

PATIENT IDENTIFICATION STICKER

PATIENT SAFETY EVENT REPORT

A **Patient Safety Event** is an unexpected occurrence which results in an adverse effect on the patient or has the **potential** for an adverse effect.

Medication events report on Medication Safety Report.

Falls report on Falls Event Report.

Patient Room #/ Department: _____
 Date of Event: _____ Time of Event: _____
 Location of Event: _____

Description of Event and Follow-Up Treatments/ Planned Evaluations:

Type of Event (Check ONE Box)

TREATMENT/PROCEDURE/DELIVERY RELATED

- | | |
|---|---|
| <input type="checkbox"/> Wrong patient/site/procedure or treatment | <input type="checkbox"/> Burn (2 nd or 3 rd degree) |
| <input type="checkbox"/> Unplanned treatment/procedure or return to OR | <input type="checkbox"/> AMI within 48 hours of procedure |
| <input type="checkbox"/> Procedure/treatment related injury/complication | <input type="checkbox"/> Retained Foreign Body |
| <input type="checkbox"/> Transfer to higher level of care related to event | <input type="checkbox"/> Unexpected death |
| <input type="checkbox"/> Cardiopulmonary or Respiratory arrest (<i>unrelated to primary reason for admission</i>) | |
| <input type="checkbox"/> Loss or impairment of limb/organ/body function | <input type="checkbox"/> Brain/CNS impairment |
| <input type="checkbox"/> IV infiltrate/phlebitis with physician intervention | <input type="checkbox"/> Moderate sedation related event |
| <input type="checkbox"/> Hemolytic transfusion reaction | <input type="checkbox"/> Anaphylaxis |
| <input type="checkbox"/> New Tissue Trauma (<i>stage 3 or greater</i>) | <input type="checkbox"/> Birth-related injuries |

EMBOLIC RELATED EVENTS

- | | |
|--|---------------------------------------|
| <input type="checkbox"/> New (hospital-acquired) DVT | <input type="checkbox"/> Air embolism |
| <input type="checkbox"/> New (hospital-acquired) pulmonary embolism (PE) | |

OTHER

- | | |
|--|---|
| <input type="checkbox"/> Breach in HIPAA/Confidentiality | <input type="checkbox"/> Suicide/Attempted Suicide |
| <input type="checkbox"/> Alleged Abuse or Rape | <input type="checkbox"/> Elopement |
| <input type="checkbox"/> Radiation Misadministration | <input type="checkbox"/> Infant abduction/discharge to wrong family |
| <input type="checkbox"/> Power outage/termination of services impacting patient care | |
| <input type="checkbox"/> Fire | <input type="checkbox"/> Other |

EQUIPMENT

- ☐ Equipment Malfunction during Treatment
- Equipment/Device Name: _____
 Manufacturer: _____
 Model #: _____
 Serial #: _____

Patient Outcome (*circle one*)

1. No change in patient condition
 2. Injury or complication*
 3. Death*
- * If significant injury/complication or death, call Patient Safety & QI Department **immediately** at 525-1230

Contributing Factors

(*circle one, if known*)

POLICY/PROCESS

1. Procedure/protocol not followed
2. Procedure/protocol not effective

HUMAN RESOURCE FACTORS

3. Performance/Knowledge Deficit
4. Distractions
5. Shift Change
6. Staffing Level Related

ENVIRONMENT/EQUIPMENT

7. Unsafe/unsecured Environment
8. Unsafe Equipment
9. Emergency Situation

COMMUNICATION/INFORMATION

10. Patient Assessment/Monitoring
11. Communication Breakdown
12. Documentation Related

OTHER: _____

NAME (please print) OF PERSON COMPLETING REPORT*

DATE REPORT COMPLETED

*No blame will be allocated to employees or physicians following their reporting of unintended changes in health status, injury or harm.

Lawson # 200976

Significant and Sentinel Events Require Immediate Reporting to Patient Safety & Quality Improvement Department

Phone: 525-1230

Fax 525-1908

Refer to Event Reporting Policy AD-035 for examples of Significant/Sentinel Events.