

Policy: Clinical Alarm Management

ORGANIZATIONAL: Affects two or more departments.							
Folder	Organizational Choices: Patient Safety and Performance Imp			Sub-Folder (If Applicable)	n/a		
Original Effective Date	11/1/2015	Scope	What departments does this policy apply to? State "All" as is may apply to the entire organization. All				
Approved (Approver/Date)	MDPRC 5/16/2019; MEC 5/28/2019						
Last Reviewed/ Revised Date	5/28/2019	OSHA Category (If Applicable)	III	Standard (If Applicable)	NPSG.06.01.01	Number of pages	4

PURPOSE: To define an alarm management process and promote the monitoring of patients through the safe use of and response to clinical alarms

GUIDELINES:

1. Biomedical Engineering (Biomed) is responsible for performing regular preventative maintenance and testing on all alarms on patient physiological monitoring and patient care equipment except on contracted equipment, loaners or demos.
2. Clinical Directors/Managers are responsible for assuring individuals adhere to the requirements of this policy and these procedures are implemented and followed. Instances of non-compliance are reported via the event reporting process.
3. The patient's primary nurse and Centralized Telemetry Monitoring (CTM) will be made aware of changes in the alarm parameters.
4. Cardiac monitor alarm functionality is monitored through Preventative Maintenance (PM) performed by Biomed; this validation report is presented to the Environment of Care (EOC) Committee which then reports to the Quality Committee of the Board.

PROCEDURE:

- A. Medical Equipment/Device Alarms/ Patient Monitoring Alarms
 1. Staff that use equipment with alarms must check settings to ensure they are appropriate and that audible alarms will be clearly discernible relative to ambient and competing noise on their respective clinical unit.
 2. Staff that use medical equipment will know how to adjust the alarm parameters if the equipment allows them to be changed.
 3. Staff shall not bypass, shut off, or adjust alarm volumes to a level that cannot be readily heard when the alarm activates. Bypass of an alarm function is reported as a patient safety event.
 4. Hospital staff trained to treat the patient must respond to alarms.
 5. The following table, which is not all inclusive of all alarms, includes details for the equipment with alarms that have been identified as possibly contributing to alarm fatigue for SoutheastHEALTH:

Type of Equipment	What are the clinically appropriate settings for the alarm signals?	Who has the authority to set default alarm parameters?	When should alarm parameters be changed?	Who has the authority to change alarm parameters?	When should alarm parameters be disabled?	What is the PM schedule?
Central Monitor with Nihon Kohden (NK) Bedside Monitors - Telemetry (Heart Rate/Rhythm) >17 years old	Low HR 50 High HR 130	MDPRC or MEC	Per the patient condition	Designated, trained personnel (LIP and/or staff who have passed Arrhythmia Exam)	At no time in general, unless it is patient specific with physician order	Alternative equipment management (AEM)
Central Monitor with NK Bedside Monitors - Blood Pressure >17 years old	High SBP 160 Low SBP 90 High DBP 90 Low DBP 50	MDPRC or MEC	Per the patient condition	Designated, trained personnel	At no time in general, unless it is patient specific with physician order	AEM
Central Monitor with NK Bedside Monitors - Pulse Ox (Oxygen Saturation) >17 years old	High OFF Low 90	MDPRC or MEC	Per the patient condition	Designated, trained personnel	At no time in general, unless it is patient specific with physician order	AEM
Corometrics	<i>Fetal Heart Rate</i> - Default (100-180) Min/Max (30-240) <i>Blood Pressure</i> - Default (90-160/50-90) Min/Max (30-50/130/240) <i>Mom's Heart Rate</i> - Default (50-130) Min/Max (35-250) <i>Pulse Ox</i> - Default (95-100%) Min/Max (80-100%)	MDPRC or MEC	Per the mom's or baby's condition	Designated, trained personnel	Fetal Heart Rate - Fetal Demise	AEM
EtCO2 Monitor	EtCO2 – 10-55 mmHg RR – 6-40 SpO2 - <90% HR - 50-130	MDPRC or MEC	Per the patient condition	Designated, trained personnel	At no time or per a physician order	AEM
Infant Warmers	Not to exceed 37°C	Manufacturer	Per the patient condition	Physician and RN per a physician order according to the patient's condition.	At no time or per a physician order	1) RT checks air & O2 tanks every night and if needed re-stocks supplies 2) AEM
Bed Alarms / Call Lights	<i>Bed Alarms</i> : 3 levels of alarms including: Off, Starting to get up, and Out of bed <i>Call Lights</i> : First goes to the Patient Care Tech and if no response within 30 seconds it goes to the nurse and if no response within another 30 seconds it goes to the overhead <i>Emergency Room</i> – There are no settings on the call lights	Bed Alarms: Designated, trained personnel Call Lights: N/A	Bed Alarms: Per the patient condition Call Lights: At no time	Bed Alarm: Designated, trained personnel Call Lights: N/A	Bed Alarms: Getting patient out of bed for transport, bathroom, etc. Call Lights: N/A	Bed Alarms: Per manufacturer guidelines Call Lights: Per manufacturer guidelines

Type of Equipment	What are the clinically appropriate settings for the alarm signals?	Who has the authority to set default alarm parameters?	When should alarm parameters be changed?	Who has the authority to change alarm parameters?	When should alarm parameters be disabled?	What is the PM schedule?
Fluid Pumps - IV Infusion, PCA, Epidural and Kangaroo (Feeding) Need to separate PCA out of this category. PCA is required to be manufacturer PM.	Infusion complete, air in line, occlusions, low battery or pump failures	Manufacturer	At no time	N/A	At no time or per a physician order	AEM
Dialysis Machine	Venous & Arterial alarm limits are adjusted to achieve the prescribed flow rate	Critical Care Committee	Per the patient condition	Designated, trained personnel	At no time or per a physician order	Per manufacturer guidelines
BIPAP Machine	10-15% above or below the settings ordered by the physician	Critical Care Committee	Per the patient condition	Respiratory Therapy based on physician order	At no time or per a physician order	Per manufacturer guidelines
Ventilators	10-15% above or below the settings ordered by the physician	Critical Care Committee	Per the patient condition	Respiratory Therapy based on physician order	At no time or per a physician order	Every 6 months by the vendor & Biomed performs PMs per manufacturer guidelines

6. Pediatric Unit Specific Alarm Limits

- i. The below parameters will be updated on each cardiac monitor according to the child's normal vital sign range or as specified by the LIP:

Age	Respiratory Rate	Pulse	Systolic Blood Pressure
Preterm newborn	55-65	120-180	40-60
Term newborn	40-60	90-170	52-92
1 month	30-50	110-180	60-104
6 months	25-35	110-180	65-125
1 year	20-30	80-160	70-118
2 years	20-30	80-130	73-117
4 years	20-30	80-120	65-117
6 years	18-24	75-115	76-116
8 years	18-22	70-110	76-119
10 years	16-20	70-110	82-122
12 years	16-20	60-110	84-128
14-17 years	16-20	60-105	85-136

B. Alarm Failure and Alarm-Related Incidents

1. When faulty equipment is identified, staff will:
 - i. Remove the faulty equipment from service
 - ii. Submit a work order to Biomed
 - iii. Place a copy of the work order and an “out of service” sign on the faulty equipment

C. Auditing and Monitoring

1. The hospital will monitor compliance with this policy as part of its Quality, Patient Safety, and Process Improvement plan. The Regulatory Compliance Committee will define what to monitor and add the items to the Joint Commission tracers as appropriate.

D. Education

1. Appropriate staff will be educated about the purpose and proper operation of alarm systems for which they are responsible during their orientation or as needed when new equipment or new processes are put in service.

REFERENCES:

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