Uterine artery embolisation in the management of recurrent vaginal haematoma

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Postpartum haemorrhage (PPH) is commonly defined as blood loss of 500 mL or more within 24 hours of birth; severe PPH is blood loss of 1 000 mL or more within the same time. PPH affects approximately 2% of all women following delivery, resulting in nearly one-quarter of all maternal deaths globally, and it is the leading cause of maternal mortality in most low-income countries. PPH is a significant contributor to severe maternal morbidity and long-term disability.1-3

Uterine atony is the most common cause of PPH (70%), but genital tract trauma (i.e. vaginal or cervical lacerations), uterine rupture, retained placental tissue or maternal coagulation disorders may also result in PPH.4,5 Coagulation disorders especially contribute to maternal morbidity rather than mortality.

When medical measures fail to control postpartum bleeding, bilateral uterine or hypogastric artery ligation may be attempted. However, owing to bilateral collateral circulation in the pelvis, the success rate of bilateral hypogastric artery ligation ranges from 40% to 100%6,7 and uterine artery ligation has been shown to have a success rate of 92%.8,9 Both these techniques aim to leave the uterus intact and preserve fertility. In 1979, transcatheter embolisation to control postpartum bleeding was initially described by Brown et al.10 Since then, several authors have reported the usefulness of this technique as first-line treatment for atomic PPH in patients who are refractory to conservative treatment,11,12 but its use in puerperal haematomas in which surgical intervention fails to control bleeding is still under-reported. We present this case report to highlight the use of transcatheter uterine artery embolisation in cases of postpartum haemorrhage due to vaginal haematoma that did not respond to surgical intervention.

Case report

A 28-year-old woman, G2P1L1, was admitted in active labour at 39 weeks' gestation. Four hours later, she delivered a live baby weighing 3 250 g by spontaneous vaginal delivery with second-degree laceration. Oxytocin 10 U intramuscularly was given as a bolus followed by an infusion of 10 U in 500 mL 0.9% saline over 4 hours. The delivery was uneventful and blood loss was estimated at 300 mL.

The patient was initially stable with a blood pressure of 120/80 mmHg, pulse rate of 90 bpm and respiratory rate of 18/ min. Two hours later she complained of excessive vaginal pain. Examination showed a blood pressure of 110/70 mmHg, a pulse rate of 100 bpm and a respiratory rate of 22/min. The uterus was well retracted, but vaginal examination revealed an 8 × 6 cm haematoma in the left lateral vaginal wall that was separate from the stitch line. The patient was moved to theatre and intravenous sedation was given. The haematoma was explored, perimal stitches were removed and about 500 mL of blood clots were removed. The vagina was closed with deep continuous interlocking sutures. After assessment of the patient's condition, two units of blood were transfused and she was monitored hourly.

However, 6 hours later, the patient again experienced the same symptoms with accompanying hypotension. On examination, she was found to be pale with a blood pressure of 100/60 mmHg, a pulse rate of 108 bpm and a respiratory rate of 24/min. Abdominal examination showed the uterus to be well retracted, but there was a left ischiorectal fossa swelling. Vaginal examination again revealed a 12 × 12 cm haematoma in the left lateral vaginal wall, which was about 3 cm short of the left fornix. The anterior and other vaginal walls were free. On rectal examination, the mass was palpable in the rectovaginal space with the rectal mucosa free. After informed consent, the patient was transferred to the operating theatre for exploration and evacuation under general anaesthesia. During exploration about 1 000 mL of blood clots were removed and resutting was done. All bleeding points were identified and ligated. Dead space was obliterated with deep figure-of-eight sutures. Vaginal packing was done after complete haemostasis was secured. During surgery, the patient was transfused two units of blood and three units of fresh-frozen plasma (FFP) while the oxytocin infusion was continued. At the end of the operation, the patient was haemodynamically stable. In the recovery room after 1 hour her blood pressure dropped to 80/40 mmHg. Her heart rate was 110 bpm and oxygen saturation 99%. Examination revealed active bleeding from the vagina with a well-retracted uterus. A series of
haematological investigations were done, showing a haemoglobin concentration of 7.2 g/dL and a platelet count of 78 × 10⁹/L. Coagulation studies showed an international normalised ratio of 1.2 and activated partial thromboplastin time (APTT) of 30.1 seconds (normal 26.2 seconds). After much discussion, embolisation of the uterine artery was planned.

The patient was transferred to the cardiac catheterisation laboratory. An injection of 10 mL xylocaine (2%) was infiltrated and the left femoral artery was cannulated with a 6F sheath. Angiogram using ioxol contrast showed active bleeding from the left uterine artery, and no other bleeding point (Fig. 1). Embolisation was performed successfully with 500 μm polyvinyl alcohol (PVA) as shown in Fig. 2. A repeat angiogram showed no further bleeding (Fig. 3). The patient was transferred back to the intensive care unit and further blood products were transfused as required according to laboratory results.

Thirty-six hours after haematoma formation, the haematological parameters returned to acceptable limits (haemoglobin 9.6 g/dL, platelets 92 × 10⁹/L, INR 1.2, APTT 33 seconds). Chest X-ray showed no sign of acute lung injury. A total of six units of packed red cells and five units of FFP were transfused. The patient’s condition improved and she was discharged from hospital 8 days after her operation. At follow-up 4 weeks later, she remained well with no further complaints.

Discussion

Approximately 30% (in some countries over 50%) of direct maternal deaths worldwide are due to haemorrhage, mostly in the postpartum period.[11,12] In India, PPH is currently the most common direct cause of maternal death.

During pregnancy the uterus, vagina, and vulva have rich vascular supplies that are at risk of trauma during the birth process; trauma to these highly vascular areas may result in formation of a haematoma. Puerperal haematomas occur in 1:300 - 1:1500 deliveries and, rarely, do not lie in the superficial fascia. Trauma to these vessels can therefore lead to a large accumulation of blood in the paravaginal space or ischiorectal fossa, resulting in considerable delay in diagnosis.

Transcatheter arterial embolisation has been reported as a method of haemorrhage control since the 1960s, and is widely used in the control of pelvic haemorrhage due to malignancy, trauma and radiation. This technique has recently been successfully used in the control of PPH.[13] Advantages of this procedure are easy identification of the bleeding site, preservation of the uterus and fertility, while rebleeding from collateral vessels is rare. Complications occur in only 8.7% of cases, the most common being low-grade fever, and rarer ones including pelvic infection and groin haematoma.[13]

Conclusion

Our case shows that selective arterial embolisation of the uterine arteries is an effective method of treating puerperal haematoma refractory to conventional first-line therapy, and an effective alternative way of achieving haemostasis with minimal morbidity.