ROUGH DRAFT

Sixty-first Legislative Assembly of North Dakota

HOUSE BILL NO.

Introduced by

Representative Delmore

1 A BILL for an Act to create and enact chapter 19-09.1 of the North Dakota Century Code,

2 relating to safe cosmetics.

3 **BE IT ENACTED BY THE LEGISLATIVE ASSEMBLY OF NORTH DAKOTA:**

4 SECTION 1. Chapter 19-09.1 of the North Dakota Century Code is created and

5 enacted as follows:

6 <u>19-09.1-01. Definitions.</u> As used in this chapter, unless context or subject matter
7 <u>otherwise requires:</u>

means any agency or formally organized program or group
lepartment as being authoritative for the purpose of identifying
se cancer or reproductive toxicity.
as causing cancer or reproductive toxicity" means a chemical
noritative body as any of the following:
sted as known or reasonably anticipated to be a human
national toxicology report on carcinogens;
iven an overall carcinogenicity evaluation of group 1, group 2A,
y the international agency for research on cancer;
lentified as a group A, group B1, or group B2 carcinogen, or as
ely carcinogen by the United Stated environmental protection
lentified as having some or clear evidence of adverse
l, male reproductive, or female reproductive toxicity effects in a
xpert panel of the national toxicology program's center for the
isks to human reproduction.

Sixty-first

Legislative Assembly

1	<u>3.</u>	"Cosmetics" has the same meaning as that term is defined in United States Code,				
2		title 21, chapter 9, subchapter II, section 321, and includes:				
3		a. Articles intended to be rubbed, poured, sprinkled, or sprayed on, introduced				
4		into, or otherwise applied to the human body or any part thereof for cleansing,				
5		beautifying, promoting attractiveness, or altering the appearance; and				
6		b. Articles intended for use as a component of any such articles, excluding soap.				
7	<u>4.</u>	"Department" means the department of health.				
8	<u>5.</u>	"Ingredient" has the same meaning as that term is defined in Code of Federal				
9		Regulations, title 21, chapter 1, part 700, section 700.3, subdivision e, and does				
10		not include any incidental ingredient as defined in Code of Federal Regulations,				
11		title 21, chapter 1, part 701, section 701.3, subdivision 1.				
12	<u>6.</u>	"Manufacturer" means any person whose name appears on the label of a cosmetic				
13		product pursuant to the requirements of Code of Federal Regulations, title 21,				
14		section 701.12.				
15 <u>19-09.1-02. Safe cosmetics program.</u>						
16	<u>1.</u>	Beginning January 1, 2010, the manufacturer of any cosmetic product subject to				
17		regulation by the federal food and drug administration that is sold in this state shall,				
18		on a schedule and in electronic or other format, as determined by the department,				
19		provide the department with a complete and accurate list of its cosmetic products				
20		that, as of the date of submission, are sold in the state and that contain any				
21		ingredient that is a chemical identified as causing cancer or reproductive toxicity,				
22		including any chemical that meets either of the following conditions:				
23		a. A chemical contained in the product for purposes of fragrance or flavoring; or				
24		b. A chemical identified by the phrase "and other ingredients" and determined to				
25		be a trade secret pursuant to the procedure established in Code of Federal				
26		REgulations, title 21, part 702, section 720.8, and part 20. Any ingredient				
27		identified under this subsection must be considered confidential trade secret				
28		information and is not public.				
29	<u>2.</u>	Any information submitted pursuant to subsection 1 must identify each chemical				
30		both by name and chemical abstract service number and must specify the product				
31		or products in which the chemical is contained.				

Sixty-first Legislative Assembly

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1	<u>3.</u>	If an ingredient identified under this section subsequently is removed from the				
2		product in which it was contained or is no longer a chemical identified as causing				
3		cancer or reproductive toxicity by an authoritative body, the manufacturer of the				
4		product containing the ingredient shall submit the new information to the				
5		department. Upon receipt of new information, the department after verifying the				
6		accuracy of that information, shall revise the manufacturer's information on record				
7		with the department to reflect the new information. The manufacturer is not under				
8		obligation to submit subsequent information on the presence of the ingredient in				
9		the product unless subsequent changes require submittal of the information.				
10	<u>4.</u>	This section shall apply to cosmetic products that may also be regulated as a drug				
11		by the federal food and drug administration.				
12 <u>19-09.1-03. Investigations.</u>						
13	<u>1.</u>	In order to determine potential health effects of exposure to ingredients in				
14		cosmetics sold in the state, the department may conduct an investigation of one or				
15		more cosmetic products that contain chemicals identified as causing cancer or				
16		reproductive toxicity or other ingredients of concern to the department.				
17	<u>2.</u>	An investigation conducted pursuant to subsection 1 may include a review of				
18		available health effects data and studies, worksite health hazard evaluations,				
19		epidemiological studies to determine the health effects of exposures to chemicals				
20		in various subpopulations, and exposure assessments to determine total				
21		exposures to individuals in various settings.				
22	<u>3.</u>	If an investigation is conducted under subsection 1, the manufacturer of any				
23		product subject to the investigation may submit relevant health effects data and				
24		studies to the department.				
25	<u>4.</u>	In order to further the purposes of an investigation, the department may require				
26		manufacturers of products subject to the investigation to submit to the department				
27		relevant health effects data and studies available to the manufacturer and other				
28		available information as requested by the department, including the concentration				
29		of the chemical in the product, the amount by volume or weight of the product that				
30		comprises the average daily application or use, and sales and use date necessary				
31		to determine where the product is used in the occupational setting.				

Sixty-first Legislative Assembly

1	<u>5.</u>	The department shall establish reasonable deadlines for the submittal of
2		information required pursuant to subsection 4. Failure by a manufacturer to submit
3		the information in compliance with the requirements of the department is a violation
4		of this section.
5	<u>6.</u>	If the department determines pursuant to an investigation that an ingredient in a
6		cosmetic product is potentially toxic at the concentrations present in the product or
7		under the conditions used, the department shall immediately make the findings
8		public.
9	<u>7.</u>	A violation of this section by a manufacturer is subject to a civil penalty not to
10		exceed five thousand dollars per violation.