2014 -- H 7644

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STATE OF RHODE ISLAND

IN GENERAL ASSEMBLY

JANUARY SESSION, A.D. 2014

AN ACT

RELATING TO INSURANCE - ACCIDENT AND SICKNESS INSURANCE POLICIES

Introduced By: Representatives McNamara, Valencia, Bennett, and Shekarchi

Date Introduced: February 27, 2014

Referred To: House Corporations

It is enacted by the General Assembly as follows:

SECTION 1. Chapter 27-18 of the General Laws entitled "Accident and Sickness 1 2 Insurance Policies" is hereby amended by adding thereto the following section: 3 27-18-82. 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 4 5 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 6 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. (2) The World Health Organization further states that any discharge of genotoxic waste 16 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

healthcare activities should provide for the costs of managing healthcare waste. This is the duty of

1	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
6	with the National Institute for Occupational Safety and Health ("NIOSH") and the joint
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
26	and contain extremely hazardous excretions for a period of time as determined by the United
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	<u>chapter.</u>
33	(e) Receipt of notice from the party administering chemotherapy drugs or their agent is

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	hazardous drugs, as the same may be updated or amended from time to time.
7	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:
9	27-18.5-11. Cancer patient safety and environmental protection
10	(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
11	pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in
12	concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same
13	are defined in the Rhode Island department of environmental management groundwater quality
14	rules and the rules and regulations for hazardous waste management.
15	(b) Findings. (1) It is acknowledged by medical experts that bodily wastes of patients
16	undergoing chemotherapy treatment may contain levels of chemicals that are toxic, carcinogenic,
17	mutagenic or teratogenic for a certain period of time, to such an extent that the World Health
18	Organization defines genotoxic waste as chemotherapy drug waste including urine, feces and
19	vomit from patients, which may contain potentially hazardous amounts of the administered
20	cytostatic drugs or of their metabolites, and which should be considered genotoxic for at least
21	forty-eight (48) hours and sometimes up to one week after drug administration.
22	(2) The World Health Organization further states that any discharge of genotoxic waste
23	into the environment could have disastrous ecological consequences. The World Health
24	Organization core principles require that all personnel associated with financing and supporting
25	healthcare activities should provide for the costs of managing healthcare waste. This is the duty of
26	care. The World Health Organization places the responsibility for genotoxic waste on the chief
27	pharmacist and further states that the chief pharmacist also has the special responsibility of
28	ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely.
29	(3) The federal Occupational Safety and Health Administration ("OSHA") is the main
30	federal agency charged with the enforcement of safety and health legislation. OSHA, in concert
31	with the National Institute for Occupational Safety and Health ("NIOSH") and the joint
32	commission on healthcare, an independent, not-for-profit organization that accredits and certifies
33	more than twenty thousand (20,000) healthcare organizations and programs in the United States,
34	stated in a 2011 letter to every hospital in the country that "Islome 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
6	general assembly shall require that standards are set forth pursuant to this section to address this
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
18	States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription
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
25	responsible for proper disposal of the hazardous wastes by the prescribing pharmacist or chief
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

1	(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
2	pollutants into the groundwaters and water systems of the state, or otherwise to be discharged in
3	concentrations which are known to be toxic, carcinogenic, mutagenic, or teratogenic as the same
4	are defined in the Rhode Island department of environmental management groundwater quality
5	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.
20	(3) The federal Occupational Safety and Health Administration ("OSHA") is the main
21	federal agency charged with the enforcement of safety and health legislation. OSHA, in concert
22	with the National Institute for Occupational Safety and Health ("NIOSH") and the joint
23	commission on healthcare, an independent, not-for-profit organization that accredits and certifies
24	more than twenty thousand (20,000) healthcare organizations and programs in the United States,
25	stated in a 2011 letter to every hospital in the country that "[s]ome of these drugs have been
26	known to cause cancer, reproductive and developmental problems, allergic reactions, and other
27	adverse effects that can be irreversible even after low-level exposures"; and
28	(4) The American Cancer Society has published a comprehensive list of safety
29	precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy
30	and their families. Therefore, for the protection of both the public health and the environment, the
31	general assembly shall require that standards as set forth pursuant to this section be observed to
32	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	administer chemotherapy treatment shall:
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	<u>chapter.</u>
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
6	Organization core principles require that all personnel associated with financing and supporting
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
19	(4) The American Cancer Society has published a comprehensive list of safety
20	precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy
21	and their families. Therefore, for the protection of both the public health and the environment, the
22	general assembly shall require that standards as set forth pursuant to this section be observed to
23	address this serious safety issue.
24	(c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other
25	healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or
26	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	chemotherapy drug or drugs;
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	insert(s); and
2	(3) Provide for safe and proper disposal of said collected extremely hazardous excretions.
3	(d) All expenses incurred as a result of this section shall be paid by Medicare, Medicaid
4	or any private insurance company providing healthcare insurance and licensed pursuant to this
5	chapter.
6	(e) Receipt of notice from the party administering chemotherapy drugs or their agent
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
17	(a) Purpose. It is the policy of the state of Rhode Island not to permit the introduction of
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	ensuring that genotoxic products are used safely, and that genotoxic waste is managed safely.
2	(3) The federal Occupational Safety and Health Administration ("OSHA") is the main
3	federal agency charged with the enforcement of safety and health legislation. OSHA, in concert
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
6	more than twenty thousand (20,000) healthcare organizations and programs in the United States,
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
0	(4) The American Cancer Society has published a comprehensive list of safety
1	precautions regarding the in-home personal hygiene for individuals undergoing chemotherapy
2	and their families. Therefore, for the protection of both the public health and the environment, the
.3	general assembly shall require that standards as set forth pursuant to this section be observed to
4	address this serious safety issue.
.5	(c) Chemotherapy precautions following treatment. All physicians, pharmacists, or other
6	healthcare professionals licensed in the state of Rhode Island authorized to prescribe and/or
7	administer chemotherapy treatment shall:
8	(1) Provide written notice from the prescribing pharmacist to each patient undergoing
9	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
22	chemotherapy drug or drugs;
23	(2) Provide a sufficient collection method so that providers and patients can safely collect
24	and contain extremely hazardous excretions for a period of time as determined by the United
25	States Food and Drug Administration ("FDA") and referenced on the relevant FDA prescription
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
80	<u>chapter.</u>
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
33	pharmacist shall satisfy the responsibility of the prescribing pharmacist hereunder.
34	(f) For the purposes of this section, "extremely hazardous excretions" shall mean any

- 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 hazardous drugs, as the same may be updated or amended from time to time.
- 5 SECTION 6. This act shall take effect upon passage.

LC004708

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.