Patient addressograph



Postpartum Haemorrhage Management Checklist

Stage 0	St	age 1			
PPH Risk Assessment	>500ml on-going blood loss				
Complete for all women in labour notes (including LSCS)	Vaginal deliveries including instrumental				
Most recent Hb=Plts: Date:	Get Help		Time	Initial	
☐ Antenatal Clexane; last dose:	Notify midwife in charge				
Antenatal "Increased risk" (if any of the following are met)	Name:	_ time arrived:	:		
Anaemia or bleeding disorder (Hb <95; Plt <100)	Request support to assist	t with measurement			
BMI <18 or >35; Booking weight <55kg if low weight/BMI: do you need to calculate the circulating blood volume?	Other staff present	Designation	Time arrived	Initial	
≥5 previous vaginal births					
Previous Postpartum Haemorrhage >1I					
Estimated fetal weight >4.5kg					
Multiple pregnancy	Act	Performed by	Time	Initial	
Polyhydramnios	Measure blood loss				
Previous uterine surgery	(cumulative measurement)				
Abnormal placental implantation	Record observations On MEOWS every 10 minutes				
Known abruption or AN haemorrhage	IV access				
Please make an on-going assessment of the following risk factors throughout labour and delivery	at least 16 Gauge				
	What is the caus	e of bleeding?	•		
Perinatal "Increased risk" (if any of the following are met) Suspicion of chorioamnionitis / Sepsis	Tone / Trauma / Tiss	ue / Thrombin (circl	e cause(s))	
Labour augmented with Syntocinon	Treat	Performed by	Time	Initial	
Prolonged labour	Uterine massage				
Instrumental delivery	Give uterotonics				
Retained products of conception	record on over page & prescribe				
Plan to measure & record all blood loss	Inspect genital tract				
(for pool deliveries estimation may be required)	Empty bladder				
Act If women at increased risk is:	Check placenta & mem- branes				
She suitable for El blood or 2 units Xmatch? Yes / No IV access required? Yes / No	Bimanual compression				
Treat Planned active 3rd stage management? Planned 40IU Syntocinon IV slow infusion? Yes / No	If bleeding stopp Record measured blood	d loss here:		ml	
Completed by (name): Date:Time:Location:	Completed by (name): Date:Time:				

Otage 2							(0 1	/40mmHg; SpO ₂ <	• •	
	Prog	ress	to here from stage	1 if va	ginal	delivery	. Re-start here afte	er stage 0 if LSCS		
Get He	lp	Name	e	Time arriv	ved	Othe	r staff present	Designation	Time arriv	ved
MW in charg	je			:					:	
Obstetrician	ı			:					:	
Anaesthetis	t			:					:	
Care assista	int / scribe			:					:	
Act							Performed	i by	Time	Initial
Measure & r	ecord cumula	ative	blood loss							
Lie patient f	lat and give o	xyge	en							
Record obs	ervations on	MEO\	WS every 5 minutes							
2nd IV acce	ss (at least 16	Gau	ige) & fluid bolus							
Take	Point of care to	ests	venous lactate; venous	Hb						
bloods	Lab tests		FBC; Coag; XMatch; U& Serum Fibrinogen	ßΕ;						
	Ini		venous blood test r	esult	S		Initial Fibrino	gen or Rotem Test re	sults	
Time:	Hb=		Lactate=			FIBTE	// A5 (Aim ≥ 12mm) =	EXTEM CT (Aim < 75sec) =		
Review	the caus	ses	of bleeding (d	circle all i	dentified)		Tone / Traur	na / Tissue / Thron	nbin	
Treat			Performed by	Time	Initial			Performed by	Time	Initial
	rotonics record	d on				Empty	bladder			
page 3 & prescri	ibe					Foley	catheter inserted			
Give tranex	amic acid					Inspec	t genital tract			
1g IV, if no contr	raindications					Repair	genital tract			
Bimanual c	ompression						placenta &			
Consider O	meprazole					membi	anes			
If ble	eding stop	pec	d ensure PPH pos	t-ever	nt che	cklist co	mpleted and Mana	gement plan written in	notes	
Complet	ed by (name): _				Dat	e:	Time:	Location:)
If bleedi	ng on-goii	ng ti	ransfer patient t	to th	eatre			Time arrived:		
Stage 3	>15	00n	nl blood loss Ol	R on	qoin	a clini	cal concern			
Act						J		rformed by	Time	Initial
Communic	ate current m	easu	red blood loss to tea	am						
Activate M	OH protocol									
Inform Obstetric and Anaesthetic consultants										
	d and coagul discuss the case w		products as per pro ematologist?	tocol						
Review	the causes	s of	bleeding (circle all id	lentified)			Tone / Trauma	/ Tissue / Thrombi	n	
Treat							Pe	rformed by	Time	Initial
Review ute	erotonics recor	d on pa	age 3 & prescribe							
Consider repeat tranexamic acid if bleeding ongoing 1g IV, if I			IV, if no	Cl's						
			techniques record on pa							
Other staff			signation		arrived			Designation	Time arriv	ived
Name:						Name				

Name:

Date:

____Time: ___

Once bleeding stopped ensure PPH post-event checklist completed and Management plan written in notes

Completed by (name): __

Name:

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Record of Uterotonics used Please record all uterotonics used here and prescribe as per protocol						
Drug	Dose (circle route)	Time	Drug	Dose(circle route)	Time	
Oxytocin	10 units IM or 5 units IV		Carboprost (caution in asthma)	250 mcg IM (repeat up to every 15 minutes)		
			Carboprost	250 mcg IM		
Ergometrine (caution in HTN/PET)	500 microgram IV or IM		Carboprost	250 mcg IM		
Syntometrine (caution in HTN/PET)	500 mcg/5units IM or IV		Carboprost	250 mcg IM		
Oxytocin infusion	40 units over 4 hr IV		Carboprost	250 mcg IM		
			Carboprost	250 mcg IM		
Misoprostol			Carboprost	250 mcg IM		
Misoprostol						

Blood & blood products transfused						
Time	Product given	Time	Product given			

Measured cumulative blood loss				
Time	Blood loss (ml)	Running total (ml)		
Total measured blood loss m				

Record of further blood test results Please do not duplicate records of blood results recorded in stage 2						
	Further venous	blood test results	Further Fibrinogen or Rotem Test results			
Time:	Hb=	Lactate=	FIBTEM A5= (Aim ≥ 12mm)	EXTEM CT= (Aim < 75sec)		
Version 2 May 2000 Provided A						

Date/Time	Documentation of co	ncerns, de	viations & oth	er information			
PPH pos	st-event checklist						
WHO sign-o	ut completed?	Yes / No	NA (patient did n	ot require care in theatre)			
Have all dru	gs been prescribed and signed for?	Yes / No	NA:				
Post-event	Re-bleed assessment						
Syntocinon i	infusion running or required?	Yes / No	Time expected to	o finish:			
Vaginal pacl	k insitu?	Yes / No	Planned removal	time:			
Intrauterine	ntrauterine tamponade balloon insitu? Yes / No Planned removal time:						
Can NSAID be given?		Yes / No	Not yet				
Thromboprophylaxis plan		LMWH	Yes / No	Time first dose:			
		TEDS	Yes / No				
Post-event	Monitoring requirements						
Level of pos	t-event care required (circle applicable)	Level 1	Level 2 (ECLW)				
Post-op bloc	ods <i>(FBC/Coag/U&E)</i> to be taken at	Time:		Plan to transfuse if Hb <			
PV loss mor	nitoring required?	Yes / No	Frequency of mo	nitoring:			
Urine output	monitoring required?	Yes / No	Frequency of mo	nitoring:			
MOH stand	down	Yes / No	N/A				
Any blood/pr	roducts to return to blood bank?	Yes / No	N/A				
If the MOH protocol was activated before stage 3 or not activated at stage 3 then please detail reason(s) why:							
Does a Dati	x form need completing?						
If yes record	t:						
	Datix form number						
Pe	erson responsible for completing Datix form						
Duty of candour and team debriefing							
	nt been discussed with the patient?	Yes / No					
	information been provided to the patient?	Yes / No					
Does a form	nal team debrief need to take place?	Yes / No					
Completed by	(name): Da	te:	Time:	Location:			