Massive Transfusion

Judy Mikhail RN, MSN, MBA Program Manager Michigan Trauma Quality Improvement Program (MTQIP) University of Michigan

Definition(s)

Classic

Complete replacement of a patient's blood volume within 24 hours (about 10 units/ 70kg adult)
Greater than 50 units within the first 24 hours
Greater than 50 units within the first 48 hours
Comparisons made difficult

"Reality"

- Typical patient initially bleeds without any replacement at scene
- Receives crystalloid en route
- Only then receives blood in ED or OR
- How does one determine when transfusion becomes massive?



Reality

 Hemorrhage models show that 70% of estimated blood volume is lost even before the 1st unit is given



New Definition Required

Discrepancy between bleeding and replacement
Behind the eight ball
Proactive anticipatory "working" definition:

Administration of 4 units of blood within first hour, with anticipation of ongoing usage?
Administration of 4 units of blood within first 4 hours?

Massive Transfusion Protocol Development:

All players at the table

- Surgeons
 Emergency Medicine
 Anesthesia
 Lab
 Blood Bank
- Nursing
 Trauma Program Manager
 IS
 Medical Records
 Admitting

MTP: Logistics

Hospital Specific (Resource Dependent)

- Proximity of Blood Bank to ED/OR
- Point of care testing
- Sophisticated Tube system
- Blood fridge in ED/OR
- Runner service
- Lab personnel on trauma activations
- Transfusion specialists 24/7
 Pathology Residents
- None of the above



MTP: Process Issues

Primary Goal:

- Timely, coordinated, access to adequate blood supply
- Secondary Goals:
 - Standardization
 - Avoid surgeon distraction from operative field
 - Accountability built into the process
 - Minimize product wastage

Massive Transfusion Activation



Formal Activation vs "The Slide..."

- Overhead announcement of Massive Transfusion
- Beeper
- 2 way radio
- QI Tip: Incorporate activation time into ED flowsheet & anesthesia record

Who Can Activate?

- Surgeon
- Anesthesia
- Emergency Medicine?
- PA / NP / Nurse?
- Also responsible for timely <u>deactivation</u>

Indications for Activation

- Class IV Shock
- Class III Shock with anticipated blood loss requiring 10 unit replacement and likelihood of continued hemorrhage
- Or 4 units within one hour & more anticipated

 The actual loss of blood does not have to occur before the judgment is made that such loss is imminent
 Never based on laboratory values!

Patient Identification CAUTION: Mislabel risk high in multiple trauma Sophisticated John Doe system ■ SPEED and SAFTEY Blood Draws Ideal if Lab personnel respond to activations ■ Nurse drawn & labeled ■ (prelabeled tubes John Doe packet) Adequate non hemolyzed specimen Tip: May obtain from externally dripping wound as last resort

Emergency Uncrossmatched (immediate)

- Ideally in ED before patient
- O Positive
 - For males
 - 36% donor pool
- O Negative
 - For childbearing females & children
 - Only 8% donor pool



QI: Delay to Specimen

✓ If more than 4 U of Emergency O blood given before specimen obtained ✓ Leads to subsequent difficulty in compatibility testing ✓ Stay with type O blood ✓ Review cause of delay



Blood Progression

- 1. Emergency Uncrossmatched (immediate)
- **2. Type Specific (10 mins)**ABO and Rh compatible
- 3. Crossmatched (40 mins)
 ABO and Rh type
 Other known antibodies



Specimen transportation
Lab Personnel
Tube system (caution)
Runner (baton pass off)
Eye to eye
Differentiate this specimen from others



Blood Containers ED/OR Blood Refrigerator (stationary) Ice Chest (advantage of being portable) Typical Massive Transfusion Pack: ■ 6-10 units PRBC's ■ 4-6 units FFP ■ 1 pheresis Platelets unit (6-8 random)

Massive Transfusion Pack

Product	# Units	Container	Time out	Re-Issue
		Temp	Of BB	Ability
PRBC	6-10	Ice Chest	3-8 hours	Yes
FFP	4-6	Ice Chest	3-8 hours	Yes
Platelets	1 pheresis	Room	3 hours	Yes
		Temp		

FFP or Thawed Plasma

Policy:

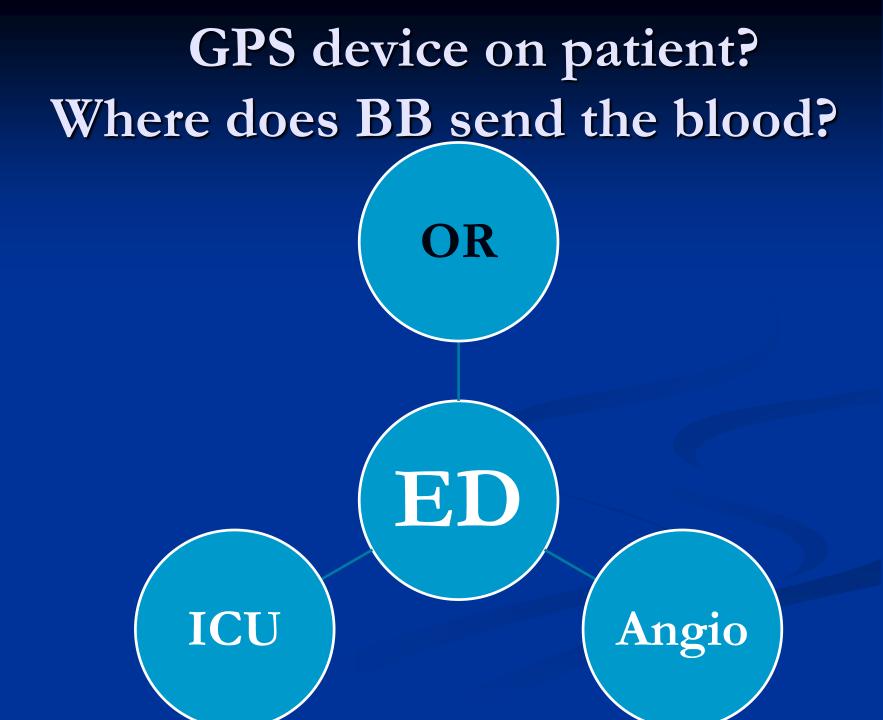
Thaw time variable (equipment dependent)
Typical: 2 units in 20 minutes

 TIP: Most Blood Banks keep 4-6 pre-thawed plasma at all times

Re-cycle daily

■ Monitor for wastage

Consider sending it with initial chest of blood



Blood Bank Accountability

- Mobilize immediate staffing (cross train)
 Stay 6-10 U ahead at all times
- Continuous communication with anesthesia/surgeon
 Monitor and maintain inventory



BB Inventory Maintenance

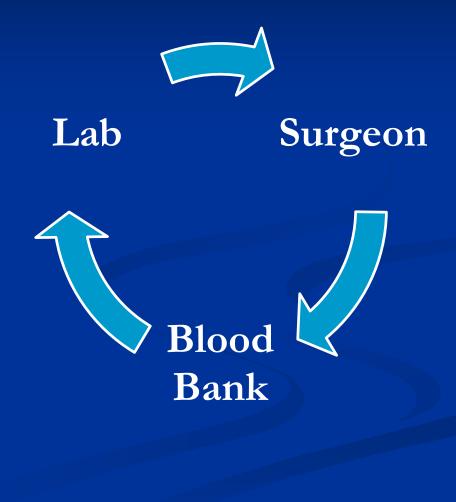
- Automatic switch of patients to ABO compatible blood as blood supply dwindles:
- Male Patient A Neg O+, A+, then A-, then O-
- Blood Bankers always think about the current patient as well as the next one coming in



Communication Logistics

- BB/Lab direct phone contact to Surgeon/Anesth
- Intercom
- 2 Way Radio
- Red "Hot Line" Phone





- Suggested Coagulation Tests
 PT-INR
 PTT
 Platelets
 - FibrinogenIonized Calcium



■ Suggested Frequency: q 30 – 60 min



Do you remember when <u>whole blood</u> was given?
Phased out in 1970's to component therapy
Initial additives: blood out date was 21 days
Then extended to 35 days (late 1970's)
Then extended to 42 days (1980's)

Blood Additives Affect on Coagulopathy

CPDA1

- 35 day out date
- > Hct 75%
 - No AdeninePreferred in Neonates

 Most hospitals have a mix of blood additives due to distribution issues

- Adsol AS1
- Nutricel-AS3
- Optisol AS5
 - 42 day outdate
 - > Hct 60%
 - 50cc plasma
 - 100cc additive saline

Farringer J of Trauma 1993

Historically 3 MTP Management Strategies



1st Strategy

Transfuse coagulation components according to a formula based upon # units RBC transfused
✓ Prophylactic administration = not supported by research





2nd Strategy

 Defer component transfusion until microvascular
 bleeding is identified in the wound





^{3rd} Strategy

Directed replacement protocol based on coagulation monitoring





Reality...

Challenge is how to operationalize
How to administer products in a timely manner?
With pending labs results?
Order labs and components simultaneously
When lab results known product available

Directed Component Replacement

FFP

- PT or aPTT > 1.5 times control
- INR <u>alone</u> not recommended to base MTP decisions
 - INR elevations are not indicative of the risk of bleeding unless they are > 1.8 and are associated with an elevated PTT
- Reality:
 - If PT-INR/PTT pending with ongoing blood loss
 - Give 2 units FFP for every 5-6 units of blood given

Directed Component Replacement

Platelets

Platelet count less than 50,000 with ongoing bleeding

 1 pheresis platelet unit = 6-8 random donor units

 Cryoprecipitate

 Fibrinogen less then 80-100 mg/dl
 Give 10 u Cryo

Conclusion

- Transfusion triggers for RBC and Platelets have been abandoned
- RBC administration =
 - physiologic instability and on going blood loss
- Platelets administration =
 - thrombocytopenia with on going blood loss
- **FFP**
 - Relationship to # units given
 - Replacement ratios vary among trauma centers

Ideal

RBC to Plasma Ratio?

Majority of studies show survival advantage
1:1 or 1:2

No Prospective randomized controlled trials



EAST Guidelines RBC Transfusion in Adult Trauma and Critical Care 2009

Level I

RBC transfusion is indicated for patients with:
 Evidence of shock
 Acute hemorrhage and hemodynamic stability
 Postrictive strategy is as effective as liberal

Restrictive strategy is as effective as liberal

SPEED is Important

1. PT prolongation

- Occurs early!
- If FFP delayed >3u RBC PT crosses hemostatic threshold regardless of amt of FFP subsequently given

2. Fibrinogen

Fibrinogen depletion easier to correct Suggested Formula
FFP/PRBC ratio
1:1 or 1:2
Give 2 u FFP with first units of PRBC to start

Fibrinogen depletion is prevented when the FFP/PRBC ratio of 4:5 was used

Model Recommendations

3. Platelets

- Even if replacement of platelets is delayed until10 units of PRBC given
- Critical platelet dilution is prevented
- Reinforces: prophylactic administration of PLT is unnecessary

Suggested Formula
PLT: Blood Ratio
8:10

NOT A PERMANENT PART OF THE CHART Massive Transfusion Tracking Sheet

Automatic Numerical Order of Transfusions

Suggested Sheet Use: Follow the numbers and cross off units as you give them

Chest Shipment		RBC's	Thawed Plasma	PLT's 1 Jumbo Apheresis unit = 6 (old style)	CRYO 10 units 1 pooled unit = 5 single unit's	rFVIIa Dose 6 mg dose for > 100 kg 5mg dose for 65-100 kg (Pediatric dosing per physician)	
-	Chest 1	1	2				
l ü		3	4				
Class I Trauma		5	6				
ra	Chest 2	7	8				
OE		9	10				
		11	12				
	Chest 3	14	15	13			
		16	17				
		18	19				
	Chest 4	23	24		20, 21	22	
		25	26			From Pharmacy	
D		27	28				
0	Chest 5	30	31	29			
S		32	33				
.2		34	35				
Massive Transfusion	Chest 6	39	40		36, 37	38 From Pharmacy	
ä		41	42				
-		43	44				
	Chest 7	46	47	45			
Ve		48	49				
		50	51				
22	Chest 8	55	56		52, 53	Ask for order 54	
3		57	58				
Z		59	60				
	Chest 9	62	63	61			
		64	65				
	<u></u>	66	67				
	Chest 10	71	72		68, 69	Ask for order 70	
		73	74				
		75	76				
			Continue	as necessary			

**NOT A PERMANENT PART OF THE CHART*

Is There A Limit to Massive Transfusion After Trauma?



There is no threshold for stopping blood in a massive transfusion

Duration and severity of shock are predictors of mortality not volume of blood given

Blood Costs (MI \$)

Product	Packaged	Cost		
PRBC	1 unit	\$850		
FFP	1 unit	\$250		
Platelets	1 random unit	\$250		
	1 pheresis unit	\$1000		
Cryo	1 unit	\$100		
	10 units pooled	\$900		
Example	22 u PRBC	One Patient		
MTP	16 u FFP	\$25,800		
	3 ph PLTs	Total		
	1 Cryo	Wow!		

Complications of Massive Transfusion Theoretical vs Genuine?

Depletion 2,3 DPG (*Theoretical*)

Decreased O2 off load

■ Hypocalcemia (*Genuine*)

- Significant citrate load
- Monitor ionized calcium
- Hyperkalemia (Theoretical)
 - Storage lesion

- Coagulopathy (*Genuine*) Multifactorial:
 - Coagulation abnormalities
 - Excessive fibrinolysis
 - Hypothermia, Acidosis,
 - Dilutional coagulopathy
- Hypothermia (Genuine)
 - Rapid infusers 42 C
 - Hypothermia protocol

Emerging Effects

Trauma -Independent risk factor:

- Death
- Perioperative infection
- SIRS
- ICU admission
- Age of blood (MOF)
 - # u > 14 days
 - # u > 21 days



Emerging Effects of Blood

Detrimental Immunomodulation

- Increased risk for cancer recurrence
- Stimulation of cytokine release (> 14 days)

Enhanced acute inflammatory response



Do you know the age of blood transfused in your trauma center?

- Blood bankers always release "short date" (older blood) first on a first come first served basis
 Average age of RBC's transfused is 21 days
 40% > 28 days
- Huge ramifications on blood supply



Supportive Guidelines

- Pleural Autotransfusion
- Intra Operative Cell Saver
- Hypothermia control
- ✓ Damage control surgery
- ✓ Recombinant Factor VIIa
- ✓ Blood substitutes (future...)



MTP Case: ED Analysis

Class I Pedestrian vs car 3 min notice, EMS: no IV or ETT Open Book Pelvis, Liver Lac

Time	2048	2058	2108	2120	2130	2140	2150	2205	To OR
BP	51/33	68/33	57/38	89/38	105/53	105/55	78/56	78/50	
Р	133	122	126	124	131	131	116	118	
Tx's	ETT	Hgb 13.9 Unable to start IV Plain Films Subclavian Cordis Placed		pH 7.15 BD –16 To CT (BP 105/55) Angio Notified Did not activate MTP!				Total: 16 L 12 U	
IV			3L	3L	3L	2L	2L	1L	
Blood			4u	2u		2u	2u	2u	
			O+ 20min!	O+		O+	O+	O+	

GSW Chest

- Found down in snow bank
- Single GSW to Lt Chest
- EMS: load & go
- Total EMS time: 8 min
- 2 min prior notice to
 - Tr Center
- Surgeon present on arrival

- Signs of life
 ETT @ 2 min
 Chest opened @ 3 min
 Femoral Cordis 9F@ 7 min
 Blood specimen to lab @ 7 min
- 1st unit up on rapid infuser
 <u>8 MINs</u>

Cost Savings

T&S vs T&C in trauma program ■ T&S for majority of trauma ■ T&C reserved for those only at high risk: ■Class I activations Can we do even better? ■Are there predictors of blood use in trauma or is it just common sense? Physiologic and Mech of Injury

shifthappens

In Summary

- MTP serves as a barometer of Trauma Program Maturity
- Reflects the highest level of multidisciplinary collaboration working together in a timed event
- Challenge: to continuously update your protocols to reflect current literature
- Critique every massive transfusion as another opportunity for improvement!