



Rep. Mary E. Flowers

**Filed: 4/6/2011**

09700HB1476ham002

LRB097 06656 CEL 53877 a

1 AMENDMENT TO HOUSE BILL 1476

2 AMENDMENT NO. \_\_\_\_\_. Amend House Bill 1476, AS AMENDED, as  
3 follows:

4 on page 1, immediately below line 3, by inserting the  
5 following:

6 "Section 3. The Hospital Licensing Act is amended by adding  
7 Section 9.7 as follows:

8 (210 ILCS 85/9.7 new)

9 Sec. 9.7. Health care facility requirements to report  
10 adverse events resulting in patient death or serious  
11 disability.

12 (a) Definitions. As used in this Act:

13 (1) "Death" means a patient's death related to an  
14 adverse event and not related solely to the natural course  
15 of the patient's illness or underlying condition. Events

1 otherwise reportable shall be reported even if the death  
2 might have otherwise occurred as the natural course of the  
3 patient's illness or underlying condition.

4 (2) "Serious disability" means a physical or mental  
5 impairment, including loss of a body part, related to an  
6 adverse event and not related solely to the natural course  
7 of the patient's illness or underlying condition, that  
8 substantially limits one or more of the major life  
9 activities of an individual or a loss of bodily function,  
10 if the impairment or loss lasts more than 7 days prior to  
11 discharge or is still present at the time of discharge from  
12 an inpatient health care facility.

13 (b) Notwithstanding any other reporting requirements of  
14 State law or regulation, each health care facility shall report  
15 to the Department, in the form and manner required by the  
16 Department, the occurrence of any of the adverse health care  
17 events described in subsections (c) through (h) of this Section  
18 no later than 30 days after discovery of the event. The  
19 Department shall determine from the information provided in the  
20 report if there is a basis for further investigation to  
21 determine if there were any violations of the standards and  
22 procedures covered by this Act.

23 (c) Surgical events reportable under this subsection are:

24 (1) surgery performed on a wrong body part that is not  
25 consistent with the documented informed consent for that  
26 patient; reportable events under this clause do not include

1 situations requiring prompt action that occur in the course  
2 of surgery or situations whose urgency precludes obtaining  
3 informed consent;

4 (2) surgery performed on the wrong patient;

5 (3) the wrong surgical procedure performed on a patient  
6 that is not consistent with the documented informed consent  
7 for that patient; reportable events under this clause do  
8 not include situations requiring prompt action that occur  
9 in the course of surgery or situations whose urgency  
10 precludes obtaining informed consent;

11 (4) retention of a foreign object in a patient after  
12 surgery or other procedure, excluding objects  
13 intentionally implanted as part of a planned intervention  
14 and objects present prior to surgery that are intentionally  
15 retained; and

16 (5) death during or immediately after surgery of a  
17 normal, healthy patient who has no organic, physiologic,  
18 biochemical, or psychiatric disturbance and for whom the  
19 pathologic processes for which the operation is to be  
20 performed are localized and do not entail a systemic  
21 disturbance.

22 (d) Product or device events reportable under this  
23 subsection are:

24 (1) patient death or serious disability associated  
25 with the use of contaminated drugs, devices, or biologics  
26 provided by the health care facility when the contamination

1 is the result of generally detectable contaminants in  
2 drugs, devices, or biologics regardless of the source of  
3 the contamination or the product;

4 (2) patient death or serious disability associated  
5 with the use or function of a device in patient care in  
6 which the device is used or functions other than as  
7 intended; "device" includes, but is not limited to,  
8 catheters, drains, and other specialized tubes, infusion  
9 pumps, and ventilators; and

10 (3) patient death or serious disability associated  
11 with intravascular air embolism that occurs while being  
12 cared for in a health care facility, excluding deaths  
13 associated with neurosurgical procedures known to present  
14 a high risk of intravascular air embolism.

15 (e) Patient protection events reportable under this  
16 subsection are:

17 (1) an infant discharged to the wrong person;

18 (2) patient death or serious disability associated  
19 with patient disappearance for more than 4 hours, excluding  
20 events involving adults who have decision-making capacity;  
21 and

22 (3) patient suicide or attempted suicide resulting in  
23 serious disability while being cared for in a health care  
24 facility due to patient actions after admission to the  
25 health care facility, excluding deaths resulting from  
26 self-inflicted injuries that were the reason for admission

1       to the health care facility.

2       (f) Care management events reportable under this  
3 subsection are:

4           (1) patient death or serious disability associated  
5 with a medication error, including, but not limited to,  
6 errors involving the wrong drug, the wrong dose, the wrong  
7 patient, the wrong time, the wrong rate, the wrong  
8 preparation, or the wrong route of administration,  
9 excluding reasonable differences in clinical judgment on  
10 drug selection and dose;

11           (2) patient death or serious disability associated  
12 with a hemolytic reaction due to the administration of ABO  
13 incompatible blood or blood products;

14           (3) maternal death or serious disability associated  
15 with labor or delivery in a low-risk pregnancy while being  
16 cared for in a health care facility, excluding deaths from  
17 pulmonary or amniotic fluid embolism, acute fatty liver of  
18 pregnancy, or cardiomyopathy; and

19           (4) patient death or serious disability directly  
20 related to hypoglycemia, the onset of which occurs while  
21 the patient is being cared for in a health care facility  
22 for a condition unrelated to hypoglycemia.

23       (g) Environmental events reportable under this subsection  
24 are:

25           (1) patient death or serious disability associated  
26 with an electric shock while being cared for in a health

1 care facility, excluding events involving planned  
2 treatments such as electric countershock;

3 (2) any incident in which a line designated for oxygen  
4 or other gas to be delivered to a patient contains the  
5 wrong gas or is contaminated by toxic substances;

6 (3) patient death or serious disability associated  
7 with a burn incurred from any source while being cared for  
8 in a health care facility that is not consistent with the  
9 documented informed consent for that patient; reportable  
10 events under this clause do not include situations  
11 requiring prompt action that occur in the course of surgery  
12 or situations whose urgency precludes obtaining informed  
13 consent;

14 (4) patient death associated with a fall while being  
15 cared for in a health care facility; and

16 (5) patient death or serious disability associated  
17 with the use of restraints or bedrails while being cared  
18 for in a health care facility.

19 (h) Physical security events reportable under this  
20 subsection are:

21 (1) any instance of care ordered by or provided by  
22 someone impersonating a physician, nurse, pharmacist, or  
23 other licensed health care provider;

24 (2) abduction of a patient of any age;

25 (3) sexual assault on a patient within or on the  
26 grounds of a health care facility; and

1           (4) death or significant injury of a patient or staff  
2           member resulting from a physical assault that occurs within  
3           or on the grounds of a health care facility."