GENERAL ASSEMBLY OF NORTH CAROLINA SESSION 2003

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(Public)

SESSION 2003 SENATE DRS15036-LN-7A (1/10)

Short Title: Health Care Information Privacy.

Sponsors:Senator Reeves.Referred to:

1	A BILL TO BE ENTITLED
2	AN ACT TO PROTECT HEALTH INFORMATION PRIVACY BY PROHIBITING
3	USE OR DISCLOSURE OF PROTECTED HEALTH INFORMATION FOR
4	PURPOSES OF MARKETING WITHOUT WRITTEN AUTHORIZATION OF
5	THE INDIVIDUAL, AND BY LIMITING THE USE OR DISCLOSURE OF
6	PROTECTED HEALTH INFORMATION WITHOUT INDIVIDUAL
7	AUTHORIZATION FOR CERTAIN PUBLIC HEALTH-RELATED ACTIVITIES.
8	The General Assembly of North Carolina enacts:
9	SECTION 1. The General Statutes are amended by adding the following
10	new Chapter to read:
11	" <u>Chapter 132A.</u>
12	"Privacy of Health Information.
13	" <u>Article 1.</u>
14	"Use or Disclosure of Protected Health Information.
15	" <u>§ 132A-1. Purpose.</u>
16	The purpose of this Article is to provide greater patient privacy protections regarding
17	the use or disclosure of protected health information for marketing purposes than are
18	provided under the Health Insurance Portability and Accountability Act and Privacy
19	Standards, and regulations adopted thereunder.
20	" <u>§ 132A-2. Definitions.</u>
21	(a) Unless otherwise defined in this Article, each term used in this Article has the
22	meaning and application assigned by the Health Insurance Portability and
23	Accountability Act and Privacy Standards.
24	(b) As used in this Article:
25	(1) <u>'Health Insurance Portability and Accountability Act and Privacy</u>
26	Standards' means the privacy requirements of the Administrative
27	Simplification subtitle of the Health Insurance Portability Act of 1996

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GENERAL ASSEMBLY OF NORTH CAROLINA

SESSION 2003

1			$(\mathbf{D}_{\mathbf{r}}, \mathbf{L}_{\mathbf{r}}, 104, 101)$ and the final value of anti-d $\mathbf{D}_{\mathbf{r}}$ and 102 2000 as
1			(Pub. L. 104-191) and the final rules adopted December 28, 2000, as
2		$\langle \mathbf{O} \rangle$	modified August 14, 2002, and any subsequent amendments.
3		<u>(2)</u>	'Marketing' means to make a communication about a product or
4			service to encourage recipients of the communication to purchase or
5			use the product or service, but does not include communications made
6			as part of the treatment of a patient for the purpose of furthering
7			treatment unless the covered entity receives direct or indirect
8	U.C. 100 A	2 D	remuneration from a third party for making the communication.
9			tection of private health information.
10	$\frac{(a)}{a}$	-	pt in accordance with subsection (b) of this section, a covered entity
11	<u>shall not</u>		Disclose and the life information to one outite for models in the
12		<u>(1)</u>	Disclose protected health information to any entity for marketing the
13		$\langle \mathbf{O} \rangle$	products or services of the entity; or
14		<u>(2)</u>	Use protected health information in its possession to provide
15	(1)		marketing services to any entity.
16	<u>(b)</u>		overed entity may provide marketing services to a pharmaceutical
17	<u>company</u>		covered entity:
18		<u>(1)</u>	Provides clear and conspicuous notice to the individual involved
19			concerning its disclosure practices for all protected health information
20			collected or created with regard to the individual; and
21		<u>(2)</u>	Obtains the consent of the individual involved to use the information
22			and that consent is manifested by an affirmative act in a written
23			communication which only references and applies to the specific
24			marketing purpose for which the information is to be used.
25	<u>(c)</u>		icability. – This Article does not affect the validity of another law of this
26	State that	at prov	ides greater confidentiality for information made confidential by this
27	Article.		
28	" <u>§ 132A</u>		e or disclosure of protected health information for public health
29			ities and purposes to persons subject to the jurisdiction of the Food
30		and I	Drug Administration.
31	A co	vered e	entity shall not use or disclose protected health information without the
32	written c	onsent	of the individual who is the subject of the protected health information
33	to persor	<u>is subje</u>	ect to the jurisdiction of the Food and Drug Administration except for the
34	<u>followin</u>	<u>g publi</u>	<u>c health activities or purposes:</u>
35		<u>(1)</u>	To report adverse events (or similar reports with respect to food or
36			dietary supplements), product defects or problems (including problems
37			with the use or labeling of a product), or biological product deviations
38			if the disclosure is made to the person required or directed to report the
39			information to the Food and Drug Administration;
40		<u>(2)</u>	To track products if the disclosure is made to a person required or
41			directed by the Food and Drug Administration to track the product;
42		(3)	To enable product recalls, repairs, or replacement (including locating
43			and notifying individuals who have received products of product
44			recalls, withdrawals, or other problems); or

GENERAL ASSEMBLY OF NORTH CAROLINA

SESSION 2003

1	(4) To conduct post-marketing surveillance to comply with requirements
2	or at the direction of the Food and Drug Administration.
3	" <u>§ 132A-5. Enforcement.</u>
4	(a) Injunctive Relief. – The Attorney General of this State may institute an action
5	for injunctive relief to restrain a violation of this Article.
6	(b) <u>Civil Penalties. – In addition to the injunctive relief provided by this section</u> ,
7	the Attorney General may institute an action for civil penalties against a covered entity
8	for a violation of this Article. A civil penalty assessed under this section may not exceed
9	three thousand dollars (\$3,000) for each violation. If the court in which an action under
10	this subsection is pending finds that the violations have occurred with a frequency as to
11	constitute a pattern or practice, the court may assess a civil penalty not to exceed two
12	hundred fifty thousand dollars (\$250,000).
13	(c) Disciplinary Action. – In addition to the penalties prescribed under this
14	section, a violation of this Article by an individual or facility that is licensed by an
15	agency of this State is subject to investigation and disciplinary proceedings, including
16	probation or suspension by the licensing agency. If the licensing agency finds evidence
17	that the violations of this Article constitute a pattern or practice, the licensing agency
18	may revoke the individual's or facility's license.
19	(d) <u>Availability of Other Remedies. – This Article does not affect any right of a</u>
20	person under other law to bring a cause of action or otherwise seek relief with respect to
21	conduct that is a violation of this Article."
22	SECTION 2. This act becomes effective January 1, 2004.