23-01393

SENATE STATE OF MINNESOTA NINETY-THIRD SESSION

RSI/KA

S.F. No. 1138

 (SENATE AUTHORS: KLEIN, Seeberger and Lucero)

 DATE
 D-PG
 OF

 02/02/2023
 Introduction and first reading Referred to Judiciary and Public Safety

OFFICIAL STATUS

1.1	A bill for an act
1.2 1.3 1.4	relating to consumer data protection; requiring direct-to-consumer genetic testing companies to provide disclosure notices and obtain consent; proposing coding for new law in Minnesota Statutes, chapter 325F.
1.5	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF MINNESOTA:
1.6	Section 1. [325F.995] GENETIC INFORMATION PRIVACY ACT.
1.7	Subdivision 1. Definitions. (a) For purposes of this section, the following terms have
1.8	the meanings given them.
1.9	(b) "Biological sample" means any material part of a human, discharge from a material
1.10	part of a human, or derivative from a material part of a human, including but not limited to
1.11	tissue, blood, urine, or saliva, that is known to contain deoxyribonucleic acid (DNA).
1.12	(c) "Consumer" means an individual who is a Minnesota resident.
1.13	(d) "Deidentified data" means data that cannot reasonably be used to infer information
1.14	about, or otherwise be linked to, an identifiable consumer and that is subject to:
1.15	(1) administrative and technical measures to ensure the data cannot be associated with
1.16	a particular consumer;
1.17	(2) public commitment by the company to (i) maintain and use data in deidentified form,
1.18	and (ii) not attempt to reidentify the data; and
1.19	(3) legally enforceable contractual obligations that prohibit any recipients of the data
1.20	from attempting to reidentify the data.

Section 1.

1

2.1	(e) "Direct-to-consumer genetic testing company" or "company" means an entity that:
2.2	(1) offers consumer genetic testing products or services directly to consumers; or (2) collects,
2.3	uses, or analyzes genetic data that was (i) collected via a direct-to-consumer genetic testing
2.4	product or service, and (ii) provided to the company by a consumer. Direct-to-consumer
2.5	genetic testing company does not include an entity that collects, uses, or analyzes genetic
2.6	data or biological samples only in the context of research, as defined in Code of Federal
2.7	Regulations, title 45, section 164.501, that is conducted in a manner that complies with the
2.8	federal policy for the protection of human research subjects under Code of Federal
2.9	Regulations, title 45, part 46; the Good Clinical Practice Guideline issued by the International
2.10	Council for Harmonisation; or the United States Food and Drug Administration Policy for
2.11	the Protection of Human Subjects under Code of Federal Regulations, title 21, parts 50 and
2.12	<u>56.</u>
2.13	(f) "Express consent" means a consumer's affirmative response to a clear, meaningful,
2.14	and prominent notice regarding the collection, use, or disclosure of genetic data for a specific
2.15	purpose.
2.16	(g) "Genetic data" means any data, regardless of the data's format, that concerns a
2.17	consumer's genetic characteristics. Genetic data includes but is not limited to:
2.18	(1) raw sequence data that results from sequencing a consumer's complete extracted
	DNA or a portion of the extracted DNA;
2.19	DIVA of a portion of the extracted DIVA,
2.20	(2) genotypic and phenotypic information that results from analyzing the raw sequence
2.21	data; and
2.22	(3) self-reported health information that a consumer submits to a company regarding
2.23	the consumer's health conditions and that is (i) used for scientific research or product
2.24	development, and (ii) analyzed in connection with the consumer's raw sequence data.
2.25	Genetic data does not include deidentified data.
2.26	(h) "Genetic testing" means any laboratory test of a consumer's complete DNA, regions
2.27	of a consumer's DNA, chromosomes, genes, or gene products to determine the presence of
2.28	genetic characteristics.
2.29	(i) "Person" means an individual, partnership, corporation, association, business, business
2.30	trust, or legal representative of an organization.
2.31	Subd. 2. Disclosure and consent requirements. (a) To safeguard the privacy,
2.32	confidentiality, security, and integrity of a consumer's genetic data, a direct-to-consumer
2.33	genetic testing company must:

Section 1.

01/13/23	REVISOR	RSI/KA	23-01393	as introduced

3.1	(1) provide clear and complete information regarding the company's policies and
3.2	procedures governing the collection, use, maintenance, and disclosure of genetic data by
3.3	making available to a consumer:
3.4	(i) a high-level privacy policy overview that includes basic, essential information about
3.5	the company's collection, use, or disclosure of genetic data; and
3.6	(ii) a prominent, publicly available privacy notice that includes at a minimum information
3.7	about the company's data collection, consent, use, access, disclosure, maintenance, transfer,
3.8	security, retention, and deletion practices;
3.9	(2) obtain a consumer's consent to collect, use, and disclose the consumer's genetic data,
3.10	including at a minimum:
3.11	(i) initial express consent that clearly (A) describes the uses of the genetic data collected
3.12	through the genetic testing product service, and (B) specifies who has access to the test
3.13	results and how the genetic data may be shared;
3.14	(ii) separate express consent to (A) transfer or disclose the consumer's genetic data to
3.15	any person other than the company's vendors and service providers, or (B) use genetic data
3.16	beyond the primary purpose of the genetic testing product or service and inherent contextual
3.17	uses;
3.18	(iii) separate express consent to retain any biological sample provided by the consumer
3.19	following completion of the initial testing service requested by the consumer;
3.20	(iv) informed consent in compliance with federal policy for the protection of human
3.21	research subjects under Code of Federal Regulations, title 45, part 46, to transfer or disclose
3.22	the consumer's genetic data to a third-party person for research purposes or research
3.23	conducted under the control of the company for publication or generalizable knowledge
3.24	purposes; and
3.25	(v) express consent for marketing by (A) the direct-to-consumer genetic testing company
3.26	to a consumer based on the consumer's genetic data, or (B) a third party to a consumer based
3.27	on the consumer having ordered or purchased a genetic testing product or service. For
3.28	purposes of this clause, "marketing" does not include customized content or offers provided
3.29	on the websites or through the applications or services provided by the direct-to-consumer
3.30	genetic testing company with the first-party relationship to the customer;
3.31	(3) require valid legal process to disclose genetic data to law enforcement or any other
3.32	governmental agency without a consumer's express written consent;

	01/13/23	REVISOR	RSI/KA	23-01393	as introduced
4.1	(4) devel	op, implement, and	d maintain a comp	rehensive security progr	am to protect a
4.2	consumer's	genetic data agains	t unauthorized acc	ess, use, or disclosure; a	nd
4.3	<u>(5) provi</u>	de a process for a d	consumer to:		
4.4	(i) access	s the consumer's ge	enetic data;		
4.5	(ii) delet	e the consumer's ac	count and genetic	data; and	
4.6	<u>(iii) requ</u>	est and obtain the	destruction of the o	consumer's biological sa	mple.
4.7	<u>(b) Notw</u>	vithstanding any otl	ner provisions in th	nis section, a direct-to-co	onsumer genetic
4.8	testing comp	pany is prohibited f	from disclosing a c	onsumer's genetic data v	without the
4.9	consumer's	written consent to:	(1) any entity offe	ring health insurance, lif	e insurance, or
4.10	long-term ca	are insurance; or (2) any employer of	the consumer.	
4.11	<u>Subd. 3.</u>	Enforcement. The	e commissioner of	commerce may enforce t	his section under
4.12	section 45.0	27.			
4.13	Subd. 4.	Limitations. This	section does not a	oply to:	
4.14	(1) protect	cted health informa	tion that is collected	d by a covered entity or b	usiness associate,
4.15	as those term	ns are defined in C	ode of Federal Reg	gulations, title 45, parts	160 and 164;
4.16	<u>(2) a pub</u>	lic or private instit	ution of higher edu	ucation; or	
4.17	<u>(3) an en</u>	tity owned or oper	ated by a public or	private institution of high	gher education.