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Analysis of Patients Undergoing Peripartum Hysterectomy for Obstetric Causes According to Delivery Methods: 13-Year Experience of a Tertiary Center

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Abstract

Objective: The aim of this study was to examine the maternal and fetal outcomes of patients undergoing peripartum hysterectomy (PH) after vaginal delivery (VD) and cesarean section (C/S).

Methods: The files of patients undergoing PH following postpartum hemorrhage (PPH) between January 2005 and November 2018 were reviewed retrospectively. Patients undergoing PH were divided into two groups as C/S and VD. Age, parity, gestational weeks, time between delivery and hysterectomy, estimated blood loss, duration of operation, number of blood transfusions, hospitalization time, APGAR scores of the fetus at the 1st and 5th minutes, previous C/S histories, fetal and maternal mortality, indications for PH, additional surgeries performed during PH, and pre-op and post-op complications were recorded retrospectively and the groups were compared.

Results: A total of 147 patients who underwent PH for postpartum PPH were identified. Of the patients included in the study, 77 underwent PH after VD and 70 underwent PH after C/S. There was no statistically significant difference between the groups in terms of age, parity, time between delivery and hysterectomy, estimated blood loss, number of blood transfusions, hospitalisation time, and maternal mortality rates. The gestational weeks of the patients in the VD group were higher than that of the patients in the C/S group ($P = 0.003$). Mean duration of operation of the C/S group was longer than that of the VD group ($P < 0.001$). APGAR scores of the fetus at the 1st and 5th minutes were higher in the VD group compared to the C/S group ($P < 0.001$, $P < 0.001$, respectively). The most common indication for PH was uterine atony in the VD group ($n: 54, 70.1\%$) and uterine rupture in the C/S group ($n: 24, 34.2\%$). Disseminated intravascular coagulopathy (DIC) was the most common complication in both groups.

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Conclusion: While fetal mortality and morbidity are higher in patients undergoing hysterectomy after C/S, long-term effects caused by C/S (previous C/S, placenta accreta, placenta previa) increase PH risk. However, it should also be considered that PH risk may increase after VD as well.

Keywords: Peripartum hysterectomy, cesarean section, vaginal delivery, postpartum hemorrhage

Obstetrik Nedenlerle Peripartum Histerektomi Yapılan Hastaların Doğum Şekillerine Göre Analizi: Tersiyer Merkezin 13 Yıllık Deneyimi

Öz

Amaç: Bu çalışmada vajinal doğum (VD) sonrası ve sezaryen doğum (C/S) sonrası peripartum histerektomi (PH) uygulanan hastaların maternal ve fetal sonuçlarını incelemek amaçlandı.

Yöntemler: Ocak 2005 ile Kasım 2018 tarihleri arasında tersiyer bir merkezde postpartum kanama (PPK) sonrası PH olan hastaların dosyaları retrospektif olarak incelendi. PH olan hastalar C/S sonrası ve VD sonrası olmak üzere iki gruba ayrıldı. Tüm hastaların yaşları, parite sayıları, gebelik haftaları, doğum ile histerektomi arası geçen süreleri, tahmini kan kayıpları, operasyon süreleri, kan transfüzyonu sayıları, hastanede yatış süreleri, fetüsün 1.ve 5.dakika APGAR skorları, geçirilmiş C/S öyküleri, fetal ölüm ve maternal ölüm durumları, PH endikasyonları, PH operasyonu sırasında yapılan ek cerrahiler, cerrahi sırasında veya sonrasında olan komplikasyonlar retrospektif olarak kaydedildi ve gruplar birbiri ile karşılaştırıldı.

Bulgular: Doğum sonrası PPK nedeni ile PH yapılan 147 hasta tespit edildi. Çalışmaya dahil edilen hastaların 77'sine VD sonrası ve 70'ine C/S sonrası PH uygulandı. Grupların yaş, parite, doğum ile histerektomi arası geçen süreleri, tahmini kan kayıpları, yapılan kan transfüzyonu sayıları, hastanede yatış süreleri, maternal ölüm oranları arasında istatistiksel olarak fark izlenmedi. VD grubundaki hastaların gestasyonel haftaları, C/S grubundaki hastaların gestasyonel haftalarına oranla daha yüksekti (p:0.003). C/S grubunun operasyon süreleri VD grubuna göre daha uzundu (p <0.001). VD grubundaki hastaların bebeklerinin 1.ve 5.dakika APGAR skorları, C/S grubuna oranla daha yüksekti (sırasıyla p <0.001, p<0.001). VD grubundaki hastaların en sık (n:54, %70,1) uterin atoni nedeniyle, C/S grubundaki hastaların ise en sık (n:24, %34,2) uterin rüptür nedeniyle PH olduğu görüldü. Her iki grupta da en sık görülen komplikasyonun ise dissemine intravasküler koagülopati (DİK) olduğu tespit edildi.

Sonuç: C/S sonrası histerektomi olan hastalarda fetal mortalite ve morbidite daha fazla iken, C/S operasyonun neden olduğu uzun dönemli sonuçlar (geçirilmiş C/S, plasenta akreata, plasenta previa) PH riskini artırmaktadır. Ancak VD sonrası da PH riskinin artabileceği göz önünde bulundurulmalıdır.

Anahtar kelimeler: Peripartum histerektomi, sezaryen, vajinal doğum, postpartum hemoraji.

INTRODUCTION

Postpartum hemorrhage (PPH) is a potentially preventable obstetric emergency occurring after both vaginal delivery (VD) and cesarean section (C/S). PPH is the main cause of maternal mortality worldwide^{1,2}. Maternal mortality due to PPH vary between 1% and 5% in all deliveries³. Since PPH is preventable, accurate and early diagnosis is crucial to prevent maternal mortality. PPH is defined as blood loss of more than 500 ml in the first 24 hours after VD, or more than 1000 ml after C/S⁴.

Peripartum hysterectomy (PH) is usually performed to prevent maternal mortality in life-threatening obstetric hemorrhage cases and is therefore considered as "near-miss"⁵. PH is generally used as a life-saving treatment of massive bleeding when other medical or conservative surgical treatments fail. Its incidence is 0.4 per 1000 births in developed countries, while in underdeveloped countries the incidence is⁵ per 1000 births⁶. Known risk factors for PH are advanced maternal age, abnormal placentation, high parity number, and C/S history in current or previous

pregnancies^{7,8}. While uterine atony and uterine rupture were the most common causes of PH in previous years, these rates started to decrease with intrapartum and postpartum follow-up methods and increasing C/S rates, placental invasion anomalies, and placenta previa became the most common cause of PH in recent years⁹⁻¹¹.

The aim of this study was to compare the indications, postoperative or intraoperative complications, and maternal and fetal outcomes of patients who underwent PH after VD and C/S.

METHODS

The files of patients who underwent PH due to PPH in a tertiary center between January 2005 and November 2018 were reviewed retrospectively. Ethics committee approval for the study (ethics committee number: 182) was obtained from our hospital. Patients who underwent hysterectomy for gynecological indications, patients referred to our clinic due to PPH after VD or C/S in an external center, patients whose gestational week was less than 24 weeks, and patients with incomplete or insufficient hospital records were excluded from the study. Patients with PPH within 24 hours after VD or C/S who underwent PH after failure to control bleeding with medical treatment (oxytocin, methergine, misoprostol) and surgical treatment (uterine fundal massage, postpartum uterine curettage, Bakri balloon, b-Lynch, Hayman suture, etc.) were included in the study. Patients undergoing PH were divided into two groups as post C/S and post VD. Age, parity, gestational weeks, time between delivery and hysterectomy, estimated blood loss, duration of operation, number of blood transfusions, hospitalization time, APGAR scores of the fetus at the 1st and 5th minutes, previous C/S histories, fetal and maternal mortality, indications for PH, additional surgeries performed during PH, and complications during or after surgery were recorded retrospectively from patient files and

the groups were compared. The gestational week of the patients was calculated based on the last menstrual date (LMD). For patients who did not remember their LMDs, first trimester ultrasonography recordings were used to calculate the gestational week. Hospitalisation time was defined as the time between the day of delivery and the day of discharge. Patients receiving blood transfusions were those who periodically lost more than 20% of their total blood volume (patients with postoperative hemoglobin values below 7 g/dl or patients with symptoms of anemia).

Statistical Package for Social Sciences 20.0 (SPSS Inc., Chicago, IL, USA) was used for statistical analysis. The distribution of the data was evaluated by Kolmogorov-Smirnov test. Descriptive statistical methods (mean, standard deviation) were used. Pearson chi-square test was used for comparison of groups. A p value of < 0.05 was accepted as statistically significant.

RESULTS

Between January 2005 and November 2018, a total of 289,793 deliveries were performed in our center, including 218,437 VD and 71,356 C/S deliveries. There were 147 patients who underwent PH after delivery. Of the patients included in the study, 77 underwent PH after VD and 70 underwent PH after C/S. Table I lists the demographic data and operational characteristics of the groups. There was no statistically significant difference between the groups in terms of age, parity, time between delivery and hysterectomy, estimated blood loss, number of blood transfusions, hospitalisation time, and maternal mortality rates. The gestational weeks of the patients in the VD group were higher than that of the patients in the C/S group ($P = 0.003$). Mean duration of operation of the C/S group was longer than that of the VD group ($P < 0.001$). APGAR scores of the fetus at the 1st and 5th minutes were higher in the VD group compared

to the C/S group (P <0.001, P <0.001, respectively). The rate of fetal mortality and hypogastric artery ligation (HGAL) during PH were significantly higher in the C/S group.

Table 1: Demographic Data and Operation Characteristics of Groups

Characteristics	VD (n=77)	C/S (n=70)	p-value
Age	33.7±5.6	34.1±6.1	0.70
Parity	4.8±2.7	4.6±2.3	0.68
Gestational Week	38.5±2.6	36.8±3.8	0.003
Time between delivery and hysterectomy (min)	158±138	105±108	0.12
Estimated blood loss (ml)	2603±844	2571±1009	0.83
Duration of operation (min)	105.1±44.8	135.5±55.2	<0.001
Blood Transfusion	5.1±2.1	5.2±1.9	0.71
1st Minute APGAR score	5.6±2.4	3.6±2.7	<0.001
5th Minute APGAR score	7.9±2.5	5.7±3.5	<0.001
Hospitalization (days)	5.5±2.6	5.3±2.4	0.58
HGAL	Yes 13/77 (16.9%)	22/70 (31.4%)	0.03
Previous caesarean section	Yes 0/77 (0%)	24/70 (34.3%)	<0.001
Fetal mortality	Yes 6/77 (7.8%)	21/70 (30%)	0.001
Maternal mortality	Yes 3/77(3.9%)	1/70 (1.4%)	0.06

HGAL: Hypogastric artery ligation

Table II lists the indications for PH, additional surgeries during the operation, and complications. The most common indication for PH was uterine atony in the VD group (n: 54, 70.1%) followed by uterine rupture (n: 23, 29.8%), and uterine rupture in the C/S group (n: 24, 34.2%) followed by uterine atony (n: 23, 32.8%). While the majority of patients with VD and C/S delivery did not undergo any additional surgery during PH (76.6% and 67.1%, respectively), HGAL was the most commonly used procedure (16.8% and 31.4%,

respectively). Complications were not seen in the majority of both groups after PH. Disseminated intravascular coagulopathy (DIC) was the most common complication in both groups.

Table 2: PH Indications, Additional Surgical Operations, and Postoperative Complications

Characteristics		VD (n=77)	C/S (n=70)
Indication for Hysterectomy	Atony	54 (70.1%)	23 (32.8%)
	Uterine rupture	23 (29.8%)	24 (34.2%)
	Placenta previa	-	2 (2.8%)
	Placenta accrete	-	14 (20%)
	Placental abruption	-	5 (7.1%)
	Incision site bleeding	-	2(2.8%)
	Additional surgeries	None	59 (76.6%)
USO		1 (1.2%)	-
HGAL+USO		3 (3.8%)	1 (1.4%)
HGAL+BSO		1 (1.2%)	-
Complications	None	39 (50.6%)	23 (32.8%)
	Bladder injury	1 (1.2%)	9 (12.8%)
	DIC	11 (14.2%)	11 (15.7%)
	Fever	5 (6.4%)	7 (10%)
	Wound site Infection	10 (12.9%)	9 (12.8%)
	ARF	-	3 (4.2%)
	PTE	4 (5.1%)	1 (1.4%)
	Ureteral injury	2 (2.5%)	2 (2.8%)
	Ileus	5 (6.4%)	5 (7.1%)

ARF: acute renal failure USO: unilateral salpingo-oophorectomy, BSO: bilateral salpingo-oophorectomy PTE: pulmonary thromboembolism DIC: Disseminated intravascular coagulopathy

DISCUSSION

In this study, maternal and fetal outcomes, demographic data, and operation characteristics of patients undergoing PH after VD and C/S were compared. While the VD group had higher APGAR scores at the 1st and 5th minutes, fetal mortality rate and HGAL during PH were higher in the C/S group. The most common indication for PH was uterine atony in the VD group (n: 54, 70.1%) and uterine rupture in the C/S group (n: 24, 34.2%). In addition, fetal mortality and morbidity were higher in the C/S group.

PH is the last life-saving treatment for PPH cases that cannot be controlled by medical or surgical treatment¹². It can be applied in PPHs after C/S (cesarean hysterectomy) or VD (postpartum hysterectomy). Although the rate of PH varies by country, it has been reported in rates ranging from 0.2 to 2.7 per 1000 births¹³⁻¹⁵. While the rates in Europe range from 0.2 to 1 per 1000 births¹⁶, studies in Turkey report rates of 4.68/1000 and 5.09/1000^{14,17}. In our study, the rate of PH in the specified time interval was found to be 5 in 10,000 births (147 in 289,793 births). This rate is lower than those reported in the literature. This low rate can be attributed to several reasons: Since our clinic is a tertiary center and hospital conditions are favourable, PH is not preferred initially for patients with PPH and procedures such as Bakri balloon, B-lynch, Hayman suture, HGAL, uterine artery ligation, and segmental resection are applied liberally. Therefore, PH rates may be lower because many patients benefit from the procedures without having to undergo PH.

In the present study, the most common cause of PH after VD was uterine atony with 70.1% and the most common cause of PH after C/S was uterine rupture with 34.2%. It has been reported in the literature that PH rates due to uterine atony indication are decreasing, whereas PH rates due to placental accreta are increasing¹⁸. The most important reason for

this has been reported to be the increasing C/S ratios in recent years¹⁸. These indication differences may be due to differences in C/S rates and approaches to obstetric hemorrhage in various countries. It is also remarkable that 29.8% of VD patients underwent PH due to uterine rupture. In a study conducted in Turkey, it was reported that 69.2% of patients with unscarred uterine rupture after VD underwent PH¹⁹. The low rate in the present study may be due to the difference in demographics data between the studies.

The most important point that should be kept in mind in patients undergoing PH is the preparation of necessary and sufficient blood products especially in patients with C/S history with placental previa or placental invasion anomaly since there may be excessive blood loss due to obstetric bleeding¹⁹. Studies have reported that patients undergoing PH may require an average of 4-6 units of erythrocyte suspension (ES)²⁰. In the present study, no statistically significant difference was observed in the number of blood transfusions between the two groups (P = 0.71). However, in the VD and C/S groups, 5.1 ± 2.1 and 5.2 ± 1.9 units of ES transfusion were performed, respectively. This rate is consistent with the literature. In addition, HGAL was performed in 31.4% of the C/S group and 16.8% of the VD group to control bleeding during PH. This ratio is higher in the C/S group since indications for PH include placenta previa (2.8%), placenta accreta (20%) and placental abruption (7.1%), and therefore the need for HGAL may have arisen.

PH can cause many complications, such as increased number of massive transfusions, DIC, urinary tract injury, and febrile morbidity^{4,21}. In the present study, the most common complication in the C/S group was DIC, while the second most common complication was bladder injury. In the VD group, the most common complication was again DIC. DIC secondary to excessive blood loss as a result of

obstetric bleeding is an expected condition. In addition, due to the fact that our center is a referral hospital, the time taken during the transfer of patients to our clinic may have increased the blood loss and thus led to DIC being the most common complication.

In the present study, none of the patients in the VD group had a history of C/S. In the C/S group, 34% of the patients had a previous C/S history ($P < 0.001$). C/S history may have an effect on the higher rate of bladder injury in the C/S group. In the literature, the rate of bladder injury during PH varies between 9-15% 4,14. The rate of bladder injury in the present study was 12.8% in the C/S group and 1.2% in the VD group. The rate of bladder injury in the C/S group is consistent with the literature and the probable cause is adhesions in the vesicouterine pouch due to C/S.

In a study conducted in Turkey, maternal mortality rate was reported as 16.7% 14. Maternal mortality rate was reported as 0.6% in the UK 11, and 23 821% in Nigeria²². In the present study, 3 cases (3.9%) in the VD group and 1 case (1.4%) in the C/S group resulted in maternal mortality and this was not associated with the mode of delivery ($P = 0.06$). However, it should be kept in mind that maternal mortality rates are high in obstetric bleeding. These rates may be associated with differences in geography, low antenatal follow-up, differences in socio-economic level, and time taken during the transfer of patients to advanced centers.

When the fetal outcomes of the groups were examined, APGAR scores at the 1st and 5th minutes were lower ($P < 0.001$) and fetal mortality rate (30%) was significantly higher in the C/S group ($P = 0.001$). To the best of our knowledge, there are no publications in the literature comparing fetal outcomes with PH. The low APGAR scores in the C/S group in our study and the high fetal mortality rate may be due to the higher rate of uterine rupture in the

C/S group compared to the VD group. In addition, while patients with placenta previa, placenta accreta, and previous C/S history were non-existent in the VD group, the presence of these patients only in the C/S group may have increased fetal mortality and resulted in a low APGAR score. In addition, the significantly higher duration of operation in the C/S group may have an effect on the delivery time of the fetus, as previous C/Ss may increase the duration of the operation, resulting in low APGAR scores and increased fetal morbidity. Another reason was that gestational week in the C/S group was significantly lower compared to the VD group. This may have increased fetal morbidity and mortality.

Retrospective nature of this study is a limitation. Prospective studies with broader patient groups can be performed. However, the high number of patients and the evaluation of a 13-year period are the strengths of this study.

CONCLUSION

While fetal mortality and morbidity are higher in patients undergoing hysterectomy after C/S, long-term effects caused by C/S (previous C/S, placenta accreta, placenta previa) increase PH risk. However, it should also be considered that PH risk may increase after VD as well.

Ethics Committee Approval: Ethics committee approval for the study (ethics committee number: 182) was obtained from our hospital.

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