

PATIENT INFORMATION

Weight	kg
	Weight

Allergies

	Blood Component (Cryoprecipitate) Order Set	м	к	(
rders rocessed	Criteria						
ate	All adult inpatients and emergency adult patients			1			
d/mm/yyyy)	Note: All STAT orders are to be called to your site Transfusion Medicine Laboratory			I			
	Not Recommended For			I			
	Massively bleeding or unstable bleeding patients						
me (hhmm)	Note: This would ONLY be patients being transfused for resuscitation due to critical bleeding or the			I			
	Massive Transfusion Protocol was activated						
	Operating Room or Recovery Room patients						
Pre-Transfusion Patient History							
	Admitting diagnosis			1			
tatus	☑ Allergies/Sensitivities related to Transfusion						
alus	Patient consent completed						
	Vitals/Monitoring	11					
ocessing	Vitals			I			
eviewed by	I Height and Weight on admission to be documented in Meditech						
	Temp, HR, RR, BP, SpO ₂ – Pre-Transfusion, at 15 minutes, and Post-Transfusion						
	(Document along with time on Transfusion Product Issue/Nursing Documentation Form)						
atus	IV Fluids	<u> </u>					
	IV Fluid			I			
	□ sodium chloride 0.9% at mL/h						
axed by	Other IV Orders						
	□ Saline Lock						
	Pre-Transfusion Medications						
	\square diphenhydr AMINE mg \square PO OR \square IV x 1 prior to transfusion if history of allergic reactions			I			
	\square furosemide mg \square PO OR \square IV x 1 prior to transfusion			I			
	Note: Consider furosemide in patients at risk for transfusion associated circulatory overload. It is						
	preferable to give furosemide before the transfusion if the patient is not hypovolemic and is						
	hemodynamically stable						
	Pre-Transfusion Lab Investigations	11					
	Group and Screen (GPS) if required (Once per admission)						
Telephone	Order			-			
	Ordering Practitioner, Designation Signature Date/Time (dd/mm/yyy	y hhr	nm)	-			
Telephone (Read	-	ŀ			
	Ordering Physician Date (dd/mm/yyyy) Time (hhmm)						
			1 0	1			



PATIENT INFORMATION

_ cm	Weight	kg
	_ cm	_cm Weight

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	Blood Component (Cryop	precipitate) C	Order Set	м	К	o
Orders Processed	Cryoprecipitate Transfusion					
Date (dd/mm/yyyy)	Pre-Transfusion Lab Investigation					
<u> </u>	Indications and Dosing for Transf	usion of Cryo	precipitate			
Fime (hhmm)	Clinical Setting					
	Diagnosis/Indication	Fibrinogen g/L	Recommendation and Dose			
	Microvascular bleeding	Less than 1	1 dose (8 – 12 units)			
3y Status	Uremic bleeding	Any	For patients that do not respond to desmopressin or desmopressin is contraindicated 1 dose (10 units)			
Processing Reviewed by Status	 1 dose increases the fibrinogen by 0.5 g/L Pre-transfusion fibrinogen g/L (N Does the patient have microvascular bleeding No Yes Does the patient have uremic bleeding? No Yes 	lot applicable for ure	emia)			
axed by	Administration Patient weight (kg): ☑ Transfuse units, pool over (e.g. 1 dose over 10 – 30 minutes per dose m Medicine Laboratory)	naximum 4 hours fro	m issue time from the Transfusion			
	Post-Transfusion Lab Investigatio ⊠ FIB after dose to reassess the need for add		emia excluded)			
] Telephone	Order			1		L
f Telephone (Ordering Practitioner, Designation	Signature	Date/Time (dd/mm/yy	yy hhr Read		
	Ordering Physician Date (dd/r	nm/yyyy)	Time (hhmm)	-	-	
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	Blood Compone	ent (Cryop	orecipitate) C	Order Set	м	ĸ	c
Orders Processed Date	Cryoprecipitate Transfu	ision Conti	nued				
(dd/mm/yyyy)	Further Lab Investigatio	ons					
Time (hhmm)	Additional Orders						
Ву							
Status							
Processing Reviewed by							
Status							
Faxed by							
Telephone	Order						
	Ordering Practitioner, Desig	gnation	Signature	Date/Tir	me (dd/mm/yyyy h	hmm))