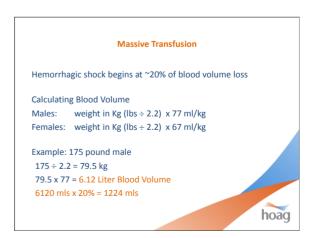




Massive Transfusion • Massive Transfusion (>10 Units in 24 hours) - 7% of military trauma patients - 3% of civilian trauma patients - 30 - 60% mortality rate • 40% of 14 million annual transfusions • Cause of Death in Trauma Patients - 40% Uncontrolled Hemorrhage/Exsanguination - Second only to CNS injury



Clinical Signs of Acute Hemorrhage							
Blood loss (ml)	0-750	750-1500	1500-2000	>2000			
% of total blood volume	0 -15%	15-30%	30-40%	>40%			
Pulse rate	< 100	> 100	> 120	> 140			
Blood pressure	Normal	Normal	↓	↓			
Pulse pressure	Normal / ↑	↓	↓	↓			
Orthostasis	Absent	Minimal	Marked	Marked			
Capillary Refill (perfusion)	Normal	Delayed	Delayed	Delayed			
Respiratory Rate	14 - 20	20 - 30	30 - 40	> 34			
Urine Output (ml/hr)	> 30	20 - 30	5 - 15	< 5			
CNS Mental Status	Slight Anxiety	Mild Anxiety	Anxious/ Confused	Confused/ Lethargic			
Cardiac Index L/min (△ %)	↓ 0-10%	↓ 20-50%	↓ 50-75%	↓ >75%			
				noag			

The "Lethal Triad of Trauma" • Hypothermia • Acidosis • Coagulopathy

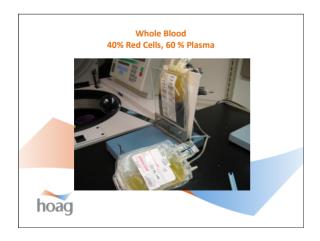
The Lethal Triad of Trauma Hypothermia

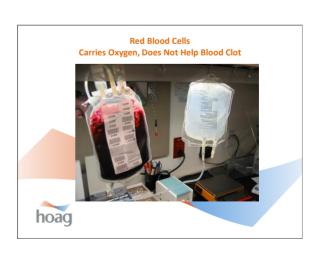
- Causes of Hypothermia
 - Environmental factors: extrication and transport time
 - IV fluids and ongoing blood loss
 - Alteration of normal heat producing metabolism
- Effects of Hypothermia
 - Decreases platelet aggregation and adhesion
 - Decrease coagulation factor activity by 10% for each degree decrease in core temperature.
 - Both R (Rx Time) & K (Fibrin) prolonged on TEG
 - 100% fatal when core temperature reaches < 32° €
- Coagulation assays are run at 37 ° C.!

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The Lethal Triad of Trauma Acidosis

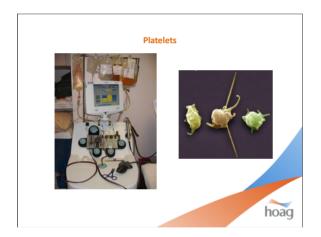
- Causes of Acidosis
 - Decreased perfusion leads to anaerobic metabolism and lactic acid production.
 - RL pH 6.0, normal saline 4.5, no buffering capacity
 - Red cells at two weeks have pH < 7.0
- Effects of Acidosis
 - Reduced clot formation demonstrated by TEG
 - Spherical platelets devoid of pseudopods
 - Reduced fibrinogen levels, platelet counts & Xa
- · Prevention of Acidosis
 - Dependent on restoration of perfusion
 - Exogenous bicarb has mixed results

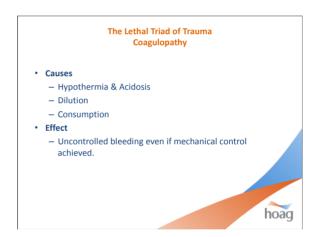


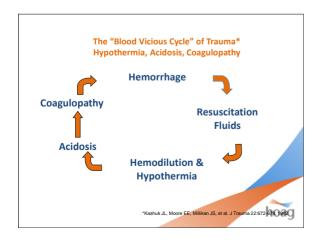


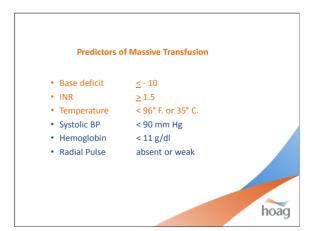










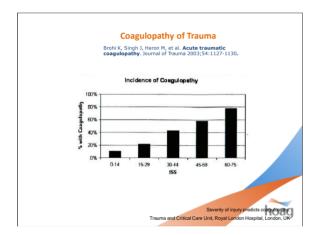


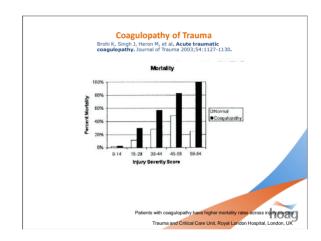
Traditional Treatment of Acute Hemorrhage ATLS Resuscitation Protocol

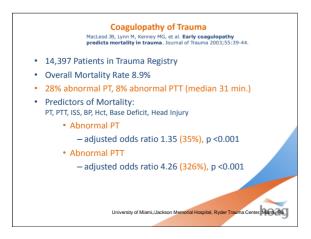
- · Insert two large bore IVs.
- · Crystalloids to support volume and blood pressure
 - ATLS: 2 L crystalloid if systolic BP <100
 - ACLS: 3 ml of crystalloid/1 ml of blood loss.
- · Red cells as an oxygen carrier
 - If systolic BP remains or falls back to <100
 - If bleeding > 100 ml/min
- Platelets, FFP and Cryo if coag tests abnormal
 - INR > 1.5
 - Platelets < 50 K
 - Fibrinogen < 100 mg/dl

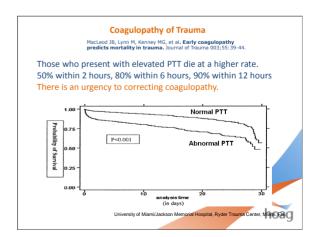
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Coagulopathy of Trauma Brohi K, Singh J, Heron M, et al. Acute traumatic coagulopathy. Journal of Trauma 2003;54:1127-1130. 1088 consecutive trauma patients 24% had a significant coagulopathy on admission — PT >18, PTT >60 More severely injured you are, the worse the coagulopathy. Mortality rate higher in those with coagulopathy across range of injury severity







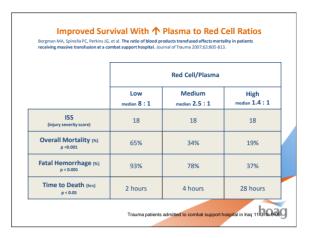


2005 US Army Institute of Surgical Research International Symposium on Massive Transfusion

Holcomb JB, Hess JR: Early massive trauma transfusion: State of the art. J Trauma 60:S1-S2, 2006

- · 2005: International Consensus Conference
 - Sponsored by US Army Institute of Surgical Research
 - 46 experts from US and Europe
- Conclusions
 - Transfusion practices and survival rates vary.
 - Increased plasma and platelet to red cells ratios associated with better survival.
 - Guidelines should aim for 1:1:1 ratio.

US Army Institute of Surgical Research, Ft. Sam Houston, San Antonio, Tx.

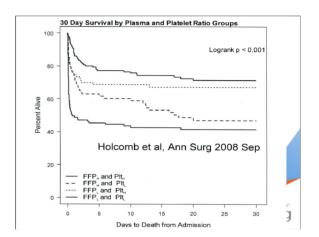


Improved Survival With ↑ Platelet to Red Cell Ratios

Holcomb JG, Wade CE, Michalec JE, et al. **Increased plasma and platelet to red cell transfusion rations improves outcome in 466 massively transfused civilian trauma patients**. Annals of Surger 2008;248:447-458,

- 16 US Level I trauma centers
- 466 of 1574 (30%) patients massively transfused
- Divided into four groups:
 - High Plasma (red cell/plasma <2:1)
 - High (<2:1) & Low (>2:1) Platelet
 - Low Plasma (red cell/plasma > 2:1)
 - High (<2:1) & Low (>2:1) Platelet

16 US Level I Trauma Centers,



Decreased Red Cell:Plasma Ratios Associated With Improved Mortality Rates in Trauma Resuscitation Patients

Nine Studies Oct 2007 – Feb 2009					
study	n	outcome			
Borgman et al 2007 J Trauma 63:805-807	246	r/p 8:1 mortality 65% r/p 2.5:1 mortality 34% r/p 1.4:1 mortality 19%			
Duchesne et al 2008 J Trauma 65:272-276	385	r/p > 1:1 mortality 88% r/p ≤ 1:1 mortality 26%			
Margele et al 2008 Vox Sang 95:112-119	713	r/p > 1.1 mortality 24.6% r/p 0.9-1.1 mortality 9.6% r/p < 0.9 mortality 3.5%			
Holcomb et al 2008 Ann Surg 248:447-458	466	r/p > 1:2 mortality 60% $r/p \le 1:2$ mortality 40%			
Kashuk et al 2008 J Trauma 65:986-993	133	r/p 4:1 in non-survivors r/p 2:1 in survivors			
Sperry et al 2008 J Trauma 65:986-993	415	r/p > 1.5:1 mortality 12.8% r/p < 1.5:1 mortality 3.9%			
Snyder et al 2008 J Trauma 66:358-362	134	r/p > 2:1 mortality 58% r/p < 2:1 mortality 40%			
Gunter et al 2007 J Trauma 65:527-534	213	r/p > 3:2 mortality 62% r/p ≤ 3:2 mortality 41%			
Johansson 2009 Vox Sang 96:111-118	832	no protocol mortality 31.5% 5:5:2 r/p/plt mortality 20.4%			

The Problem of Survivor Bias

- 50% of MTP patients die within 24 hours
- 25% die within first 4 hours, many within one hour.
- 1:1 red cell/plasma only applies to ~ 5% of patients.
- Patients that died before receiving plasma counted in non-survivor groups.
- "Does the plasma save the life or does plasma transfusion happen to those who live?"
 Jeannie Calum, MD, Toronto, AABB Annual Meeting 2008, Montreal, Canada

The PROMMT Study

Holcomb JB, del Junco D, Fox E, et al. The prospective, observational, multicenter major trauma transfusion (PROMMT) study: comparative effectiveness of a time-varying treatme with competing risks. Arch Surg 2012 Oct 15:1-10. doi: 10.10.1001/2013.jamasurg.387. [Epub ahead of print]

- · Prospective, multicenter observational trial
- 10 Level I trauma centers, 905 patients
- · Goal of the study design was to eliminate survivor bias
 - Real time data collection from time of admission
 - Not limited to massive transfusion patients
 - Ratios computed at 14 consecutive time intervals.
 - Data analyzed using a time-dependent proportional hazard regression analysis.



The PROMMTT Study

Holcomb JB, del Junco D, Fox E, et al. The prospective, observational, multicenter major trauma transfusion (PROMMTT) study: comparative effectiveness of a time-varying treatment with comprisks. Arch Surg 2012 Oct 15:1-10. doi: 10.10.1001/2013.jamasurg.387. [Epub ahead of print]

- Hemorrhagic Cause of Death:
 - 60% within 3 hours, 94% occur within 24 hours
 - 81% of patients that died within 6 hours bled to death.
- Red Cell/Plasma & Platelet/Red Cell ratios >1:2
 - 3-4 times less likely to die in the first 6 hours
 - Benefit not seen after 24 hours
 - · cause of death shifts to head injury, respiratory distress, organ failure and infection

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Are MTPs Effective in Non-Trauma Cases?

Mell MW, O'Neil AS, Callcut RA, et al. Effect of early plasma transfusion on mortality in pati abdominal aortic aneurysm. Surgery 2010 Apr 6, Epub, in press

- · Non-elective ruptured AAA repair
 - 128 patients received >10 units during OR
 - 30 day mortality 22.6%, 11 intra op deaths
 - -2 groups: p/r > 1:2 and p/r <1:2
- · High plasma group
 - 30 day mortality 15% vs 39%
 - Colon ischemia 22.4% vs. 41.1%

Division of Vascular Surgery, Stanford University

The Changing Resuscitation Paradigm "Damage Control Resuscitation

- Goal: Prevention of the "lethal triad" of acidosis, hypothermia and coagulopathy.
 - Tolerance of moderate hypotension (~90 systolic) and minimal crystalloid use.
 - Delay surgery if possible until hypothermia, acidosis and coagulopathy are treated.
 - Short surgical procedures to control bleeding and minimize contamination.
 - Give plasma, platelets and cryoprecipitate earlier and in increased amounts.
 - Best achieved with a massive transfusion protoc

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Growth of Massive Transfusion Protocols

- · 2006: 3 academic trauma centers in the US
- 2010: 85% of 186 trauma centers

 - Most begin with 1:1:1 ratio - All include plasma by second delivery
 - 37% include Factor VIIa as part of their protocol

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California Maternal Quality Care Collaborative (CMQCC)

- UCLA Maternal Quality Indicator Group evolved into the California Perinatal Quality Care Collaborative (CPQCC)
- · 2004 CDPH and CPQCC formed the CMQCC
- Mission End preventable morbidity and mortality and racial disparities in Califronia maternity care by sharing data, facilitating collaborations and defining clinical best preactices realted to obstetrical care.
- 2009 Hemmorhage Task Force practice guidelines

2009 CMQCC Hemorrhage Task Force Practice Guidelines

- · Maternal deaths in California on the increase
 - 6 per 100,000 in 1996
 - 16 per 100,000 in 2006 (54 in African Americans)
- In US, transfusions in OB patients have increased 92% between 1998 and 2005.
- In California, 2% of all deliveries are complicated by hemorrhage.
- Obstetric hemorrhage is the leading cause of maternal death

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2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Categorized into 1 of 4 stages with actions defined for:
 - Patient Assessment
 - Medication
 - Procedures
 - Transfusion Support
- Detailed protocols, slide presentations, charts available at http://www.cmqcc.org

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2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 0 Assess for risk factors for hemorrhage
 - Low Risk: Hold Clot
 - · No previous uterine incision
 - Singleton pregnancy
 - ≤4 previous births
 - · No known bleeding disorder
 - No history of PPH

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2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 0 Assess for risk factors for hemorrhage
 - Medium Risk: Type & Screen
 - Prior C-Section or uterine surgery
 - Multiple gestation
 - ≥4 previous vaginal births
 - Chorioamnionitis
 - History of previous PPH
 - Larger uterine fibroids
 - Estimate fetal weight > 4 Kg
 - Morbid obesitiy (BMI >35)

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2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 0 Assess for risk factors for hemorrhage
 - High Risk: Type & Crossmatch
 - Placenta previa, low lying placenta
 - Suspected placenta accreta of percreta
 - Hematocrit < 30 and other risk factors
 - Platelets < 100,000
 - Active bleeding (greater than show) on admission
 - Known coagulopathy

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2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 1
 - Blood loss >500 ml (vaginal) or 1000 ml C-section
 - Vital sign changes
 - HR > 110
 - BP < 85/45
 - O2 Sat < 95%
 - Blood Bank Recommendations
 - Ensure Type & Cross for 2 units

2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 2
 - Continued bleeding, total blood loss under 1500 ml
 - Blood Bank Recommendations
 - · Deliver 2 units red cells to bedside
 - Transfuse per clinical signs, do not wait for labs
 - Consider thawing 2 units FFP
 - Give FFP if thawing > 2 units red cells
 - Determine availability of additional red cells & "coag products"

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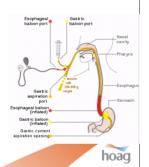
2009 CMQCC Hemorrhage Task Force Practice Guidelines

- Stage 3 (CODE RBC Activated)
 - Total blood loss over 1500 ml
 - > 2 units red cells given
 - Vital signs unstable or suspicion of DIC
 - Blood Bank
 - Massive "Hemorrhage Pack"
 - Near 1:1 red cell/plasma
 - 1 platelet
 - "unresponsive Coagulopathy" after 10 units red cells and "full coagulation factor replacement"
 - Consider Factor VIIa

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April 2008 Case Review

- 45 year old male
- Uncontrolled esophageal varices.
- Blakemore tube placement.
- Esophageal rupture during procedure.
- Received 37 blood components



Hoag Hospital

- · Community not-for-profit hospital
- Opened 1953, 75 beds
- Two campus system: 500 and 50 beds
- Specialties in Oncology, Heart & Vascular, Orthopedics, Neurosciences and Women's Health
- Annual Statistics
 - 6000 deliveries
 - 10,000 inpatient surgeries, 400 open heart
 - 70,000 ED visits
 - 25,000 blood components transfused





CODE RBC Team - August 2010

Arell Shapiro MD, Transfusion Medicine Greg Super MD, Director, ED Jennifer Keiner MD, Internal Medicine Pau Lee MD, Gl Lab Victor Beretta MD, Anesthesiology Grete Porteous MD, Anesthesiology Tamerou Asrat MD, Perinatology Rosemary O'Meeghan, MD, Critical Care Dale Braithwaite, MD, Obstetrics Stephanie Waldman, MD, Anesthesiology

Randy German, CLS, Transfusion Service Carol Vanderree, CLS, Transfusion Service

Sherry Lemasters, RN, Performance Imprvt Marilyn Lang, RN JD, Performance Imprvt Carlene Green, Performance Imprvt Tammy Valencia RN, ED
Molly Hewett RN, VP, Patient Care Svcs
Carole Metcalf RN, Director, Periop Svcs
Kelly Parra RN, Critical Care
Jamie Lynch, RN, Labor & Delivery
Kim Mikes RN, Director, Short Stay Unit
Kim Mullen RN, Exec Dir. Women's Health
Debbie Lepman, RN, Director, Critical Care
Debra Burgnski, RN, Nurse Educator, ICU

David Godoy, Support Services/Transport Michele VanRy, Supervisor, Comm/PBX

Heather Paradee, Respiratory Therapy

Stephanie Chao, Mgr, Pharmacy Dong Dao, Pharmacy Resident

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Focus Group Outcomes

- Poor (and excessive) communication
- Empirical and non-standardized physician orders
- Transport delays
- Laboratory testing turn around time too slow to guide therapy.
- No blood warmer/rapid infusion device available
- Inexperienced and/or insufficient staff at the bedside
- Excessive paperwork
- No defined roles or protocols in the Transfusion Service.
- Differing protocols being developed in different areas
- No guidelines for the use of Activated Factor VIIa



CODE RBC Goals

- Improve communications between the Nursing Unit and the Transfusion Service.
- Rapidly deploy equipment, blood & personnel to the bedside.
- Transfuse using a standardized ratio of blood components in accordance with the current Massive Transfusion literature.
- Prevent or minimize the "lethal triad of trauma"

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CODE RBC Goals

- Improve patient monitoring and treatment through a customized Code RBC order set.
- Meet or exceed the CMQCC Hemorrhage Task Force Recommendations
- · Universal protocol for all areas of the hospital.
- Develop guidelines for the use of Activated Factor VIIa.

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Medical Subgroup

- Determine Blood component transfusion ratios
- · Develop Order Sets
 - Nursing Care
 - Medications
 - Frequency and Type of Laboratory Monitoring

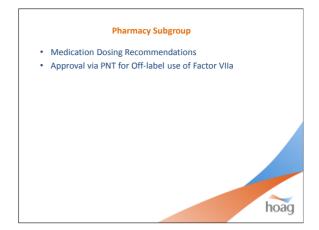
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Response Subgroup

- Assemble CODE RBC Kits
- Develop Hospital Policies
- CODE RBC Documentation Form
- Conduct Training and Education

Transfusion Service Subgroup

- · Select and validate coolers
- Develop multi-unit Transfusion Record
- Validate 5 day plasma
- · Define internal protocol and train staff
- · Obtain Belmont Rapid Infuser
- Develop computer workaround for Rh Negative patients
 - Switched to Rh Positive after Wave 3 (12 red cells)



Metrics Subgroup

- Development and Monitoring of Patient Metrics
- Ongoing Case Review and Quality Improvements

Ongoing case neview and Quanty improvements

CODE RBC Response

- Nursing Units
 - Call operator and announce "CODE RBC, Patient Location"
 - Call Blood Bank with
 - · medical record number
 - ordering MD
 - Order CODE RBC testing panels in HIS:
 - CODE RBC Blood Bank Panel
 - CODE RBC Diagnostic Panel
 - Retrieve CODE RBC Kit from Crash Cart

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CODE RBC Response

- Rapid Response Team to patient locations
- Respiratory to patient location
- Blood Bank prepares Wave One blood components.
- Transport to the Transfusion Service
 - Pick-up & deliver Wave One and Belmont Rapid Infuser.
 - Pick-up and deliver Wave Two
- Baseline labs drawn
 - Respirator runs gases on nearest POC instrument
 - Coagulation tests sent to Lab in green bag.
- MD completes paper order set

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Order Set: Blood Components

☑ Wave 1: 6 red cells in a cooler

Belmont Rapid Infuser

Wave 2: 1 platelet, 10 cryo

- Wave 3: 6 red cells, 6 plasma (cooler)
- Wave 4: 6 red cells, 6 plasma, 1 platelet, 10 cryo
- Wave 5: 6 red cells, 6 plasma, 1 platelet
- Wave 6: 6 red cells, 6 plasma, 1 platelet, 10 cryo

Continue alternating Wave 5 & 6

CODE RBC Kits Located on All Crash Carts

- · Code RBC Protocol Flowchart
- · Code RBC Order Set
- · Code RBC Documentation Form
- · Factor VIIa Order Form
- TEG Order Form
- Key Contact Numbers



Blood Draw Kits

- · Carried by RRT & on Belmont Infuser
- Green specimen bag
- 20 ml syringe
- · 21 bauge butterfly
- Blood transfer devices
- · Pre-filled 10 ml saline flush
- Blood Draw Tubes
 - 3 ml light green top
 - 10 ml lavender top
 - 3 ml lavender top









Order Set – Nursing Care

☑ Establish IV Access (Large Bore, Multiple lines if indicated)

- · Apply Bair Hugger PRN temp _____ degrees C.
- \boxtimes Strict Intake and Output, save all blood product and IV fluid bags.
- Pulse, respirations and blood pressure every 5 minutes.
- Temp every 30 minutes, core temp if possible.

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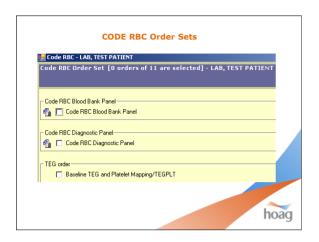
Order Set – Laboratory Monitoring

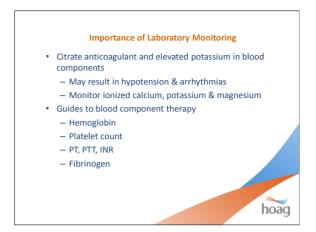
区ode RBC Blood Bank Panel

XM (10) Plasma (6) Plt (1) Cryo (10) Blood Issue Request

区ode RBC Diagnostic Panel

- ABGs with Lytes (Point of Care on GEM 4000)
 - ABGs, Hgb, Na, K, Cl, Ionized Ca, Glucose, Lactic Acid
- Code RBC Coag Panel
- Plt Count, PT/APTT, Fibrinogen, Mg
- TEG (consultation required)

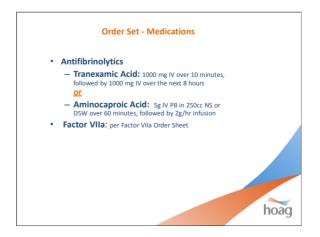




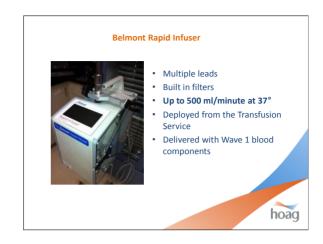














Risk of Uncrossmatched Red Cell Transfusions Frequency or Red Cell Alloimmunization · Risk related to pre-existing red cell alloantibodies No transfusions or pregnancies 0% Healthy Blood Donors 0.2% General patient population 1.0 - 1.5% Previous transfusions • 5 units 1.0% • 10 units 2.4% • 20 units 3.4% • 30 units 5.8% • 40 units 6.5 % - Previously Pregnancy - ? (lower red cell exposure) hoag

Risk of Uncrossmatched Red Cell Transfusions

Risk of hemolytic transfusion reactions following emergency-release rbc transfusion. Goodell P, UN L, Mohammed M, Powers American Journal of Clinical Pathology 2010, 134:202-206.

262 patients (265 episodes), 1002 red cell transfusions.

Clinically significant antibodies 17/265 (6.4%)

15 incompatible units to 7 patients 7/265 (2.6%)

1 delayed hemolytic reaction 1/1002 (0.1%/unit)

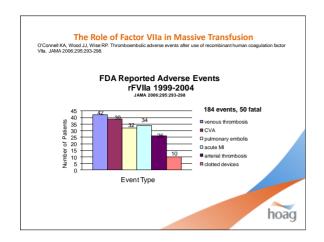
— Anti c, Jk(a), E in plasma and eluate

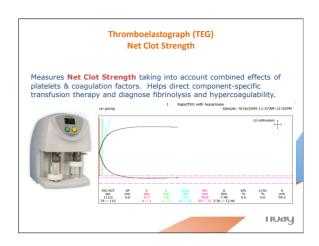
— 36 hours following transfusion

• LD 1057, T Billi 2.2, Haptoglobin < 20

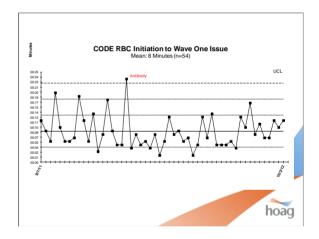
— No clinical sequelae

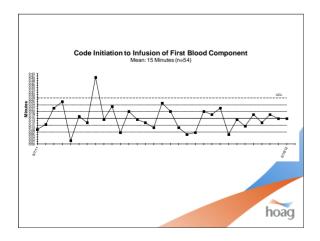
FDA-lab	peled indications		Dose (IV)	Frequency
	Bleeding episodes in patients with known Hemophilia A or B with inhibitors to Factor VIII and IX	90 mcg/kg IV bolus every 2 hours until hemostasis achieved or treatment judged ineffective	90 mcg/kg	
	Surgical bleeding prophylaxis in patients with known Hemophilia A or B with inhibitors to Factor VIII and IX	90 mcg/kg IV bolus immediately before surgery, repeat every 2 hours for duration of surgery and up to 48 hours, then every 2-6 hours until healed. In major surgery, may continue every 2 hours x 5 days, then every 4 hours until healed.	90 mcg/kg	
	Acquired hemophilia	70-90 mcg/kg IV bolus every 2-3 hours until hemostasis achieved	90 mcg/kg	
	Congenital Factor VII deficiency	15-30 mcg/kg IV bolus every 4-6 hours until hemostasis achieved	30 mcg/kg	
Uses in patients without hemophilia or Factor VII deficiency: Massive hemorrhage refractory to standard therapy secondary to surgery or trauma		Must write new order for each repeat dose desired for uses below:		
	Peri- and post-operative bleeding	15-45 mcg/kg IV, may repeat after 30 minutes for up to 3 doses (Due to thromboembolic risks, the lowest possible dose is recommended)	☐ 15 mcg/kg IV x 1	
	Trauma-associated bleeding	15-90 mcg/kg IV, may repeat after 20 minutes for up to 3 doses (Due to thromboembolic risks, the lowest possible dose is recommended)	IS INCO	Ng IV X I
	ses requiring a Medical Director	r, Transfusion Medicine; Pathology; or		
	Post-partum hemorrhage	50-100 mcg/kg IV bolus every 2 hours until hemostasis achieved	☐ 30 mcg	kg IV x 1
П	Reversal of anticoagulation therapy	Non-emergent bleeding: 15-40mcg/kg IV bolus, may repeat Emergent/life-threatening bleeding: 41-90mcg/kg IV bolus, may repeat	□ 45 mag	kg IV x 1
	Intracerebral hemorrhage	40-80mcg/kg IV bolus x 1 dose	_ ic incg	g
	Intracerebral hemorrhage Severe thrombocytopenia	40-80mcg/kg IV bolus x 1 dose 50-100mcg/kg IV bolus, may repeat		

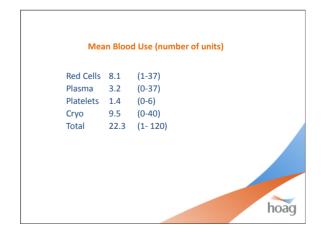


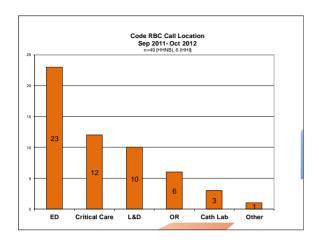


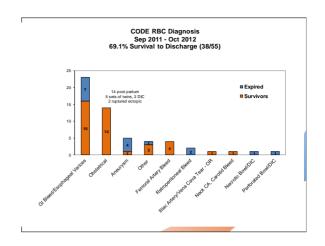


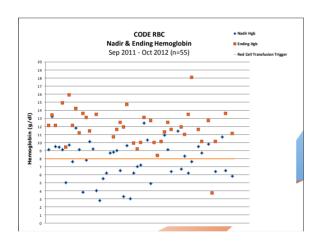


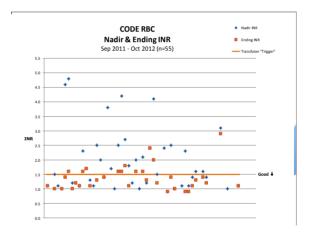


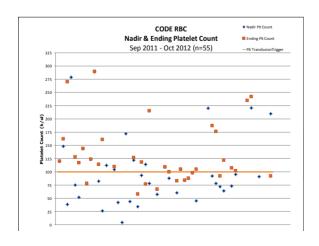


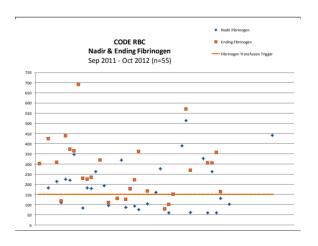












Lessons Learned

- Literature review key to physician member buy-in
- · Initial and ongoing education critical
- · Ongoing case review for continuous improvement
- · Some physicians still "want what we want"
- Additional components may be indicated.
- · Poor compliance with laboratory monitoring
- · Staff love standardized approach, order from chaos.
- · Success breeds acceptance!
- A massive transfusion protocol can be very beneficial in non-trauma center hospitals!

hoag

Case Review - Tumor Debulking

- 54 y/o female hysterectomy and tumor debulking.
- Estimate blood loss 6200 ml
- 1.5 L Cell Saver Blood
- 47 Blood Components
 - 13 red cells, 9 plasma, 5 platelets, 20 cryo
- Nadir Labs
 - Hgb 9.5, Plt 239, Fibrinogen 182, INR 1.5
- Post-op Day One Labs
 - Normal kidney and liver function
- Discharge post-op day 7

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Case Review - Placenta Accreta

- 29 y/o g1p1
- Mild preeclampsia
- · Rupture of membranes at 35 weeks.
- Started on Pitocin and IV antibiotics Normal delivery, apgar 9 & 9
- No placental delivery after 30 minutes or after attempts at manual extraction
- Probable placenta accreta
 - abnormally deep attachment of the placenta, through the endometrium and into the myometrium

hoaq

Case Review - Placenta Accreta

- 750 ml blood loss, became tachycardia and hypotensive
- · Taken urgently to the OR and CODE RBC called.
- Second attempt at manual extraction, only partially successful
- Exploratory lap and abdominal hysterectomy performed.
- 3000 ml blood loss.

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Case Review - Placenta Accreta

- 48 total blood components
 - 21 red cells, 13 plasma, 4 platelets, 10 cryo
- 5000 unit factor VIIa
- Labs
 - Hemoglobin 4.3 9.3
 Platelet 25 72
 Fibrinogen 274 274
 INR 1.4 1.2
- Discharged day 5

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Case Review - Back Procedure

- 55 year old female
- Elective L4-L5 laminectomy, discectomy and expandable cage inter-body fusion using minimally invasive technique
- · Vena cava and iliac artery laceration
- 10,000 ml EBL
- 50 total blood components
 - 18 red cells, 9 plasma, 3 platelets, 20 cryo
 - 3600 ml cell saver blood

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