GENERAL ASSEMBLY OF NORTH CAROLINA SESSION 2025

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Toxic-Free Medical Devices Act of 2025.

Short Title:

HOUSE BILL DRH40354-NM-33A

	Sponsors:	Rej	presentative Reeder.
	Referred to:		
1			A BILL TO BE ENTITLED
2 3			OHIBIT THE MANUFACTURING, SELLING, AND DISTRIBUTING OF OUS SOLUTION CONTAINERS AND INTRAVENOUS TUBING THAT
4	ARE INTENTIONALLY MADE WITH DEHP.		
5	The General Assembly of North Carolina enacts:		
6	SECTION 1. Chapter 130A of the General Statutes is amended by adding a new		
7	Article to rea		
8			" <u>Article 19C.</u>
9			"DEHP Hazard Management.
10	" <u>§ 130A-453</u>	3.33.	Legislative finding.
11	The Gen	eral A	assembly finds all of the following:
12	<u>(1</u>	<u>1)</u>	DEHP and other ortho-phthalates are toxic chemicals used primarily to
13			produce flexibility in plastics, mainly polyvinyl chloride (PVC).
14	<u>(2</u>	<u>2)</u>	DEHP is the most common plasticizer used in medical devices, including
15			intravenous solution containers, which are also known as IV bags, and
16			intravenous tubing.
17	<u>(3</u>	<u>3)</u>	Over the course of its shelf life, DEHP leaches from IV bags and tubing made
18			from DEHP into the solutions being held in the medical devices.
19	<u>(</u> 4	<u>4)</u>	DEHP is classified by the United States Environmental Protection Agency as
20			an endocrine-disrupting compound since it can:
21			a. Interfere with the hormonal system in humans and animals.
22			b. <u>Mimic or block the actions of hormones, leading to adverse effects on</u>
23	,	-\	reproductive health, development, and metabolism.
24	<u>(:</u>	<u>5)</u>	DEHP exposure has been associated with adverse effects on reproductive
25			organs and fertility. DEHP can also disrupt normal reproductive development,
26	(reduce sperm quality, and affect hormone levels in both males and females.
27	<u>((</u>	<u>6)</u>	DEHP is metabolized in the liver and can accumulate in the body over time.
28			Prolonged exposure to high levels of DEHP has been shown to cause liver and
29	(**	7)	kidney damage in animal studies. Inhelation or investion of DEUP con course respiratory imitation and allowed.
30 31	<u>C</u>	<u>7)</u>	Inhalation or ingestion of DEHP can cause respiratory irritation and allergic
32			reactions in some individuals, particularly those with preexisting respiratory
33	(6	9)	conditions or sensitivities. Studies have suggested a potential link between DEHP exposure and contain
33	(3	<u>8)</u>	Studies have suggested a potential link between DEHP exposure and certain types of cancer, including breast, liver, lung, and testicular cancer.
35	((9)	The United States Environmental Protection Agency has determined that
36	\overline{c}	<u> </u>	DEHP is a probable human carcinogen.
30			DETIT IS a probable numan caremogen.



The leaching of DEHP from medical devices at varying concentrations has 1 (10)2 been linked to multidrug resistance in breast cancer cells, inhibiting the effectiveness of breast cancer drugs. This phenomenon has been observed at 3 4 both high and low concentrations of DEHP, highlighting the potential impact 5 of DEHP leaching on cancer treatment outcomes. 6 Exposure to DEHP has been linked to multidrug resistance in triple-negative <u>(11)</u> 7 breast cancer cells, inhibiting the apoptosis mechanism induced by breast 8 cancer drugs, such as tamoxifen, and increasing cell proliferation. 9 DEHP has been suggested to serve as a mitogenic factor for estrogen (12)10 receptor-positive breast cancer cells, potentially making them multidrug 11 resistant. "§ 130A-453.34. Definitions. 12 13 The following definitions apply in this Article: 14 (1) DEHP. – Di(2-ethylhexyl) phthalate. 15 (2) Health care practitioner. – An individual who is authorized to practice some component of the healing arts by a license, permit, certificate, or registration 16 issued by a State licensing agency or board. 17 Intentionally added DEHP. – Any of the following: 18 (3) 19 DEHP that a manufacturer has intentionally added to a product and <u>a.</u> 20 that has a functional or technical effect on the product. 21 DEHP that is an intentional breakdown product of an added chemical. 22 <u>(4)</u> Intravenous solution container. – A container used to house medicine, fluid, 23 or nutrition therapy that is intravenously delivered to a patient in a hospital, 24 outpatient facility, or other health care facility. 25 Intravenous tubing. - Tubing used to intravenously administer fluids, (5) 26 medication, or nutrients directly to an adult, child, or infant. 27 Ortho-phthalate. – A class of chemicals that are esters of ortho-phthalic acid, (6) 28 including DEHP or any of the following: 29 Benzyl butyl phthalate (BBP). <u>a.</u> 30 b. Dibutyl phthalate (DBP). 31 Dicyclohexyl phthalate (DCHP). <u>c.</u> 32 <u>d.</u> Diethyl phthalate (DEP). 33 Diisobutyl phthalate (DIBP). <u>e.</u> 34 <u>f.</u> Diisodecyl phthalate (DIDP). 35 Diisononyl phthalate (DINP). g. 36 Di-n-hexyl phthalate (DnHP). h. 37 <u>i.</u> Di-n-octyl phthalate (DNOP). 38 Di-n-pentyl phthalate (DnPP). į. 39 Diisoheptyl phthalate (DIHP). k. 40 "§ 130A-453.35. Prohibitions.

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- Intravenous Solution Containers. Beginning January 1, 2030, a person or entity shall not manufacture, sell, or distribute into commerce in the State of North Carolina intravenous solution containers made with intentionally added DEHP.
- Intravenous Tubing. Beginning January 1, 2035, a person or entity shall not manufacture, sell, or distribute into commerce in the State of North Carolina intravenous tubing made with intentionally added DEHP.
- Replacement. A person may not replace DEHP, pursuant to this Article, with (c) another ortho-phthalate in a new or revised medical device.
- Maximum Quantity. An intravenous solution container or intravenous tubing product shall not have unintentionally added DEHP present at a quantity at or above 0.1 percent weight per weight (w/w).

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- (e) Exemptions. The following items, as described in Title 21 of the Code of Federal Regulations, are exempt from these provisions:
 - (1) Human blood collection and storage bags.
 - (2) Apheresis and cell therapy blood kits and bags, including integral tubing." **SECTION 2.** G.S. 130A-22(b3) reads as rewritten:

"(b3) The Secretary may impose an administrative penalty on a person who violates Article 19A or 19B Article 19A, 19B, or 19C of this Chapter or any rules adopted pursuant to Article 19A or 19B Article 19A, 19B, or 19C of this Chapter. Each day of a continuing violation is a separate violation. The penalty shall not exceed five thousand dollars (\$5,000) for each day the violation continues for Article 19A of this Chapter. The penalty shall not exceed five thousand dollars (\$5,000) for each day the violation continues for Article 19B of this Chapter. The penalty shall not exceed five thousand dollars (\$5,000) for each day the violation continues for Article 19C of this Chapter. The penalty authorized by this section does not apply to a person who is not required to be certified under Article 19A or 19B."

SECTION 3. Except as otherwise provided, this act is effective when it becomes law.

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