1. **Purpose**

This document outlines the procedure details for the use of balloon tamponade in managing primary major postpartum haemorrhage at the Women’s.

This procedure has been developed to assist clinicians managing primary postpartum haemorrhage in order to reduce the morbidities and mortality associated with major blood loss after childbirth. This procedure is linked to the following guideline and procedures:

- [Postpartum Haemorrhage](#)
- [Postpartum Haemorrhage - Immediate and Ongoing Postnatal Care after Major PPH](#)
- [Postpartum Haemorrhage - Prostaglandin F2 Alpha](#)


2. **Definitions**

**Bakri balloon** is a balloon tamponade indicated for women not responding to uterotonics and uterine massage.

It is used to control haemorrhage due to uterine atony in the upper segment of the uterus and to control bleeding in the lower uterine segment secondary to placental implantation in the lower uterine segment.

3. **Responsibilities**

**Obstetric and midwifery staff** are responsible for recognising and promptly managing postpartum haemorrhage, for collaborating with other clinicians necessary for the woman’s care, escalating to senior clinicians in cases of major PPH.

**Senior medical staff (on-call Obstetric Consultant)** are responsible for attending all cases of major PPH or on request.

**Anaesthetic staff** are responsible for providing and advising on clinical care in cases of major PPH when intensive monitoring and resuscitation are required.

The **Haematology Consultant** should be consulted early to prevent development of DIC; all cases of developing/actual coagulopathies; resuscitation with blood products required.

**Other available specialists** such as Gynae/Oncology Consultant should be consulted early when bleeding is intractable, where hysterectomy or ligation/embolisation of uterine arteries are being considered.

4. **Procedure**

**Balloon tamponade** is indicated for women not responding to uterotonics and uterine massage

- To control haemorrhage due to uterine atony in the upper segment of the uterus
- To control bleeding in the lower uterine segment secondary to placental implantation in the lower uterine segment, either where the placenta has been delivered complete but the placental site has not properly contracted or there is an abnormally adherent placenta in the lower uterine segment
- Before laparotomy to arrest haemorrhage in placenta accreta

### 4.1 Insertion of the Bakri Balloon

**In the case of uterine atony unresponsive to uterotonics:**

Ensure the uterus is clear of any retained placental fragments, blood clot, arterial bleeding or lacerations before inflating balloon.

Insert balloon catheter under spinal, epidural or general anaesthesia in theatre.

Introduce vaginal speculum and using sponge forceps, insert balloon catheter transvaginally into the uterine cavity under guided ultrasound.
Procedure

Postpartum Haemorrhage - Bakri Balloon Tamponade

Once in place, inflate balloon with a volume of 100 - 300 mL of warm 0.9 % sodium chloride until enough counter pressure is exerted to stop bleeding from uterine sinuses (usually fill balloon until visible in the cervix lumen).

The test result is considered successful if there is no bleeding through the cervix or through the drainage channel of the balloon catheter.

If bleeding continues, the tamponade test is unsuccessful and surgery is needed.

Document amount of fluid in balloon.

Apply gentle traction to balloon and tape balloon to the woman’s inner thigh to maintain tension.

In the case of bleeding from the lower uterine segment:

As above steps 1 & 2.

Insert balloon into lower segment with the tip of the catheter in the uterine cavity.

Inflate balloon under ultrasound guidance with up to 500 mL warm 0.9 % sodium chloride.

Pack the vagina to ensure the balloon stays in place.

Continue to observe the uterus by ultrasound scanner, the output from the Bakri catheter and the vaginal loss.

4.2 Management following insertion

High dependency or intensive care for ongoing management.

If the balloon tamponade is for uterine atony, continue 40 U oxytocin / 1000 mL 0.9% sodium chloride infusion over 4 hours until after removal of the balloon catheter (helps keep the uterus well contracted over the balloon).

Observations

Hourly urine output, blood pressure, pulse rate, respiratory rate, oxygen saturation, fundal height and vaginal blood loss (through the lumen of the catheter) until stable.

Temperature every two hours (every hour if blood transfusion in progress).

A strict fluid balance chart must be kept with input/output recordings made at least hourly. The estimated/weighed blood loss from the postpartum haemorrhage must be included to ensure this is accounted for when making decisions about fluid balance.

Antibiotics

Administer IV antibiotics (ampicillin [or amoxycillin] 2g IV initial dose then 1g IV every 4 hours, gentamicin 5 mg / kg IV as a single daily dose, metronidazole 500 mg IV every 12 hours) until after removal of the balloon catheter.

4.3 Removal of Bakri balloon

Leave balloon tamponade in place for 8 to 24 hours to allow time for blood transfusion and coagulopathy correction.2,5

Once parameters are within acceptable limits, deflate the balloon in two stages – withdraw half the 0.9 % sodium chloride, and if no significant bleeding after 30 minutes, withdraw the remaining volume to deflate and remove balloon.

Continue to observe the woman for any active bleeding.

Documentation

Time and date of Bakri balloon insertion and removal to be documented in surgical record.

5. Evaluation, monitoring and reporting of compliance to this procedure

Compliance to this procedure will be monitored, evaluated and reported through the notification of clinical incidents on VHIMS and by monthly clinical audit of PPHs greater than 1500mLs.

6. References


7. Legislation related to this procedure

Not applicable.

8. Appendices

Not applicable.

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