug administration label accompanying said 15 chemotherapy drug or drugs; 16 (2) Provide a sufficient collection method so that providers and patients can safely collect 17 and contain extremely hazardous excretions for a period of time as determined by the United States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription 18 19 insert(s); and 20 (3) Provide for safe and proper disposal of said collected extremely hazardous excretions. 21 (d) All expenses incurred as a result of this section shall be paid by Medicare, Medicaid 22 or any private insurance company providing healthcare insurance and licensed pursuant to this 23 chapter. 24 (e) Receipt of notice from the party administering chemotherapy drugs or their agent responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief 25 26 pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder. 27 (f) For the purposes of this section, "extremely hazardous excretions" shall mean any 28 excretion from a patient on a regimen of chemotherapy agents that are antineoplastic or cytotoxic, 29 during the period of administration and the time period referenced in subsection (c) of this 30 section, including, but not limited to, drugs listed in the NIOSH list of antineoplastic and other 31 hazardous drugs, as the same may be updated or amended from time to time.

32 SECTION 3. Chapter 27-19 of the General Laws entitled "Nonprofit Hospital Service
 33 Corporations" is hereby amended by adding thereto the following section:

34 27-19-73. Cancer patient safety and environmental protection. --

(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
 pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in
 concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same
 are defined in the Rhode Island department of environmental management groundwater quality
 rules and the rules and regulations for hazardous waste management.

- 6 (b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients 7 undergoing chemotherapy treatment may contain levels of chemicals that are toxic, carcinogenic, 8 mutagenic or teratogenic for a certain period of time, to such an extent that the World Health 9 Organization defines genotoxic waste as chemotherapy drug waste including urine, feces and 10 vomit from patients, which may contain potentially hazardous amounts of the administered 11 cytostatic drugs or of their metabolites, and which should be considered genotoxic for at least 12 forty-eight (48) hours and sometimes up to one week after drug administration.
- 13 (2) The World Health Organization further states that any discharge of genotoxic waste 14 into the environment could have disastrous ecological consequences. The World Health 15 Organization core principles require that all personnel associated with financing and supporting 16 healthcare activities should provide for the costs of managing healthcare waste. This is the duty of 17 care. The world health organization places the responsibility for genotoxic waste on the chief 18 pharmacist and further states that the chief pharmacist also has the special responsibility of 19 ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely.
 - (3) The federal Occupational Safety and Health Administration ("OSHA") is the main

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federal agency charged with the enforcement of safety and health legislation. OSHA, in concert with the National Institute for Occupational Safety and Health ("NIOSH") and the joint commission on healthcare, an independent, not-for-profit organization that accredits and certifies more than twenty thousand (20,000) healthcare organizations and programs in the United States, stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been known to cause cancer, reproductive and developmental problems, allergic reactions, and other

27 <u>adverse effects that can be irreversible even after low-level exposures"; and</u>

(4) The American Cancer Society has published a comprehensive list of safety
 precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy
 and their families. Therefore, for the protection of both the public health and the environment, the
 general assembly shall require that standards as set forth pursuant to this section be observed to
 address this serious safety issue.

33 (c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other
 34 healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or

1 <u>administer chemotherapy treatment shall:</u>

2	(1) Provide written notice from the prescribing pharmacist to each patient undergoing
3	such treatment as to the hazards posed to patients and their families of extremely hazardous
4	excretions, including, but not limited to, urine, vomit, and feces, for a period following treatment
5	as generally determined by the food and drug administration label accompanying said
6	chemotherapy drug or drugs;
7	(2) Provide a sufficient collection method so that providers and patients can safely collect
8	and contain extremely hazardous excretions for a period of time as determined by the United
9	States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription
10	insert(s); and
11	(3) Provide for safe and proper disposal of said collected extremely hazardous excretions.
12	(d) All expenses incurred as a result of this section shall be paid by Medicare, Medicaid
13	or any private insurance company providing healthcare insurance and licensed pursuant to this
14	chapter.
15	(e) Receipt of notice from the party administering chemotherapy drugs or their agent
16	responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief
17	pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder.
18	(f) For the purposes of this section, "extremely hazardous excretions" shall mean any
19	excretion from a patient on a regimen of chemotherapy agents that are antineoplastic or cytotoxic,
20	during the period of administration and the time period referenced in subsection (c) of this
21	section, including, but not limited to, drugs listed in the NIOSH list of antineoplastic and other
22	hazardous drugs, as the same may be updated or amended from time to time.
23	SECTION 4. Chapter 27-20 of the General Laws entitled "Nonprofit Medical Service
24	Corporations" is hereby amended by adding thereto the following section:
25	27-20-69. Cancer patient safety and environmental protection
26	(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
27	pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in
28	concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same
29	are defined in the Rhode Island department of environmental management groundwater quality
30	rules and the rules and regulations for hazardous waste management.
31	(b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients
32	undergoing chemotherapy treatment may contain levels of chemicals that are toxic, carcinogenic,
33	mutagenic or teratogenic for a certain period of time, to such an extent that the World Health
34	Organization defines genotoxic waste as chemotherapy drug waste including urine, feces and

1 vomit from patients, which may contain potentially hazardous amounts of the administered

2 cytostatic drugs or of their metabolites, and which should be considered genotoxic for at least

3 forty-eight (48) hours and sometimes up to one week after drug administration.

4 (2) The World Health Organization further states that any discharge of genotoxic waste 5 into the environment could have disastrous ecological consequences. The World Health Organization core principles require that all personnel associated with financing and supporting 6 7 healthcare activities should provide for the costs of managing healthcare waste. This is the duty of 8 care. The world health organization places the responsibility for genotoxic waste on the chief 9 pharmacist and further states that the chief pharmacist also has the special responsibility of 10 ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely. 11 (3) The federal Occupational Safety and Health Administration ("OSHA") is the main 12 federal agency charged with the enforcement of safety and health legislation. OSHA, in concert 13 with the National Institute for Occupational Safety and Health ("NIOSH") and the joint 14 commission on healthcare, an independent, not-for-profit organization that accredits and certifies 15 more than twenty thousand (20,000) healthcare organizations and programs in the United States,

16 stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been

17 known to cause cancer, reproductive and developmental problems, allergic reactions, and other

18 adverse effects that can be irreversible even after low-level exposures"; and

(4) The American Cancer Society has published a comprehensive list of safety
 precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy
 and their families. Therefore, for the protection of both the public health and the environment, the
 general assembly shall require that standards as set forth pursuant to this section be observed to

23 address this serious safety issue.

(c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other
 healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or
 administer chemotherapy treatment shall:

27 (1) Provide written notice from the prescribing pharmacist to each patient undergoing
28 such treatment as to the hazards posed to patients and their families of extremely hazardous
29 excretions, including, but not limited to, urine, vomit, and feces, for a period following treatment
30 as generally determined by the food and drug administration label accompanying said

31 <u>chemotherapy drug or drugs;</u>

32 (2) Provide a sufficient collection method so that providers and patients can safely collect
 33 and contain extremely hazardous excretions for a period of time as determined by the United
 34 States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription

1 <u>insert(s); and</u>

2 (3) Provide for safe and proper disposal of said collected extremely hazardous excretions. (d) All expenses incurred as a result of this section shall be paid by Medicare, Medicaid 3 4 or any private insurance company providing healthcare insurance and licensed pursuant to this 5 chapter. (e) Receipt of notice from the party administering chemotherapy drugs or their agent 6 7 responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief 8 pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder. 9 (f) For the purposes of this section, "extremely hazardous excretions" shall mean any 10 excretion from a patient on a regimen of chemotherapy agents that are antineoplastic or cytotoxic, 11 during the period of administration and the time period referenced in subsection (c) of this 12 section, including, but not limited to, drugs listed in the NIOSH list of antineoplastic and other 13 hazardous drugs, as the same may be updated or amended from time to time. 14 SECTION 5. Chapter 27-41 of the General Laws entitled "Health Maintenance 15 Organizations" is hereby amended by adding thereto the following section: 16 27-41-86. Cancer patient safety and environmental protection.--(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of 17 18 pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in 19 concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same 20 are defined in the Rhode Island department of environmental management groundwater quality 21 rules and the rules and regulations for hazardous waste management. 22 (b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients 23 undergoing chemotherapy treatment may contain levels of chemicals that are toxic, carcinogenic, 24 mutagenic or teratogenic for a certain period of time, to such an extent that the World Health 25 Organization defines genotoxic waste as chemotherapy drug waste including urine, feces and 26 vomit from patients, which may contain potentially hazardous amounts of the administered 27 cytostatic drugs or of their metabolites, and which should be considered genotoxic for at least 28 forty-eight (48) hours and sometimes up to one week after drug administration. 29 (2) The World Health Organization further states that any discharge of genotoxic waste 30 into the environment could have disastrous ecological consequences. The World Health 31 Organization core principles require that all personnel associated with financing and supporting 32 healthcare activities should provide for the costs of managing healthcare waste. This is the duty of 33 care. The world health organization places the responsibility for genotoxic waste on the chief 34 pharmacist and further states that the chief pharmacist also has the special responsibility of

1 <u>ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely.</u>

2 (3) The federal Occupational Safety and Health Administration ("OSHA") is the main federal agency charged with the enforcement of safety and health legislation. OSHA, in concert 3 4 with the National Institute for Occupational Safety and Health ("NIOSH") and the joint 5 commission on healthcare, an independent, not-for-profit organization that accredits and certifies more than twenty thousand (20,000) healthcare organizations and programs in the United States, 6 7 stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been 8 known to cause cancer, reproductive and developmental problems, allergic reactions, and other 9 adverse effects that can be irreversible even after low-level exposures"; and 10 (4) The American Cancer Society has published a comprehensive list of safety 11 precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy 12 and their families. Therefore, for the protection of both the public health and the environment, the 13 general assembly shall require that standards as set forth pursuant to this section be observed to 14 address this serious safety issue. 15 (c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other 16 healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or 17 administer chemotherapy treatment shall: (1) Provide written notice from the prescribing pharmacist to each patient undergoing 18 19 such treatment as to the hazards posed to patients and their families of extremely hazardous 20 excretions, including, but not limited to, urine, vomit, and feces, for a period following treatment 21 as generally determined by the food and drug administration label accompanying said chemotherapy drug or drugs; 22 (2) Provide a sufficient collection method so that providers and patients can safely collect 23 24 and contain extremely hazardous excretions for a period of time as determined by the United States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription 25 26 insert(s); and 27 (3) Provide for safe and proper disposal of said collected extremely hazardous excretions. 28 (d) All expenses incurred as a result of this section shall be paid by Medicare. Medicaid 29 or any private insurance company providing healthcare insurance and licensed pursuant to this 30 chapter. 31 (e) Receipt of notice from the party administering chemotherapy drugs or their agent 32 responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder. 33

(f) For the purposes of this section, "extremely hazardous excretions" shall mean any

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- 1 excretion from a patient on a regimen of chemotherapy agents that are antineoplastic or cytotoxic,
- 2 during the period of administration and the time period referenced in subsection (c) of this
- 3 section, including, but not limited to, drugs listed in the NIOSH list of Antineoplastic and other
- 4 <u>hazardous drugs, as the same may be updated or amended from time to time.</u>
- 5 SECTION 6. This act shall take effect upon passage.

LC005138

EXPLANATION

BY THE LEGISLATIVE COUNCIL

OF

AN ACT

RELATING TO INSURANCE - ACCIDENT AND SICKNESS INSURANCE POLICIES

This act would require that protections related to the disposal of extremely hazardous
 wastes generated by the use of toxic, carcinogenic, mutagenic, or teratogenic chemotherapy drugs
 be implemented by pharmacists, physicians, healthcare providers, and insurers in the state of
 Rhode Island.
 This act would take effect upon passage.

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