



COHESIVE HEALTHCARE MANAGEMENT & CONSULTING

Mangum Regional Medical Center

TITLE		POLICY	
Heparin Standard Dose Protocol – DVT and PE		DRP-008	
MANUAL	EFFECTIVE DATE	REVIEW DATE	
Drug Room	10-1-2020	10-1-2020	
DEPARTMENT	REFERENCE		
Drug Room	<a href="https://www.accessdata.fda.gov">https://www.accessdata.fda.gov</a>		

**SCOPE**

This policy applies to all adult patients receiving care and treatment at Mangum Regional Medical Center.

**PURPOSE**

This protocol applies to adult patients at Mangum Regional Medical Center receiving standard dose heparin intravenous (IV) therapy for Deep Venous Embolism and Pulmonary Embolism.

**PROCEDURE**

Nursing Orders:

- Weigh patient STAT. Actual body weight \_\_\_\_\_ kg
- Determine Ideal Body Weight \_\_\_\_\_ kg
- Determine Adjusted Body weight \_\_\_\_\_ kg
- Determine Heparin Dosing Weight (DW): **dose using adjusted body weight if Actual Body Weight/Ideal Body Weight is greater than 1.2. DW \_\_\_\_\_ kg**

Medications:

- Heparin Sodium \_\_\_\_\_ units IV bolus STAT (\_\_\_\_\_ mL of 10,000 unit/mL vial). Bolus dose based on Heparin 80 units/kg x Dosing Weight (**Maximum of 5,000 units**)
- Heparin Sodium 25,000 units in D5W 500mL (50 units/mL) at \_\_\_\_\_ units/hour (\_\_\_\_\_ mL/hour) begin now. Heparin 18 units/kg-hour x DW (**Maximum of 1,000 units/hr initially**) \_\_\_\_\_ units/hour

Labs:

- CBC – STAT
- CMP -STAT
- PT/INR - STAT
- PTT - STAT
- PTT every 6 hours after initiation and after every Heparin rate change
- Daily weight while on Heparin Drip
- CBC every other day while on Heparin Drip
- GUAIAAC stool as needed

Monitoring Parameters:

- Draw blood for PTT from arm that doesn't have heparin infusion. Do not draw from heparin-flushed lines.
- If there is no other access other than the heparin line, then **stop** the heparin, flush the line, aspirate 10 mL of blood to waste, and then re-flush the line prior to drawing a specimen.
- Do not interrupt heparin infusion unless ordered.
- Contact medical provider if platelet count is less than 150,000 microliter or a 50% drop from baseline; hematoma, bleeding or suspected bleeding occurs.

**REFERENCES**

[https://www.accessdata.fda.gov/drugsatfda\\_docs/label/2017/017029s140lbl.pdf](https://www.accessdata.fda.gov/drugsatfda_docs/label/2017/017029s140lbl.pdf)

**ATTACHMENTS**

None.

**REVISIONS/UPDATES**

Date	Brief Description of Revision/Change