

OUR LADY OF THE LAKE REGIONAL MEDICAL CENTER			
<b>Policy Manual:</b>	<b>Pharmacy</b>	<b>Section:</b>	Medication Use, Storage and Handling
<b>Title:</b>	<b>Medications: High Alert Medications</b>	<b>Policy Reference #:</b>	PH-03-10
		<b>Supersedes #:</b>	PH-09-07-O
<b>Date of Origination:</b>	05/23/2006	<b>Last Date Reviewed:</b>	05/07/2018
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**PURPOSE:**

To provide standardized medication safety policies for “high alert” medications identified by the Institute for Safe Medication Practices (ISMP).

**DEFINITIONS:**

**High-alert Medications:** medications that bear a higher risk of causing significant patient harm when they are used in error.

**Independent double check:** for the purpose of this policy, shall mean two authorized personnel separately checking the accuracy of the following against the prescriber’s order.

**Authorized prescriber:** for the purpose of this policy, “authorized prescriber” shall mean a practitioner recognized by Our Lady of the Lake Regional Medical Center (OLOL) as having privileges to write orders for medications within the scope of their clinical practice at OLOL.

**POLICY:**

- I. The following medications are considered high-alert at Our Lady of the Lake Regional (OLOL)
  - a. Vasopressors
    - i. Epinephrine
    - ii. Norepinephrine
    - iii. Phenylephrine
  - b. Insulin (all formulations)
  - c. Alteplase (TPA; excludes catheter occlusion doses)
  - d. Opioids for Injection
    - i. Morphine
    - ii. Hydromorphone
    - iii. Fentanyl, sufentanil, remifentanil
    - iv. Meperidine
    - v. PCA
    - vi. Epidural
  - e. Intrathecal Medications
  - f. Concentrated electrolytes for injection
    - i. Potassium chloride, potassium phosphate, potassium acetate

- ii. Sodium Chloride solutions greater than 0.9%
    - g. Heparin Infusions
    - h. Neuromuscular blockers
    - i. Chemotherapy
      - i. Vinca alkaloids
        - 1. Vincristine
        - 2. Vinblastine
        - 3. Vinorelbine
      - ii. Methotrexate
- II. The following methods shall be utilized, where appropriate, to prevent harm associated with the use of high-alert medications:
  - a. Development of order sets, pre-printed order forms, and protocols to standardize prescribing and monitoring
  - b. Use of standardized concentrations, dosage strengths, and dosage forms to minimize variability
  - c. Use of pre-mixed and unit-dosed medications whenever possible
  - d. Use of programmable pumps to administer continuous and intermittent infusions
  - e. Perform independent double checks after preparation and prior to administration
    - i. Factors to be identified during the independent double check include
      - 1. Right patient identification per our organization's policy
      - 2. Right drug (verified against current order)
      - 3. Right dose of drug (verified against current order and infusion pump; mathematical calculations, strength or concentration must be verified)
      - 4. Right route of administration
      - 5. Right time of administration
  - f. Limit prescriptive authority to authorized providers
  - g. Restrict the use of certain high-alert medications to appropriate nursing units
  - h. Utilize barcode scans during the following medication processes (where automation present): IV compounding, repackaging, dispensing, loading and refilling in automated dispensing cabinets
  - i. All defined "High Alert" medications should be labeled as such in the Electronic Medical Record in a face up manner to the administering provider
  - j. Limit the storage of high-alert medications to areas with personnel trained for safe preparation and administration.
- III. Readers of this policy should refer to the attached High Alert Restrictions reference document for detailed instructions for handling high-alert medications.

## REFERENCES

MM 01.01.03 EP1. The hospital identifies, in writing, its high-alert and hazardous medications. *The Joint Commission Medication Management Standards*

MM 01.01.03 EP2. The hospital has a process for managing high-alert and hazardous medications. *The Joint Commission Medication Management Standards*

MM 01.01.03 EP3. The hospital implements its process for managing high-alert and hazardous medications. *The Joint Commission Medication Management Standards* Meds Requiring Special Monitoring Protocol

**High Alert Medications Restrictions**  
(see *High-Alert Policy PH-03-10*)

Drug(s)	Actions to Prevent Errors											Comments	
	Prescribing				Preparation & Dispensing				Administration & Monitoring				Storage
	Authorized prescriber	Order Set	Protocol	Standard Dose	Standard Conc.	Standard Form	Double-check	Labeling	Infusion Pump	Double-check	Patient care unit restrictions		Unit storage restriction
Alteplase (excludes catheter occlusion doses)					•				•		•	•	a. Standard concentrations: see protocol b. Restricted patient care units: see <i>Meds Requiring Special Monitoring</i> document c. Orders require starting dose and continuous rate d. Infusion pump must be used to administer
Epinephrine, Norepinephrine Phenylephrine infusions					•				•		•	•	a. Standard concentrations: see protocol b. Restricted patient care units: see <i>Meds Requiring Special Monitoring</i> document, exception: code blue situations c. Orders require starting dose and titration d. Infusion pump must be used to administer e. Storage Bins will be marked as "high alert"
Insulin infusion			•		•				•	•	•		a. Insulin infusion titration protocol must be utilized b. Restricted patient care units: see <i>Meds Requiring Special Monitoring</i> document c. Standard concentration: see protocol d. Infusion pump required for administration e. Independent double check: two nurses prior to administration
Insulin pens										•		•	a. Insulin vials will be stored separately. Different formulations will be stored separately. b. Storage bins will be "red" and marked as "high alert" c. Independent double check: two nurses prior to administration d. Bedside Barcode Scanning (BCMA) should be used to verify product
Methotrexate (all formulations)				•			•	•	•	•		•	a. Standard dose: oral methotrexate will default to weekly dosage. b. Independent double check: two pharmacists before dispensing IV



phosphate, acetate)													
Heparin injections and infusions (exclude flushes/locks)				•	•				•				<ul style="list-style-type: none"> <li>a. Heparin infusion titration protocol must be utilized</li> <li>b. Infusion pump required for continuous or intermittent infusion</li> <li>c. Storage restrictions: main and satellite pharmacies</li> <li>d. Standard concentrations: see protocol</li> <li>e. Storage Bins will be marked as "high alert"</li> </ul>
Sodium chloride >0.9% inj. (e.g. 3%, 14.6%, 23.4%)					•				•	•			<ul style="list-style-type: none"> <li>a. Storage restrictions: main and satellite pharmacies, TNCC, SICU</li> <li>b. Infusion pump required for continuous or intermittent infusion</li> <li>c. Two nurses must verify order and pump settings</li> <li>d. Storage Bins will be marked as "high alert"</li> </ul>
Vincristine, vinblastine, vinorelbine (non-intrathecal)	•					•	•	•	•	•		•	<ul style="list-style-type: none"> <li>a. Authorized prescriber: medical oncologist</li> <li>b. Independent double check by two pharmacists before dispensing</li> <li>c. Vincristine and vinblastine can only be prepared and dispensed in a minimum volume of 10mL of 0.9% sodium chloride injection delivered in a minimum of a 20mL syringe or minibag</li> <li>d. Vinorelbine can only be prepared and dispensed in a minimum volume of 20mL of 0.9% sodium chloride delivered in a minimum of a 30mL syringe or minibag</li> <li>e. Syringe or minibag labeled "Fatal if given by other routes", "For IV Use Only"</li> <li>f. Syringes must also be packaged in overwrap, labeled "Do not remove covering until moment of injection."</li> <li>g. Independent double check by two registered nurses prior to administration</li> <li>h. Use of an infusion pump for administration</li> <li>i. Storage restrictions: main and satellite pharmacies</li> </ul>