

Attending and Advanced Practitioner Orientation

Risk Management/Patient Safety 2021

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Risk Management/Patient Safety Overview

Resource for organization – available 24/7

- Call ext 7475 during business hours
- Call hospital operator to obtain on-call name and pager #

Risk Oversight:

- Regulatory reporting and compliance
 - DPH - Department of Public Health
 - TJC - The Joint Commission
 - BORM - Board of Registration of Medicine
 - BORN - Board of Registration in Nursing
 - FDA - Food and Drug Administration (MedWatch)
 - Root Cause Analysis (RCA)
 - Review of event
 - Determine root cause & preventability
 - Develop action plan
 - Communicate lessons learned
 - Audit Improvements
 - Assist providers with disclosure and apology - call for guidelines & see Disclosure policy
 - Safety Reporting System - SafeSpot
 - Coordinate clinical complaints/grievances with Patient Relations
 - Terminations of patient/physician relationships - call for guidelines & see Termination policy
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High Reliability Organization - HRO

Just Culture

HRO behaviors: Culture of safety

- Preoccupation with risk of failure, mitigation, and better design
- Relentless consistency
- Team training
- Safe reporting environment
- Open communications
- Colleague respect
- Focus on zero harm

Just Culture: Building a culture of Learning

- Accountability
 - Designing better systems
 - Managing human behaviors - Human Error, At-Risk Behavior, Reckless
 - Systematic learning - Investigation
 - Finding Justice - The 3 Duties:
 - Avoid harm
 - Follow a procedural rule
 - Produce an outcome
 - Implementation - commitment of the organization
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DPH - Dept. of Public Health

- Site visit - Events/Grievances
- Serious Reportable Events (SRE): (e.g. - Falls with injury, Medication Error harm, surgical retained foreign object)
 - RCA - Each SRE requires a Root Cause Analysis
 - Review of pertinent hospital policies

BORM – Board of Registration in Medicine

- Death from elective surgery, unanticipated deaths, wrong site procedure
 - Call Risk x 7475 if you receive a letter from BORM
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Regulatory (cont'd)

FDA – Food & Drug Administration

- Death or serious injury related to equipment requires reporting via MedWatch form

CMS - Centers for Medicare & Medicaid Services

- Unannounced site visit

TJC – The Joint Commission (accrediting agency)

- Unannounced visit every three years

DEA - Drug Enforcement Administration

- Diversion or theft
 - Opioid Prescribing - Adhere to MassPAT - PMP
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Serious Reportable Event (SRE)

http://www.qualityforum.org/Topics/SREs/Serious_Reportable_Events.aspx

SRE examples reportable to DPH - serious injury or death:

Surgical or Invasive Procedure Events - Wrong side/site procedure, incorrectly placed tubes, unintended retained foreign object etc.

Product or Device - Use of contaminated drugs, devices or biologics provided by the healthcare setting, intravascular air embolism etc.

Patient Protection Events - Discharge or release of a patient, of any age, who is unable to make decisions, suicide or attempted suicide while cared for in healthcare setting

Care Management Events - Medication error, unsafe administration of blood products, falls with fracture/laceration, Stage 3 or 4 or Unstageable pressure ulcers, irreplaceable lost biological specimen, failure to follow-up or communicate test results

Serious Reportable Event (SRE)

– cont'd

Environmental Event - Electronic shock in the course of care, patient burn from any source, physical restraints

Radiologic Events - Patient or staff injury associated with introduction of a metallic object in the MRI area

Potential Criminal Events - Abduction of a patient of any age, sexual assault/abuse or physical assault of a patient or staff

Patient Experience

What Do Our Patients Want?

- Introduce yourself, your specialty, and why you are there
 - Explain tests ordered and provide timely results
 - Review discharge plans – explanations that are understandable
 - Engage patient/family in the plan of care – offer opportunity to ask questions
 - Coordination of care from the healthcare team
 - Quality and safe care
 - To be listened to
 - Accessible and or return patient calls
 - Avoid blaming staff or departments
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Safety Reporting

Web-based safety reporting system – SafeSpot

Access on MassNet home page and log-in

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MassNet



Safety Reporting (cont'd)

Reporting mechanism for occurrences

- Adverse outcome
- Potential or near miss
- Unexpected events
- Process or safety issues
- Employee events
- Compliance issues

Who should report?

- All employees are responsible to report
 - Employee with knowledge about the occurrence
 - Employee that observed event
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Safety Reporting (cont'd)

Use descriptive facts NOT opinions

Confidential, peer review protected

Not part of the medical record - Do not mention safety report was filed - (document facts of occurrence when pertinent)

What do we do with safety reports?

- Internal investigation
 - Track and trend data
 - Report to multiple Committees
 - Identify potential risk
 - Identify opportunities for improvement
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Safe Practices for Patient Safety

Chain of command

Hand-off communication

Electronic medical record (EMR) documentation – EPIC

Teamwork and rounding for coordinated consistent care

Supervision of residents

Follow Policies and Procedures

- Based on best practices
 - Regulatory requirements
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Specimen Debriefing with 2 staff – Patient is still in the room :

- Label the specimen – Patient name, DOB, & MRN
- Specimen source and laterality, if applicable
- Correct tests are ordered
- Visualize the specimen in the container

Medication Reconciliation:

- Accurate medication list – engage the patient or family member
- Compare home medication list to ordered medicines
- Resolve discrepancies
- Ensure upon discharge that patient understands which meds they should take and those that should be discontinued

Health Care Proxy Activation

- Utilize EPIC SMART Phrase
- Document name of HCP, patient incapacity to make decisions & approximate time limit

Dobhoff Insertion – Two-Step Process

- X-ray confirmation of placement and documentation before use
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Safe Practices for Patient Safety - Accessing Policies

Access policies on MassNet homepage
Policy and procedures tab – Policy Library

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Clinical Policy Library

Quality and Safety | Nursing | OR Policies | Infection Prevention and Control | ED | Trauma | Employee Health | Rehab Services | **Search**

- [-] Clinical and Administrative Manual
 - [+] 02.Administrative/Legal
 - [+] 03.Patient Care/Clinical Services

Title Search

Title Search Options: any sequence

Content Search



Documentation Guidelines

Correct medical record/patient

- Patient identification – name and DOB
- Some documents are scanned into EPIC – electronic medical record

Write legibly on any paper documents such as consents

Correcting errors or late entries/addendums – see EPIC Tip Sheet

Access EPIC tip sheet:

- MassNet home page



/ Tip Sheets & eLearnings

Documentation Guidelines (cont'd)

High risk documentation:

- critical values and tests
- condition deterioration
- transfer to higher level of care

Plan of care documentation for abnormal tests

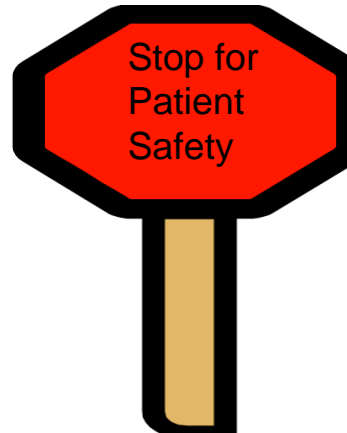
- communication for other providers
- prevent delays or missed diagnosis
- communicate incidental findings

Document discussion/disclosure after an adverse outcome -
refer to “Disclosure of Adverse Events” policy

Emails – brief and with caution

- Refer to “Information Systems: E-mail communication” policy
 - Discoverable
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Thank You



Call Risk Management early and often
Ext. 7475 (main office)

We are here to help you!
