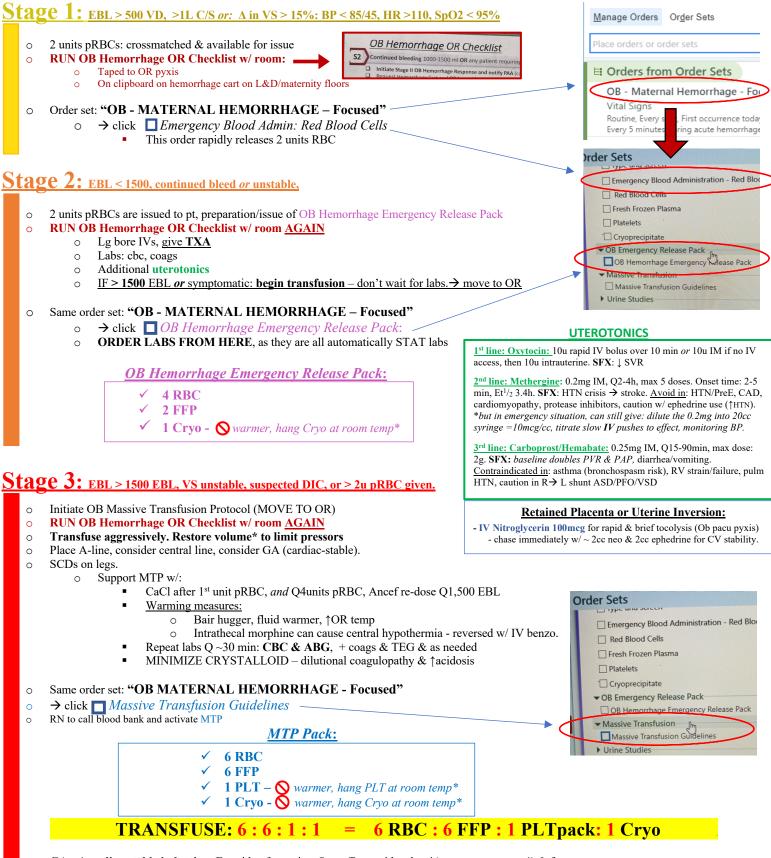
## **OB MATERNAL HEMORRHAGE**



## - Anesthesia Guide -



- GA w/ cardiac-stable induction: Etomidate/ketamine, Succs/Roc, midazolam\*(prevent awareness!) & fent
  - Anticipate airway swelling\*\* → position airway! √Consider safety of extubation if airway swollen
    - Swelling will only worsen w/ PP intravascular fluid shifts esp after MTP. Check cuff leak → ICU if unable to safely extubate.
  - Can maintain GA w/: 50% N<sub>2</sub>O <u>and</u> either ½ MAC sevo, or a propofol gtt (prop preferred in uterine atony) +/- ketamine if unstable or atony. 0
  - Parturients more susceptible to pulm edema esp Pre-E/HELLP pts <sup>2</sup>/<sub>2</sub> endothelial damage of disease process → ++ capillary leak
    - may need diuresis. (can use POCUS lung US to monitor edema & response to tx)

## **OB HEMORRHAGE NOTES:**

- Normal fibrinogen *in pregnancy* is \*\* 350-650 mg/dL \*\*
  - o normal non-pregnant is 200-400mg/dL
    - A "normal" fibrinogen of ~200 in a *pregnant* pt is *abnormally low* and suggests DIC.
- If epidural catheter in place during hemorrhage, leave in place until labs prove safety of removal (PLT wnl & fibrinogen > 300)
- ACOG defines PPH as > 1L QBL in 1st 24 hrs, or any blood loss with s/s hypovolemia, regardless of mode of delivery.
  - Vaginal > 500cc EBL is escalated, and C/S > 1000mL is escalated to Stage 1 Response.
  - General guideline: any delivery > 1500cc EBL should have received pRBCs/products other than crystalloid.
- A leading cause of delayed PPH identification, response, and tx is imprecise estimation of actual blood loss during birth & immediate PP period (ACOG)
- ~ 40% of PPH occurs in low-risk women. EVERY woman giving birth is at risk for PPH (ACOG)
- Tachycardia or Hypotension is  $\frac{2}{2}$  hemorrhage until proven otherwise and requires continuous reassessment.
- Healthy women can compensate for significant hypovolemia/blood loss... until they can't anymore. VS Δ at ~20-30% blood volume lost.
- Baroreceptive HR response to hypovolemia can be blunted by beta blockers (Pre-E pts) or neuraxial for c/s (T4 block cardiac accelerator fibers T4-T1) or just youth & good health. Unexplained HoTN absent tachycardia does not rule out hemorrhage.
- Bleeding can occur for reasons <u>OTHER THAN UTERINE ATONY</u>:
  - eg: vaginal/cervical lacerations, coagulopathy, etc.
  - o Extra-uterine, intra-abdominal, or retroperitoneal bleeding will not display vaginal bleeding, and persists regardless of uterine tone.
- Uterine clots can conceal intrauterine hemorrhage. "Adequate uterine tone" can be subjective and does not rule out hemorrhage as cause of Δ in VS.
- Allowable Blood Loss equation for estimating Hgb should use QBL instead of EBL.
  - <u>Use existing hemorrhage protocols and established trigger volumes to initiate resuscitation</u>. ABL equation can help assure we have not underresuscitated, as vitals and initial lab values can inaccurately reflect severity of blood loss.
  - Qualitative Blood Loss should replace Estimated Blood Loss as EBL is imprecise if mixed with amniotic fluid.
  - Est Blood Vol (EBV) mL/kg standards require adjustments in pregnancy, and for levels of obesity:
  - o Allowable  $QBL^* = [EBV \times (H_{initial} H_{final})] / H_{initial}$
  - Functional blood loss can be underestimated in weighed QBL if large clots present. Weight-wise, clots contain greater concentration of lost RBC/PLT/FACTORS than liquid blood\*
- Intra-op Hgb values can be slightly higher <sup>2</sup>/<sub>2</sub> hemoconcentration & compensatory hypovolemic vasoconstriction, whereas postpartum Hgb/Hct labs often see further dilutional drop as extravascular fluid shifts back into intravascular compartment.

able 2 Blood volume estimations by BMI category.  ( In pregnancy )		
NHS obesity classification	BMI range (kg m <sup>-2</sup> )	Blood volume (ml kg <sup>-1</sup> )
Healthy weight	18.5–24.9	95
Overweight	25–29.9	85
Obese	30–39.9	75
Severely obese	>40	70

- Lactate can normally reach levels of  $\sim 2.0 4.0 \text{ mmo/L}$  during labor, but > 4.0 is unusual.
  - $\circ$  (so while a vaginal delivery or F+P  $\rightarrow$  c/s may have elevated lactate, a planned c/s not in labor should not.)
- Maternal MRO<sub>2</sub> and consumption (VO<sub>2</sub>) is higher than non-pregnant levels. (VO<sub>2</sub> \(^\rightarrow\) by \(^2\rightarrow\) by \(^2\rightarrow\) by \(^2\rightarrow\).
  - Remember *SpO<sub>2</sub> can be normal*, while actual O<sub>2</sub> carrying capacity (DO<sub>2</sub>), and therefore availability and uptake (VO<sub>2</sub>) can be compromised, making the pregnant pt vulnerable to shock & acidosis in PPH.
    - $DO_{2 \text{ mL/min}} = C.O. \text{ x } [1.34 \text{ x Hgb x } SaO_2 + (0.30)] \leftarrow Arterial O_2 \text{ concentration}$

1.34 =  $O_2$  binding capacity of Hgb  $SaO_2$  as fraction: 100% = 1, 90% = 0.9, etc 0.30 = dissolved  $O_2$  in blood at  $PO_2$  of  $100_{mmHg}$ 

Admission Hemorrhage Risk Factor Evaluation			
Low (Clot only)	Medium (Type and Screen)	High (Type and Crossmatch)	
No previous uterine incision	Prior cesarean birth(s) or uterine surgery	Placenta previa, low lying placenta	
Singleton pregnancy	Multiple gestation	Suspected Placenta accreta or percreta	
≤4 previous vaginal births	>4 previous vaginal births	Hematocrit <30 AND other risk factors	
No known bleeding disorder	Chorioamnionitis	Platelets <100,000	
No history of PPH	History of previous PPH	Active bleeding (greater than show) on ac	
·	Large uterine fibroids	Known coagulopathy	
	Estimated fetal weight greater than 4 kg		
	Morbid obesity (BMI >35)	(CMQCC 2009)	