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# Management of Antepartum Haemorrhage

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# Antepartum Haemorrhage

## 1. DEFINITION

Vaginal bleeding after 24 weeks gestation. Occurs in 3-5% of pregnancies

## 2. AIMS

To reduce risk of maternal and perinatal morbidity and mortality

- Identify high risk cases
- Identify cause of bleeding
- Accurately estimate blood loss
- Adequate resuscitation including management of coagulopathy
- Expedite delivery if indicated

## 3. CAUSES

- Placenta praevia
- Placental abruption
- Vasa praevia
- Localised bleeding to vulva, vagina or cervix

## 4. DEFINITION

- Spotting – streaking, staining or blood spotting on underwear
- Minor Haemorrhage – EBL <50mls that has settled
- Major Haemorrhage - EBL 50-1000mls with no signs of clinical shock
- Massive Haemorrhage – EBL >1000mls and/or signs of shock
- Recurrent APH - >1 occasion of APH

## 5. DIFFERENTIAL DIAGNOSIS

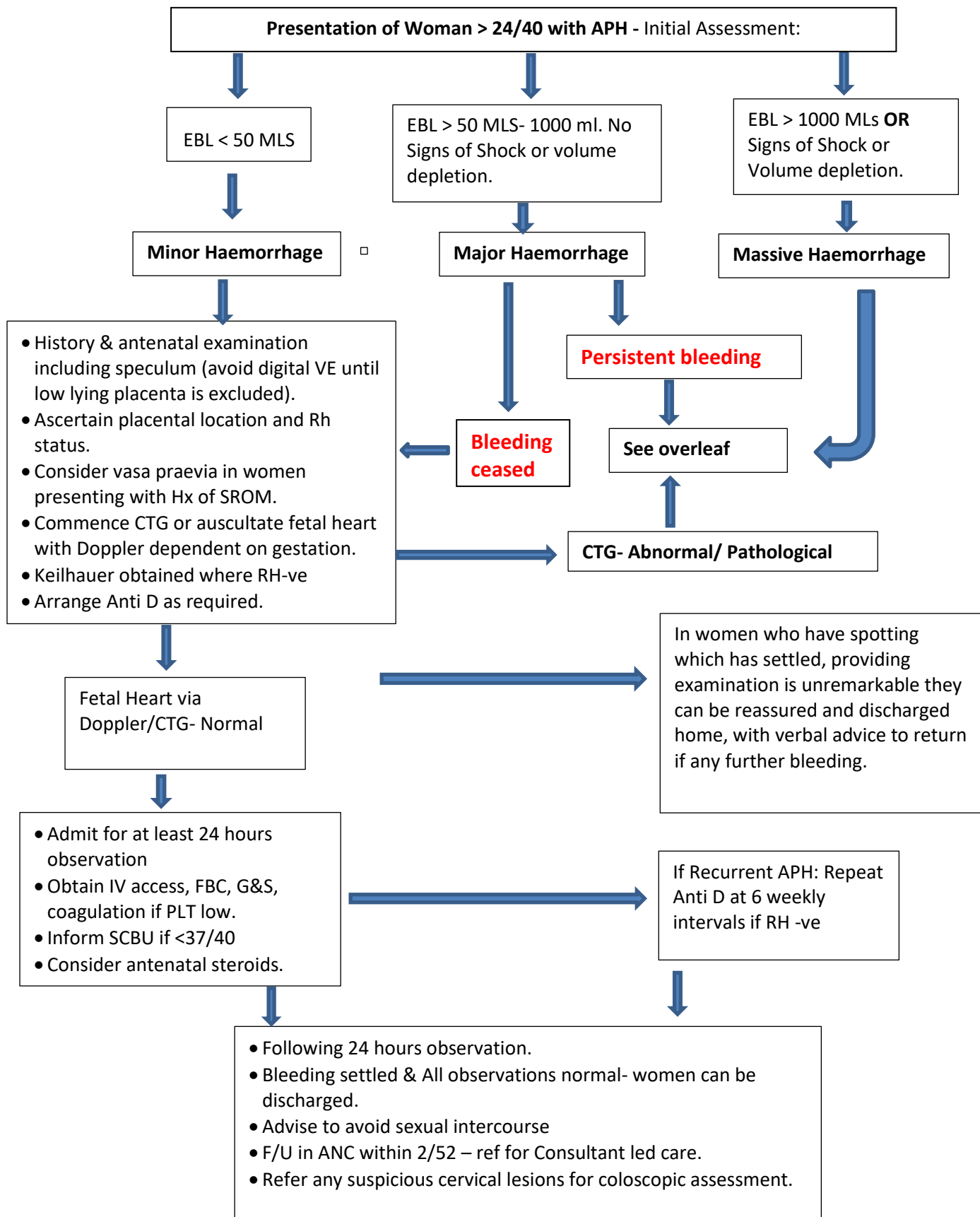
	Placental Abruption	Placenta Praevia
<b>Incidence</b>	1-2%	0.5% at term
<b>Presentation</b>	Abdominal pain Tender, tense uterus Haemorrhage may be concealed May be shock Coagulopathy	Painless bleeding, may be heavy Soft uterus Coagulopathy less common Usually stable
<b>Fetal Assessment</b>	May be abnormal CTG or IUD	CTG usually normal May be malpresentation
<b>Risk Factors</b>	<ul style="list-style-type: none"> <li>• Previous Hx of abruption</li> <li>• Pre-eclampsia</li> <li>• Fetal growth restriction</li> <li>• Non-vertex presentation</li> <li>• Polyhydramnios</li> <li>• Advanced maternal age</li> <li>• Multiparous</li> <li>• Low BMI</li> <li>• IVF</li> <li>• Intrauterine infection</li> </ul>	<ul style="list-style-type: none"> <li>• Previous Hx of praevia</li> <li>• Previous C/S</li> <li>• Hx of TOP</li> <li>• Multiparous</li> <li>• &gt;40 yrs</li> <li>• Multiple pregnancy</li> <li>• Smoking</li> <li>• IVF</li> <li>• Deficient endometrium</li> <li>• Uterine scar</li> </ul>

	<ul style="list-style-type: none"><li>• PROM</li><li>• Abdominal trauma</li><li>• Substance misuse</li><li>• First trimester bleeding</li><li>• Maternal thrombophilia</li></ul>	<ul style="list-style-type: none"><li>• Endometritis</li><li>• MROP</li><li>• Curettage</li><li>• Submucous fibroid</li></ul>
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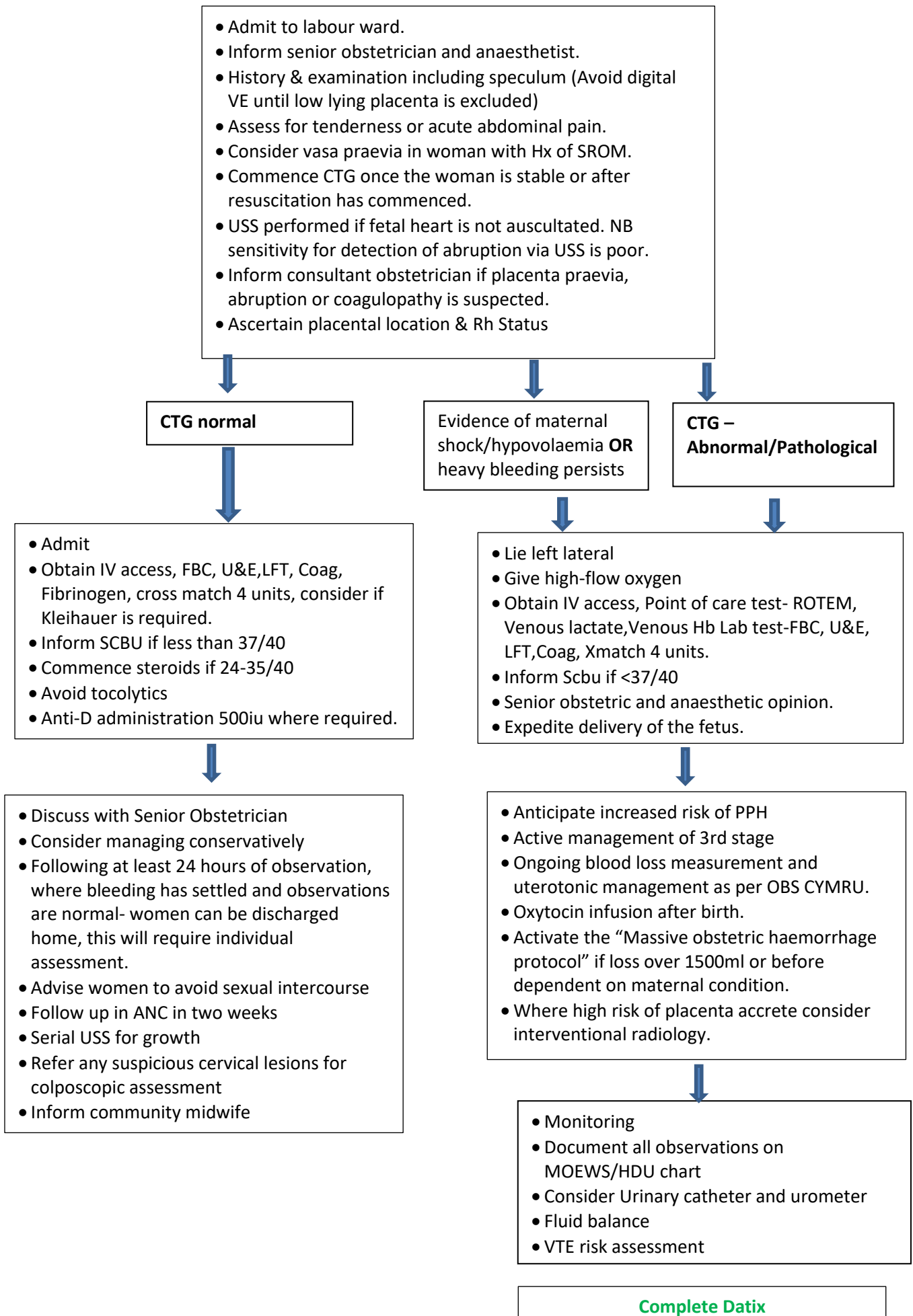
## 6. VASA PRAEVIA

- Incidence 0.1%
- Bleeding with onset of membrane rupture
- Fetal distress is a common feature
- Immediate delivery indicated as high risk of perinatal mortality from exsanguination (total blood volume of infant at term approx 250ml)

## 7. PRESENTATION OF WOMAN >24/40 WITH APH PATHWAY



## 8. MAJOR HAEMORRHAGE PATHWAY



## 9. MASSIVE HAEMORRHAGE PATHWAY

Senior Obstetrician & Senior anaesthetic involvement, haematologist and blood bank to be informed. Activate "Massive obstetric haemorrhage protocol" on clinical opinion or where EBL/MBL reaches 1500ml.

- Airway, Breathing, Circulation- commence high flow oxygen.
- Lie left lateral
- Ongoing assessment of maternal observations including Oxygen saturation and respiration
- Obtain IV access (2x 16g), Point of care test- ROTEM, venous lactate, venous Hb. Lab test- FBC, U&E, LFT, Coag, Xmatch 6 units urgently. Women who are Jehovah's witness are managed in line with Labour ward guidelines.
- Infuse 2L warmed crystalloid (Hartmanns or 0.9% Normal Saline)
- Commence CTG once mother stabilised or after resuscitation has commenced.
- USS should be performed if unable to auscultate the fetal heart. NB sensitivity for detection of abruption via USS is poor.
- Inform SCBU

- History & examination including all observations and speculum (avoid digital VE until low lying placenta is excluded).
- Assess for tenderness or acute abdominal pain.
- Ascertain placental location and Rh status (NB. Sensitivity for detection of abruption ultrasound is poor).
- Consider vasa praevia in women presenting with Hx of SROM
- Stop any anticoagulation therapy.
- Measure ongoing bleeding.

- Evidence of maternal shock or volume depletion OR
- Heavy bleeding persists OR
- Evidence of fetal compromise.

- Discuss with consultant obstetrician.
- Plan to expedite delivery of fetus
- Woman must be stabilised first, unless evidence of life threatening bleeding.
- Consider cell salvage

- Anticipate increased risk of PPH
- Active third stage
- Manage ongoing bleeding and uterotonics as per OBS CYMRU.
- Oxytocin infusion after birth.
- Contemporaneous blood loss measurement
- Activate "massive obstetric haemorrhage protocol" if loss >1500 mls or signs of hypovolemic shock.

- Consider DIC in major/massive haemorrhage. Undertake ROTEM
- If DIC suspected, discuss with haematologist.
- Infuse 6 units PRBC plus 4 units of FFP whilst waiting for blood results.

Ongoing Monitoring

Document all observations on HDU/MOEWs

Consider Urinary catheter and urometer

Strict fluid balance

VTE risk assessment

Bloods as clinically indicated

Consider need for kleihauer and Anti D.

Complete Datix

## 10. NON-ACUTE ANTENATAL MANAGEMENT OF PLACENTA PRAEVIA/ACCRETE

- Trans-vaginal USS to confirm placental site
- Colour flow Doppler if risk of accreta – if not available locally to treat as if accreta
- Patient advised to attend hospital if any abdominal pain, tightening or bleeding
- Antenatal discussion with woman regarding:
- Delivery Haemorrhage
  - Risks
  - Surgical management including hysterectomy
  - blood transfusion / cell salvage
- Antenatal discussion with Interventional Radiologist in relation to possible catheterisation
- Blood X-match
- If placenta <2cm from internal os, likely to need LSCS for delivery
- Anaesthetic referral to discuss mode of anaesthetic (likely regional). If suspected accreta refer early to anaesthetic clinic for multidisciplinary plan.
- Most senior obstetrician and anaesthetist should at the very least be available on labour ward if intervention required as either elective or emergency
- Where possible elective LSCS performed after 38 weeks gestation to minimise neonatal morbidity
- All staff should be aware of major haemorrhage guideline and drills performed at regular intervals
- Tailor antenatal care, including hospitalisation, to individual woman's needs e.g. distance between home and hospital and availability of transportation, previous bleeding episodes, haematology laboratory results, and acceptance of receiving donor blood or blood products.
- Where hospital admission has been decided, an assessment of risk factors for venous thromboembolism in pregnancy should be performed. This will need to balance the risk of developing a venous thromboembolism against the risk of bleeding from a placenta praevia or low lying placenta.



## REFERENCES:

RCOG (2011) *Antepartum Haemorrhage – Green-top guideline No. 63*, RCOG.

ABMU (2011) *Guideline for the Management of Antepartum Haemorrhage*. ABMU HB.

Winter C., Crofts J., Draycott T., Muchatuta N. (2018) *PROMPT 3<sup>rd</sup> Ed.* Cambridge University Press.

## Maternity Services

### Checklist for Clinical Guidelines being Submitted for Approval

Title of Guideline:	Management of Antepartum Haemorrhage
Name(s) of Author:	Antenatal forum
Chair of Group or Committee approving submission:	Carolyn Williams
Brief outline giving reasons for document being submitted for ratification	Document update.
Details of persons included in consultation process:	Antenatal forum/Labour ward forum members.
Name of Pharmacist (mandatory if drugs involved):	
Issue / Version No:	
Please list any policies/guidelines this document will supercede:	Guideline for the management of antepartum Haemorrhage.2011
Date approved by Group:	6 December 2019
Next Review / Guideline Expiry:	December 2022
Please indicate key words you wish to be linked to document	APH, Antepartum Haemorrhage, massive obstetric haemorrhage,
File Name: Used to locate where file is stores on hard drive	