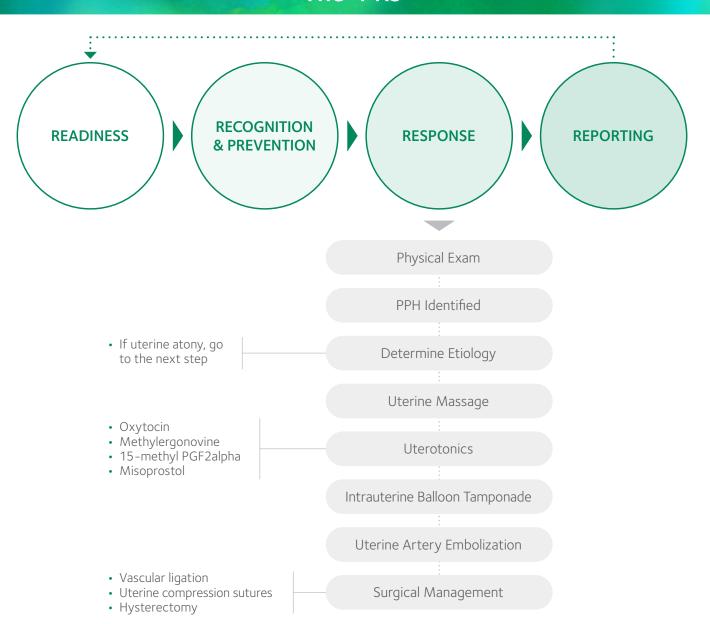


A Simple Solution for Postpartum Hemorrhage (PPH)

Sequential treatment considerations for conservative management of uterotonic-resistant PPH^{1,2,3}

The 4 Rs



The Bakri Postpartum Balloon with Rapid Instillation Components is used for temporary control or reduction of postpartum uterine bleeding when conservative management is warranted.

When inflated with sterile liquid, the Bakri Postpartum Balloon with Rapid Instillation Components applies pressure to the uterine walls that may provide management of postpartum hemorrhage.

The balloon and catheter are constructed of 100% silicone.

- Can be used after vaginal or cesarean delivery
- Includes rapid instillation components to facilitate inflation of the balloon
- May be used with B-Lynch compression sutures if clinically warranted

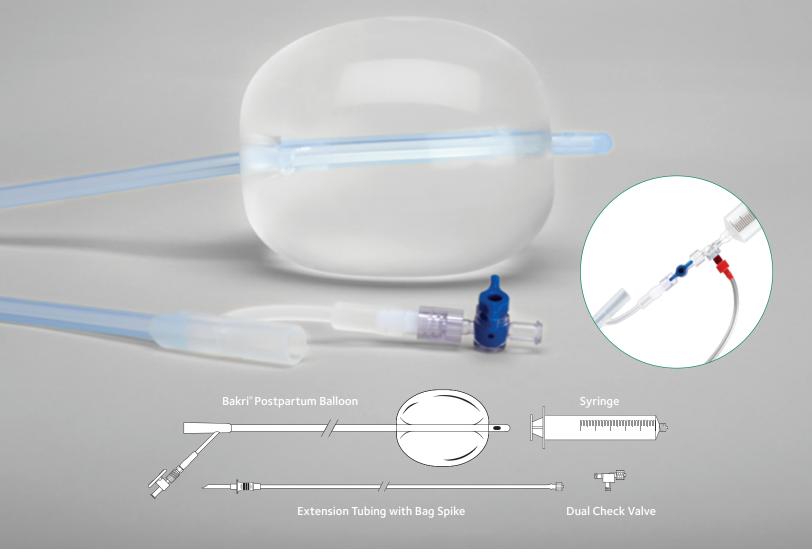
^{1.} California Maternal Quality Care Collaborative. Obstetric hemorrhage emergency management plan: table chart format. Version 2.0. Published 2014. Accessed August 31, 2016.

^{2.} Committee on Practice Bulletins-Obstetrics. Practice bulletin no. 183: postpartum hemorrhage. Obstet Gynecol. 2017;130(4):e168-e186.

^{3.} physicians.utah.edu/echo/pdfs/pregnancy-care-didactics/2016-09.09-postpartum-hemorrhage.pdf (accessed 29th October 2020).



Postpartum Balloon With Rapid Instillation Components



Ordering Information

NUMBER	REFERENCE PART #	DESCRIPTION	CATHETER	LENGTH	BALLOON VOLUME
G30673	J-SOS-100500	Bakri Balloon	24 Fr	54 cm	500 mL
G24237	J-SOSR-100500	Bakri Balloon with Rapid Instillation Components	24 Fr	54 cm	500 mL

Contact your CooperSurgical sales representative for additional information, or place your order directly with CooperSurgical Customer Service at: 800.243.2974 | 203.601.5200 | www.coopersurgical.com

IMPORTANT SAFETY INFORMATION

The Bakri® Postpartum Balloon is a single use device, intended to provide temporary control or reduction of postpartum uterine bleeding when conservative management is warranted.

CONTRAINDICATIONS: Do not use, if there is arterial bleeding requiring surgical exploration or angiographic embolization; hysterectomy is indicated; pregnancy; cervical cancer; purulent infections in vagina, cervix or uterus; untreated uterine anomaly; disseminated intravascular coagulation (DIC); or bleeding is at a site that would prohibit the device from effectively controlling bleeding.

WARNINGS: The Bakri Postpartum Balloon is indicated for use in the event of primary postpartum hemorrhage within 24 hours of delivery and should not be left indwelling for more than 24 hours; inflate with a sterile liquid to a maximum inflation of 500 mL; Do Not overinflate the balloon or use air, carbon dioxide or any other gas. Patients should be monitored closely for signs of deteriorating or non-improving condition. Patients urine output should be monitored while the balloon is in use.

IMPORTANT: Prior to placement of the balloon, the uterus should be free of all placental fragments and ensured there are no lacerations or trauma to the genital tract and source of bleeding is not arterial.

Consult the IFU prior to use of the Bakri Postpartum Balloon, for detailed instructions and potential risks. https://coopersurgical.com/bakri-ifu



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