

Review

Secondary postpartum hemorrhage: a review of the literature

Galić Tihana, Brkić Maja, Dermit Kosjenka, Pleša Ivona, Orlović Marta, Krznarić Lovošević Ana Marija, Blagaić Vladimir

Department of Gynecology and Obstetrics, Clinical Hospital "Sveti Duh", Zagreb, Croatia

Biomedicine and Surgery

ABSTRACT

Postpartum hemorrhage (PPH) may occur shortly after delivery or, less often, days later. The most common causes of secondary PPH are retained products of conception, subinvolution of the placental bed, and/or infection. The etiology of secondary postpartum hemorrhage is diverse and management is dependent on identifying the cause and tailoring treatment appropriately. The main aims of treatment are to provide basic resuscitation, establish a cause for the bleeding, and tailor the treatment (medical and/or surgical) according to the cause. Sometimes the cause cannot be determined. Surgical procedures (dilation and curettage, suction curettage) are often effective when medical management fails.

KEYWORDS: puerperium; secondary postpartum hemorrhage

Correspondence to: Galić Tihana, Clinical hospital Sv.Duh, Sv.Duh 64, HR-10000 Zagreb, Croatia, e-mail: tihanagalico@gmail.com

Date received: August 8th 2017

Date accepted: September 16th 2017

INTRODUCTION

Secondary (also called late) PPH is any abnormal or excessive bleeding from the birth canal occurring between 24 hours and 12 weeks postnatal (1-3). Unlike primary postpartum hemorrhage, there is no clear definition for quantity of blood loss and this can vary from 'increased lochia' to massive hemorrhage (4). In developed countries, two per cent of postnatal women are admitted to hospital with this condition, half of them undergoing uterine surgical evacuation; in developing countries it is a major contributor to maternal death (3,5,6). Most studies report peak incidence is at one to two weeks postpartum (3). While primary PPH is an acute condition requiring immediate management, the bleeding in secondary PPH is usually not so severe. The patient may complain of spotting on and off for days after her delivery with an occasional gush of fresh blood.

ETIOLOGY

It is usually caused by retained products of conception, subinvolution of the placental bed, and/

or infection, remaining in the uterus and causing infection or preventing the uterus from contracting (3). Rare causes include: pseudoaneurysm of the uterine artery, arteriovenous malformations, choriocarcinoma, undiagnosed carcinoma of the cervix, adenomyosis, infected polyp or submucosal fibroid, inherited or acquired bleeding diatheses, uterine diverticulum, excessive bleeding with resumption of menses, hypoestrogenism, dehiscence of a cesarean scar (1,3,7-18). The cause cannot be determined sometimes.

MANAGEMENT OF SECONDARY POSTPARTUM HEMORRHAGE

Resuscitation

For the patient who is hemodynamically unstable, aiming the immediate hemodynamic stabilization of the patient is the priority. Often, blood and plasma unit transfusion is required. Blood and blood products should be given according to blood loss, rather than waiting and using the response to initial fluid administration and hemoglobin and

DOI: 10.5281/zenodo.1219223

This is an open access article under the CC BY-NC-ND license (http://creativecommons.org/licenses/by-nc-nd/4.0/).



coagulation results as the trigger for the infusion of blood (19,20). Treatment usually falls into one of two options: surgical evacuation of the uterine cavity or medical treatment (4).

CLINICAL PRESENTATION

The amount of blood loss at presentation varies but most are hemodynamically stable. A thorough history will provide information relating to cause and should include details regarding parity, labor, mode of delivery, third-stage or puerperal complications and any relevant medical and family history. Clinical findings are nonspecific. Clinical signs and symptoms at the time of presentation may include offensive lochia, abdominal cramping, uterine tenderness, pyrexia, enlarged uterus and an open cervical os (20).

Our initial approach to management is based on the suspected cause of bleeding. No data from randomized trials are available to guide management (5).

RETAINED PRODUCTS OF CONCEPTION

Vascularity of echogenic intracavitary material is a key finding as vascularity on color Doppler suggests retained products. If no intracavitary mass, endometrial fluid, or vascularity is seen and the endometrial thickness is thin, retained products are not likely (21).

Examination under anesthetic and surgical evacuation of the uterus should be considered if retained placental tissue is suspected clinically or after ultrasound examination. This has good reported success rates, with bleeding stopping promptly in all 72 women undergoing evacuation of the uterus for secondary postpartum hemorrhage in one study, despite only 36% having proven histological evidence of retained tissue (22).

SUBINVOLUTION OF THE PLACENTAL SITE

Initial approach for suspected subinvolution of the placental site, is administration of uterotonic agents. According to "Up to date" guidelines options include methylergonovine (0.2 mg intramuscularly, repeated every two to four hours up to three doses), or intramuscular carboprost tromethamine (250 mg intramuscularly; up to eight doses at intervals at least 15 minutes apart), and/or oxytocin infusion. If uterus if firm, these agents will not be useful (1).

When medical management is unsuccessful, surgical procedures (dilation and curettage, suction curettage) are often effective (even if retained placental or membrane fragments cannot

be identified sonographically) (4,20). A study of 132 consecutive women with secondary PPH reported 75 (57%) were initially treated with surgical evacuation, which was successful in 67 (90%) (5). Of the 57 women initially managed medically, treatment was successful in 41 (72%); 16 women had continuing symptoms, of whom 12 subsequently underwent surgical evacuation. Tissue specimens were obtained at surgery in only 38 women, and just one-third of these had histological confirmation of placental tissue. The histologic diagnosis of placental subinvolution is based on dilated myometrial arteries with hyaline material replacing the medial layer, partial occlusion by thrombi of variable age, and extravillous trophoblast in and around the placental bed vessels (1,22,23).

In high-risk patients, who can be refractory to uterotonic drugs or uterine curettage, selective arterial embolization has been effective for controlling severe bleeding (2,24,25). If percutaneous therapy fails, hysterectomy may be required (1).

ENDOMETRITIS

Endometritis is more common following prolonged rupture of membranes, prolonged labor, emergency Cesarean section or with a retained placenta requiring manual removal (20). Often, the first symptoms are lower abdominal pain and uterine tenderness, followed by fever; most commonly within the first 24 to 72 h postpartum. Chills, headache, malaise, and anorexia are common. Sometimes the only symptom is a low-grade fever. Pallor, tachycardia, and leukocytosis usually occur, and the uterus is soft, large, and tender. Discharge may be decreased or profuse and malodorous, with or without blood. Under these circumstances, we prescribe broad-spectrum antibiotic therapy. However, some clinicians administer antibiotics to all patients with secondary PPH, including those without obvious signs of infection. Rare, but potentially lethal causes of endometritis include Clostridium sordellii (25-28), Clostridium perfringens (29) and streptococcal or staphylococcal toxic shock syndrome (30-33).

CONCLUSION

Our initial approach to management secondary postpartum hemorrhage is based on the suspected cause of bleeding. Whether to initially manage secondary PPH medically or surgically is still a relatively unstudied aspect of the care of these patients.



There is no randomized controlled trials to inform management of women with secondary postpartum hemorrhage. This problem deserves more attention than it has received in recent years.

REFERENCES

- 1. Belfort MA. Secondary (late) postpartum hemorrhage. Available from: https://www.uptodate.com/contents/secondary-late-postpartum-hemorrhage?source=contentShare&csi=2ab18ef1-37eb-4e16-a5e4-3652600bc911 [Accessed 15th March 2017].
- American College of Obstetricians and Gynecologists. ACOG Practice Bulletin: Clinical Management Guidelines for Obstetrician-Gynecologists. Obstet Gynecol. 2006;108(4):1039-1047.
- 3. Dossou M, Debost-Legrand A, Déchelotte P. Severe secondary postpartum hemorrhage: a historical cohort. Birth. 2015;42(2):149-155. doi: 10.1111/birt.12164.
- Neill AC, Thornton S. Secondary postpartum hemorrhage. J Obstet Gynaecol. 2002;22:119-122.
- 5. Alexander J, Thomas P, Sanghera J. Treatments for secondary postpartum hemorrhage. Cochrane Database Syst Rev. 2002;(1):CD002867. doi: 10.1002/14651858.CD002867.
- Hoveyda F, MacKenzie IZ. Secondary postpartum haemorrhage: incidence, morbidity and current management. BJOG. 2001;108(9):927-930.
- Marshall AL, Durani U, Bartley A. The impact of postpartum hemorrhage on hospital length of stay and inpatient mortality: a National Inpatient Sample-based analysis. Am J Obstet Gynecol. 2017;217:344.e1-344.e6. doi: 10.1016/j.ajog.2017.05.004.
- 8. Nanjundan P, Rohilla M, Raveendran A. Pseudoaneurysm of uterine artery: a rare cause of secondary postpartum hemorrhage, managed with uterine artery embolisation. J Clin Imaging Sci. 2011;1:14. doi: 10.4103/2156-7514.76692.
- 9. Yun SY, Lee DH, Cho KH. Delayed postpartum hemorrhage resulting from uterine artery pseudoaneurysm rupture. J Emerg Med. 2012;42(1):e11-14. doi: 10.1016/j.jemermed.2011.03.005.
- 10. Hayata E, Matsuda H, Furuya K. Rare case of postpartum hemorrhage caused by rupture of a uterine artery pseudoaneurysm 3 months after Cesarean delivery. Ultrasound Obstet Gynecol. 2010;35(5):621-623. doi: 10.1002/uog.7607.
- 11. Marnela K, Saarelainen S, Palomäki O, Kirkinen P. Sonographic diagnosis of postpartum pseudoaneurysms of the uterine artery: a report of 2 cases. J Clin Ultrasound. 2010;38(4):205-208. doi: 10.1002/jcu.20658.
- 12. Lausman AY, Ellis CA, Beecroft JR. A rare etiology of delayed postpartum hemorrhage. J Obstet Gynaecol Can. 2008;30(3):239-243. doi:10.1016/S1701-2163(16)32760-8.
- 13. Aziz N, Lenzi TA, Jeffrey RB Jr, Lyell DJ. Postpartum uterine arteriovenous fistula. Obstet Gynecol. 2004;103(5Pt2):1076-1078. doi:10.1097/01.AOG.0000123241.44401.01.
- 14. Yi SW, Ahn JH. Secondary postpartum hemorrhage due to a pseudoaneurysm rupture at the fundal area of the uterus: a case treated with selective uterine arterial embolization. Fertil Steril. 2010;93(6):2048-2049. doi: 10.1016/j.fertnstert.2009.03.0 99.
- Gürses C, Yilmaz S, Biyikli S. Uterine artery pseudoaneurysm: unusual cause of delayed postpartum hemorrhage. J Clin Ultrasound. 2008;36(3):189-191. doi: 10.1002/jcu.20372.
- 16. Wang PH, Pang YP, Chao HT. Delayed postpartum hemorrhage in adenomyosis: a case report. Zhonghua Yi Xue Za Zhi (Taipei). 1998;61:492.

- Wu MC, Hsu YP, Lin HH, Hsiao SM. Severe delayed postpartum hemorrhage due to a neglected uterine diverticulum: a case report. J Reprod Med. 2013;58(7-8):347-350.
- Zubor P, Kajo K, Dokus K, Krivus S, Straka L, Bodova KB, Danko J. Recurrent secondary postpartum hemorrhages due to placental site vessel subinvolution and local uterine tissue coagulopathy. BMC Pregnancy Childbirth. 2014;14:80. doi: 10.1186/1471-2393-14-80.
- Groom KM, Jacobson TZ. A comprehensive textbook of postpartum hemorrhage: an essential clinical reference for effective management. 2nd ed. London: Sapiens Publishing; 2012.
- Mulic-Lutvica A, Axelsson O. Ultrasound finding of an echogenic mass in women with secondary postpartum hemorrhage is associated with retained placental tissue. Ultrasound Obstet Gynecol. 2006;28(3):312-319. doi: 10.1002/ uog.2849.
- 21. King PA, Duthie SJ, Dong ZG, Ma HK. Secondary postpartum haemorrhage. Aust NZ J Obstet Gynaecol. 1989;29:394.
- Ober WB, Grady HG. Subinvolution of the placental site. Bull NY Acad Med. 1961;37:713.
- 23. Kavalar R, Arko D, Fokter Dovnik N, Takač I. Subinvolution of placental bed vessels: case report and review of the literature. Wien Klin Wochenschr. 2012;124(19-20):725-730. doi: 10.1007/s00508-012-0219-9.
- Pelage JP, Soyer P, Repiquet D. Secondary postpartum hemorrhage: treatment with selective arterial embolization. Radiology. 1999;212:385.
- 25. Park HS, Shin JH, Yoon HK, Kim JH, Gwon DI, Ko GY, Sung KB. Transcatheter arterial embolization for secondary postpartum hemorrhage: outcome in 52 patients at a single tertiary referral center. J Vasc Interv Radiol. 2014;25(11):1751-1757. doi: 10.1016/j.jvir.2014.05.009.
- 26. Hollier LM, Scott LL, Murphree SS, Wendel GD Jr. Postpartum endometritis caused by herpes simplex virus. Obstet Gynecol. 1997;89:836.
- 27. Rørbye C, Petersen IS, Nilas L. Postpartum Clostridium sordellii infection associated with fatal toxic shock syndrome. Acta Obstet Gynecol Scand. 2000;79:1134-1135.
- 28. Bitti A, Mastrantonio P, Spigaglia P. A fatal postpartum Clostridium sordellii associated toxic shock syndrome. J Clin Pathol. 1997;50:259.
- 29. Aldape MJ, Bryant AE, Stevens DL. Clostridium sordellii infection: epidemiology, clinical findings, and current perspectives on diagnosis and treatment. Clin Infect Dis. 2006;43(11):1436-1446. doi: 10.1086/508866.
- 30. Cohen AL, Bhatnagar J, Reagan S, Zane SB, D'Angeli MA, Fischer M, Killgore G, Kwan-Gett TS, Blossom DB, Shieh WJ, Guarner J, Jernigan J, Duchin JS, Zaki SR, McDonald LC. Toxic shock associated with Clostridium sordellii and Clostridium perfringens after medical and spontaneous abortion. Obstet Gynecol. 2007;110(5):1027-1033. doi: 10.1097/01. AOG.0000287291.19230.ba.
- 31. Jorup-Rönström C, Hofling M, Lundberg C, Holm S. Streptococcal toxic shock syndrome in a postpartum woman. Case report and review of the literature. Infection. 1996;24:164.
- Gibney RT, Moore A, Muldowney FP. Toxic-shock syndrome associated with post-partum staphylococcal endometritis. Ir Med J. 1983;76:90.
- 33. Gibbs RS, Blanco JD. Streptococcal infections in pregnancy. A study of 48 bacteremias. Am J Obstet Gynecol. 1981;140:405.