Risk Management House Staff Orientation to Montefiore Medical Center

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Risk Management Topics

- MMC Malpractice Insurance Coverage
- RM Administrative Policies and Procedures
 - Patient Consent, Patient Elopement, Discharge AMA
 - Occurrence Reporting
 - Tracking of Implanted Medical Devices
 - Safe Use of Medical Devices
- Office of Medical Director's Policy
 Disclosure of Unanticipated Events

Goals of Risk Management

- Promote Quality Care
 Promote Patient Safety
 Prevent Injuries
- Minimize Liability & Prevent Financial Loss to MMC

• Provide Staff Education and Inservices

When should you contact Risk Management?

- Questions re: Medical Malpractice Insurance coverage
- If you receive any legal documents
- Requests or inquires from attorneys
- Emergent medico-legal patient care issues
- Reporting accidents or potential liabilities

How to contact Risk Management

• 3328 Rochambeau Avenue

• 920-6340, 920-6733

• Office Hours: 8-30 to 5:00 weekdays

• On Call Representative 24/7 for emergent situations: Contact Administrator on Duty

MMC Malpractice Insurance

- Covers claims arising from activities within the scope of your employment at MMC
- Occurrence based policy
- Provides \$1M per occurrence, \$3M in the aggregate, plus excess layers
- Does <u>not</u> cover criminal acts, actions <u>not</u> within scope of employment, or when employed by <u>other than MMC</u>

When should you file an Occurrence Report?

- When an unanticipated event occurs resulting in injury/illness
 - Could be reportable to NY State Department of Health

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• Could result in claims against MMC

- Employee incidents involving sharps
 Complete Employee Accident/Injury Form Occupational Health Services Human Resources
 - Complete Occurrence Report Risk Management

CCUITTENCE	THE FOLLOWING IS PREP ARED FOR THE HOSP ITAL'S QUALITY ASSURANCE PROGRAM. IT IS CONFIDENTIAL AND PROTECTED FROM DISCLOSURE PURSUANT TO NEW YORK STATE EDUCATION LAW 6527 (3) AND NEW YORK STATE PUBLIC HEALTH LAW 2805-m.	ample form
Reporting	Form D: PATENT MALE AGE DAYS MONTHS VISITOR FEMALE MONTHS YEARS PROCEDURE ADMITTING DIAGNOSS PROCEDURE PERSON PREPARING REPORT LAST NAME FIRST NAME	ADDRESSOGRAPH
Form	I FALLS RELATED TO: OCCURRENCE INFORMATION Assisted to Floor Ambulating Assessed at Risk Yes No Fall Alleged Bathroom Protocol in P lace Yes No Unk Fall Witnessed Bed/Crib Patient is Oriented Confused Ambulatory Found on Floor Chair Activity Privaleges Ambulatory Non-Ambulatory Other Person Siderails Complete Pariati None Stretcher/Table Environmental Wet Debris None Unknown Restraints Yes No Unk	SEDATIVE WITHIN LAST 4 HOURS IF YES, DRUG
	2. SKIN 6. OBSTETRICS 7. PERIOPERATIVE/PERIPROCEDURAL Bum Apgar <5 @ 5 Minutes WI 30 DAYS Decubitus (Community) Brachial P ksus/Etb's ASA1 II III IV Circumcision with Repair Facial PE ksus/Etb's ASA1 II III VI J. LOST PATIENT P ROPERTY Forceps Related Date Articles/Clothing Hysterectomy-Pregnant Woman AMI	MANUFACTURER
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	Visitor Actions IL PROCEDURES Visitor Actions Langer Dialysis Pacemaker Thora/Paracentesis Biopsy Intubation PTCA Tube/Catheter Placement Cardiac Cath Laser Scope Other SPECIFY Implanted htravas cular Device	Patient D Technique Pharmacy Time 12. UNEXPECTED EVENT NON-ILLNESS RELATED Aspiration (Related to Death Loss of Limb Conscious Sedation) Impairment of Limb Surgical Interventio
	Central Line Venous Access	CAC/Respiratory Arrest Loss/Impairment Body Function

OCCURRENCE REPORTING FORM Facility ID:

Occi

Consent Issues

Risk Management

- Informed Consent
- Refusal to Consent
- Administrative Review/Patient Alone
- Miscellaneous Consent Issues

Director of Clinical Affairs

Lynn Richmond 920-7052/page operator

- DNR/DNI
- Withhold/Withdraw Life Sustaining Treatment
- Brain Death

Safe Medical Devices

<u>SMDA 1990</u> mandates reporting to the FDA or Manufacturer of all serious injuries/deaths caused by medical devices.

Medical device & malfunction are broad terms.

Whenever a device malfunction injures a patient: Care for the patient

Sequester the device and mark "out of service"

Document event in Medical Record

Complete an Occurrence Report

Call Risk Management

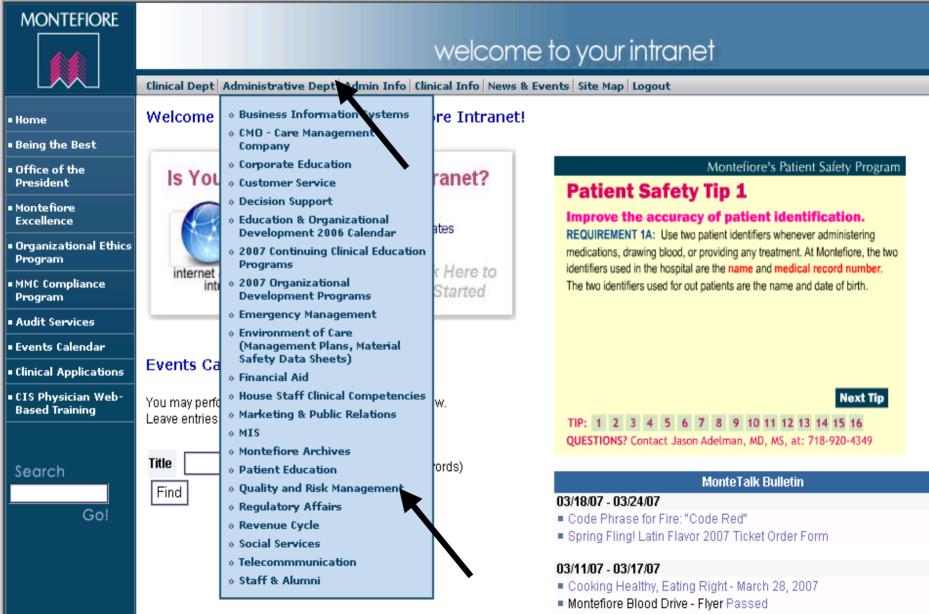
DO NOT RETURN DEVICE TO MANUFACTURER!

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Disclosure of Unanticipated Events

- 1. Recognition of unanticipated clinical event
- 2. Provision of immediate patient care
- 3. Escalation to to appropriate staff as outlined in Administrative P&P for Critical Events
- 4. Identification of needed Support Services & Investigation of event
- 5. Choosing the Individual, Timing, & Content of Disclosure
- 6. Documentation of discussion in medical record

On line resources



Shackleton Client Needs Analysis

MMC Intranet Risk Management Website

- General Info re Risk Management Department
- Insurance Information
- Risk Management Advisories Newsletters highlighting key risk related topics
- Administrative Policy and Procedure Manual



MONTEFIORE



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Being the Best

Office of the President

Montefiore Excellence

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MMC Compliance Program

Audit Services

Events Calendar

Clinical Applications

CIS Physician Web-**Based Training**

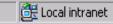
Admin Info » Policies/Procedures » Administrative Policy and Procedure Manual Refresh Search Consents Current Search: **# B**/ Eff. Date Reviewed Revised PnP Index Title/comments Current Version All Administrative Policy and Procedures were reviewed in 2006 by Policy Owner(s) and revised as necessary. Ð CONSENTS View Top 2/07 JC10.1 7/06 12/03 Consents, Informed Consent and Refusal 2/07 7/06 11/03 ** click here for English version of Consent form AND click here for Spanish version of 11/18/64 12/03 8/03 Consent form 11/03/03 12/00 9/96 7/94 JC12.1 9/06 01/03 Consent for Sterilization and Hysterectomies 06/20/78 05/17/79 01/03 Search Go!

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Department of Quality and Risk Management

Montefiore Medical Center 111 East 210th Street Bronx, NY 10467





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Risk Management Advisory Issues 8, 13, 21 = Consent



The Department of Risk Management publishes a Risk Management Advisory to highlight key risk-related topics. The **ADVISORY** is intended to be a quick reference that addresses questions we are commonly asked. **The focus of this issue is Informed Consent and Refusal to Consent. We encourage each of our readers to share this issue with the staff of your department/unit.** If there is a topic you would like to see discussed in future issues, please call us at 920-6340 or 920-6733 and let us know.

INFORMED CONSENT AND REFUSAL TO CONSENT - AN OVERVIEW

What is Informed Consent?

Informed Consent is the authorization given by a patient with decisional capacity for medical tests or

QUESTIONS ? Contact: Risk Management

920-6733

920-6340

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