Tamponade technique for postpartum hemorrhage

Refer to the Instructions for Use for complete information on product usage and a complete list of precautions, warnings, and contraindications.

Confirm before placement.

Confirm that these statements are true:

- The uterus is free of placental fragments.
- The genital tract has no trauma or lacerations.The source of the bleeding is not arterial.
- Patient does not present with any contraindications for use of this device.

Determine the uterine cavity's volume.

- Determine the uterine cavity's volume by direct or ultrasound examination.
- Place the predetermined volume of sterile fluid in a separate container.
- If you will use the rapid instillation components, note the predetermined
- volume for rapid instillation.
- The maximum balloon volume is 500 mL.

Place the balloon.

Transvaginal placement, postvaginal delivery (Fig. 1)

 Insert the balloon portion of the catheter into the uterus, making certain that the entire balloon is inserted past the cervical canal and internal ostium.

Transabdominal placement, postcesarean delivery (Fig. 2)

- Pass the uninflated balloon, inflation port first, through the cesarean incision and into the uterus and cervix. Remove the stopcock to facilitate placement, if desired.
- Have an assistant pull the balloon shaft through the vaginal canal until the base of the balloon contacts the internal cervical ostium.
- Close the incision, being careful not to puncture the uninflated balloon while suturing.

Fill the balloon with sterile liquid.

- Never inflate with air, carbon dioxide, or any other gas.
- Do not fill with more than 500 mL. Overinflation may result in the balloon being displaced into the vagina.
- Ensure that all product components are intact and that the hysterotomy is securely sutured prior to balloon inflation.
- Place a Foley catheter in the patient's bladder to collect urine and monitor urine output.
- Use the enclosed syringe or rapid instillation components to fill the balloon to the predetermined volume through the stopcock.
- Apply traction to the balloon shaft by securing it to the patient's leg or attaching a weight (not to exceed 500 grams).
- Use ultrasound to confirm that the balloon is properly placed.

Flush the lumen and monitor hemostasis.

- Connect the drainage port to a fluid collection bag to monitor hemostasis.
 The balloon drainage port and tubing may be flushed clear of clots with
- sterile isotonic saline to facilitate monitoring. • Monitor the patient for signs of increased bleeding and uterine cramping.

Remove the balloon.

- Maximum indwelling time: 24 hours.
- The attending clinician determines when the balloon is removed after bleeding is controlled and the patient is stable.
- Release the tension on the shaft and remove any vaginal packing.
 Aspirate balloon contents until the balloon is completely empty. The fluid may be removed incrementally to allow for periodic observation of the patient. In an emergency, the shaft may be cut to rapidly deflate the balloon.
- Gently retract the balloon and discard it.
- Monitor the patient for signs of bleeding.

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Illustrations for placing the Bakri balloon (step 3)

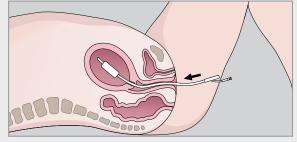


Fig. 1: Transvaginal placement, postvaginal delivery

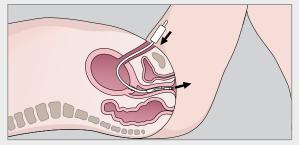


Fig. 2: Transabdominal placement, postcesarean delivery

Proper placement

A

Make sure that the entire

balloon is inserted past

the cervical canal and internal ostium.

 If necessary, pack the vagina with iodine- or antibiotic-soaked gauze.

В

• Do not extend the packing into the uterus.

use ultrasound to confirm that it is properly placed.

After the balloon

is inflated to the predetermined volume,

CONTRAINDICATIONS

- Arterial bleeding requiring surgical exploration or angiographic embolization
- Cases indicating hysterectomy
- Pregnancy
- Cervical cancer
- Purulent infections in the vagina, cervix, or uterus
- Untreated uterine anomaly
- Disseminated intravascular coagulation
- A surgical site that would prohibit the device from effectively controlling bleeding

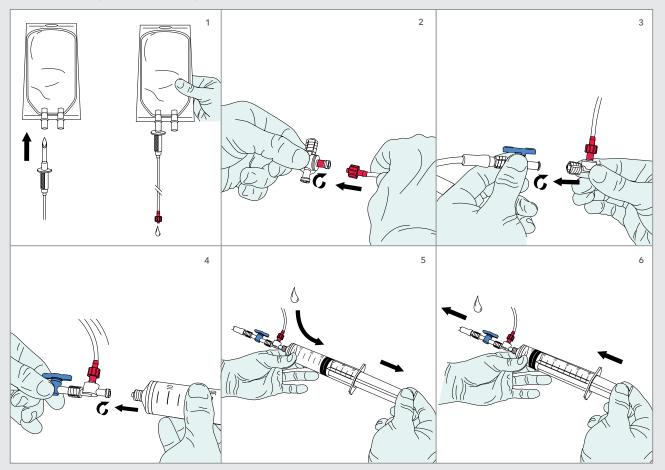
WARNINGS

- This device is intended as a temporary means of establishing hemostasis in cases indicating conservative management of postpartum uterine bleeding.
- The Bakri Postpartum Balloon is indicated for use in the event of primary postpartum hemorrhage within 24 hours of delivery.
- The device should not be left indwelling for more than 24 hours.
- The balloon should be inflated with a sterile liquid such as sterile water, sterile saline, or lactated Ringer's solution. The balloon should never be inflated with air, carbon dioxide, or any other gas.
- The maximum inflation is 500 mL. Do not overinflate the balloon. Overinflation of the balloon may result in the balloon being displaced into the vagina.
- Clinical data to support the safety and effectiveness of the Bakri Postpartum Balloon in the setting of uterine atony are limited. Patients in whom this device is being used should be closely monitored for signs of worsening bleeding and/or disseminated intravascular coagulation (DIC). In such cases, emergency intervention per hospital protocol should be followed.
- There are no clinical data to support the use of this device in the presence of DIC.
- Patient monitoring is an integral part of managing postpartum hemorrhage.
 Signs of a deteriorating or unimproving condition should lead to a more aggressive treatment and management of the patient's uterine bleeding.
- The patient's urine output should be monitored while the Bakri Postpartum Balloon is in use.
- PRECAUTIONS
- Avoid excessive force when inserting the balloon into the uterus.



Bakri® POSTPARTUM BALLOON

How to use the rapid instillation components



Customer Service

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