By: Leibowitz H.B. No. 344

A BILL TO BE ENTITLED

1	AN ACT
2	relating to reimbursement under the state Medicaid program for
3	health care services associated with certain adverse events.
4	BE IT ENACTED BY THE LEGISLATURE OF THE STATE OF TEXAS:
5	SECTION 1. Subchapter B, Chapter 32, Human Resources Code,
6	is amended by adding Section 32.0312 to read as follows:
7	Sec. 32.0312. REIMBURSEMENT PROHIBITED FOR SERVICES
8	ASSOCIATED WITH PREVENTABLE ADVERSE EVENTS. (a) In this section:
9	(1) "Health care facility" means a hospital or
10	ambulatory surgical center.
11	(2) "Health care provider" means a physician or health
12	<pre>care facility.</pre>
13	(3) "Infant" means a child younger than one year of
14	age.
15	(4) "Serious disability" means:
16	(A) a physical or mental impairment that
17	substantially limits one or more major life activities of ar
18	individual such as seeing, hearing, speaking, walking, or
19	breathing, or a loss of a bodily function, if the impairment or loss
20	lasts more than seven days or is still present at the time of
21	discharge from an inpatient health care facility; or
22	(B) loss of a body part.
23	(5) "Serious injury" means a bodily injury that
24	results in:

1	(A) death;
2	(B) permanent and serious impairment of an
3	important bodily function; or
4	(C) permanent and significant disfigurement.
5	(b) The department may not provide reimbursement under the
6	medical assistance program to a health care provider for a health
7	care service provided in association with a preventable adverse
8	event involving a recipient of medical assistance while in the
9	provider's care, including a health care service provided as a
10	result of or to correct the consequences of a preventable adverse
11	event.
12	(c) For purposes of this section, a preventable adverse
13	event is any of the following events involving a recipient of
14	<pre>medical assistance:</pre>
15	(1) surgery performed on the wrong body part that is
16	not consistent with the documented informed consent for that
17	recipient, excluding a situation requiring prompt action that
18	occurs in the course of surgery or an urgent situation that
19	<pre>precludes obtaining informed consent;</pre>
20	(2) surgery performed on the wrong person;
21	(3) the wrong surgical procedure performed on the
22	recipient that is not consistent with the documented informed
23	consent for that recipient, excluding a situation requiring prompt
24	action that occurs in the course of surgery or an urgent situation
25	that precludes obtaining informed consent;
26	(4) the unintended retention of a foreign object in
27	the recipient after surgery or another procedure;

(5) death during or immediately after surgery if the 1 2 recipient would be classified as a normal, healthy patient under guidelines published by a national association 3 4 anesthesiologists; 5 (6) death or serious disability caused by the use of a 6 contaminated drug, device, or biologic provided by a health care provider if the contamination is the result of a generally 7 8 detectable contaminant in drugs, devices, or biologics regardless 9 of the source of the contamination or product; (7) death or serious disability caused by the use or 10 function of a device during the recipient's care in which the device 11 12 is used for a function other than as intended; (8) death or serious disability caused by an 13 14 intravascular air embolism that occurs while the recipient is 15 receiving care in a health care facility, excluding a death associated with a neurological procedure known to present a high 16 17 risk of intravascular air emoblism; 18 (9) an infant being discharged to the wrong person; 19 (10) death or serious disability associated with the recipient's disappearance for more than four hours, excluding the 20 21 death or serious disability of an adult recipient who has 22 decision-making capacity; 23 (11) suicide or attempted suicide resulting in serious 24 disability while the recipient is receiving care in a health care facility if the suicide or attempted suicide is due to the 25

recipient's actions after admission to the facility, excluding a

death resulting from a self-inflicted injury that was the reason

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for the recipient's admission to the facility; 2 (12) death or serious disability caused by a medication error, including an error involving the wrong drug, 3 wrong dose, wrong patient, wrong time, wrong rate, wrong 4 5 preparation, or wrong route of administration; 6 (13) death or serious disability caused by a hemolytic reaction resulting from the administration of ABO- or 7 8 HLA-incompatible blood or blood products; (14) subject to Subsection (d), death or serious 9 disability caused by labor or delivery in a low-risk pregnancy 10 while the recipient is receiving care in a health care facility, 11 12 including death or serious disability occurring not later than 42 days after the delivery date; 13 14 (15) death or serious disability directly related to 15 hypoglycemia, the onset of which occurs while the recipient is receiving care in a health care facility; 16 17 (16) death or serious disability, including kernicterus, caused by failure to identify and treat 18 19 hyperbilirubinemia in a neonate before discharge from a health care 20 facility; (17) stage three or four pressure ulcers acquired 21

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admission;

manipulative therapy;

27 (19) death or serious disability caused by an electric

after admission to a health care facility, excluding progression

from stage two to stage three if stage two was recognized on

(18) death or serious disability resulting from spinal

- 1 shock while the recipient is receiving care in a health care
- 2 facility, excluding an event involving a planned treatment such as
- 3 electric countershock;
- 4 (20) an incident in which a line designated for oxygen
- 5 or other gas to be delivered to the recipient contains the wrong gas
- 6 or is contaminated by a toxic substance;
- 7 (21) death or serious disability caused by a burn
- 8 incurred from any source while the recipient is receiving care in a
- 9 health care facility;
- 10 (22) death or serious disability caused by a fall
- 11 while the recipient is receiving care in a health care facility;
- 12 (23) death or serious disability caused by the use of a
- 13 restraint or bed rail while the recipient is receiving care in a
- 14 health care facility;
- 15 (24) an instance of care for the recipient ordered or
- 16 provided by an individual impersonating a physician, nurse,
- 17 pharmacist, or other licensed health care professional;
- 18 (25) abduction of the recipient from a health care
- 19 facility;
- 20 (26) sexual assault of the recipient within or on the
- 21 grounds of a health care facility;
- 22 (27) death or significant injury resulting from a
- 23 physical assault of the recipient that occurs within or on the
- 24 grounds of a health care facility; and
- 25 (28) artificial insemination with the wrong donor
- 26 sperm or donor egg.
- 27 (d) For purposes of Subsection (c)(14), the death of a

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- 1 recipient of medical assistance resulting from pulmonary or
- 2 amniotic fluid embolism, acute fatty liver of pregnancy, or
- 3 cardiomyopathy during the course of labor or delivery is not
- 4 considered a preventable adverse event.
- 5 (e) The department's refusal to reimburse a health care
- 6 provider under this section does not in itself create civil
- 7 liability and is not subject to discovery or admissible in any civil
- 8 action against the provider.
- 9 SECTION 2. Not later than November 1, 2009, the executive
- 10 commissioner of the Health and Human Services Commission shall
- adopt rules necessary to implement Section 32.0312, Human Resources
- 12 Code, as added by this Act.
- SECTION 3. Section 32.0312, Human Resources Code, as added
- 14 by this Act, applies only to a preventable adverse event occurring
- on or after the effective date of the rules adopted by the executive
- 16 commissioner of the Health and Human Services Commission under
- 17 Section 2 of this Act.
- SECTION 4. If before implementing any provision of this Act
- 19 a state agency determines that a waiver or authorization from a
- 20 federal agency is necessary for implementation of that provision,
- 21 the agency affected by the provision shall request the waiver or
- 22 authorization and may delay implementing that provision until the
- 23 waiver or authorization is granted.
- SECTION 5. This Act takes effect September 1, 2009.