INTRODUCTION

Placenta accreta as described by Irving et al, is “failure of separation of the placenta from the uterine wall following delivery of the human fetus leading to the often-used term morbid placental adherence”.1 The condition is characterised by placental invasion thereby leading to torrential haemorrhage. It has been proposed that the terminology, placenta accreta spectrum (PAS), including accreta, increta, and percreta be used hence forth.2

The condition has not yet been detected in any other animal in the literature till date.3

The incidence of PAS has increased substantially from 0.8 per 1000 deliveries in the 1980s to 3 per 1000 deliveries in the past decade, a phenomenon may be because of a rising caesarean section rate worldwide.4 PAS is associated with significant maternal morbidity and mortality, majorly including obstetric haemorrhage and obstetric hysterectomy.5 Mortality rates of up to 7% associated with PAS are reported.6 The most recent confidential inquiry into maternal mortality in the United Kingdom (MMBRACE-UK, 2017) highlighted the continued high maternal mortality associated with PAS.7

The most important antenatal risk factor for PAS is the number of previous caesarean sections. In the presence of low-lying placenta (placenta previa) and three previous caesarean sections, it is estimated that a woman would have a 61% risk of PAS.8 Antenatal diagnosis is a key element in order to improve maternal and perinatal outcome. Despite ultrasound and MRI having improved antenatal diagnosis, between one half and two thirds of cases remain undiagnosed, resulting in poorer maternal outcomes.9,10
The risk factors for placenta previa are smoking, previous caesarean sections, advanced maternal age, multiparity and conception by in vitro fertilization (IVF).\textsuperscript{11}

Abortions are mainly associated with foetal pathology, congenital abnormality, low birth weight and preterm labour in subsequent pregnancies.\textsuperscript{12}

Primary and secondary uterine pathologies which have been associated with PAS include direct surgical scar including caesarean delivery, surgical termination of pregnancy, dilatation and curettage, myomectomy, endometrial resection, Asherman’s syndrome and nonsurgical scar including IVF procedures, uterine artery embolization, chemotherapy and radiation, endometritis intra-uterine device, manual removal of placenta and previous accrete.

Certain uterine anomalies associated include bicornuate uterus, adenomyosis, submucous fibroids and myotonic dystrophy.

**METHODS**

This was a cross sectional observational study done from June 2017 to June 2019 at obstetrics and gynaecology department at a tertiary care centre, Mumbai. All the antenatal patients who visited the tertiary care centre underwent ultrasonography with placental localization.

Study design - cross sectional observational study.

**Inclusion criteria**

- ANC of all age groups who were diagnosed on ultrasonography to be placenta accreta spectrum and were confirmed intra operatively.

**Exclusion criteria**

- The cases with a normally located placenta were excluded.

A detailed history and examination of the patients was done. The patients were kept for close follow up and MRI was done in some selected cases. A cross sectional study was done to analyse the most common risk factors contributing to the etiology of the placental accreta syndrome. In cases of past surgeries like check curettage, previous caesarean sections, the incidence was seen and compared with patients not having any such history and hence the role of surgeries as an etiology factor was assessed.

**RESULTS**

Placenta accreta spectrum is a major cause of maternal morbidity and mortality. It is one of the leading causes of obstetric haemorrhage. A total of 19 cases were studied.

**Age distribution**

The results support the fact that the incidence increases as the age advances. Most of the cases were found in the age group > 35 years of age (52.64%).

The cases of PAS in the age group of 20-35 years were 47.36% and in women more than 35 years age group were 52.64%. The risk of placenta percreta and accrete increases as the parity increases and is associated with increased risk of caesarean sections. When compared with placenta previa cases, a higher incidence of placenta previa was found in the age group of 20-35 years (73.75%), whereas the placenta accrete syndrome had a higher incidence in age group > 35 years (52.64%).

| Age       | No. of cases (PAS) | Percentage (PAS) | No. of previa cases | Percentage of previa cases |
|-----------|--------------------|------------------|---------------------|----------------------------|
| < 20 years| 0                  | 0                | 3                   | 3.75                       |
| 20-35 years| 9                 | 47.36            | 59                  | 73.75                      |
| > 35 years| 10                 | 52.64            | 18                  | 22.5                       |
| Total     | 19                 | 100              | 80                  | 100                        |

**Previous caesarean sections**

The cases of PAS in previously unscarred uterus were 15.78%, in previous one LSCS were 57.89%, in previous 2 LSCS were 15.78% and in women with more than previous 2 LSCS were 10.55%.

A total 16 cases had history of transfusion of blood and blood products (94.73%).

Most of the cases were shifted under critical care unit for post-operative monitoring. The perinatal outcomes were satisfactory and only 2 cases with perinatal mortality.

**Indication of previous caesarean sections**

In case of previous caesarean sections, the commonest indication of previous caesarean sections included fetal distress (n = 3), failure of induction (n = 2), and previous caesarean with a short inter-conception period (n = 2).
The current study shows that the incidence of percreta/accreta increases as the rate of caesarean sections increases as has been shown by the studies done by Ananth CV et al, Faiz AS et al, Getahun D, Oyelese Y, et al.14-16 There were 5 cases of percreta and 14 cases of accrete.

Most studies have reported a dose related response pattern of risk factors of placenta previa with increasing number of previous caesarean sections.17 The previous caesarean delivery include polyp formation, lymphocyte infiltration, capillary dilatation, and infiltration of the endometrial tissue that surround the scar by free red blood cells , thereby increasing the chances of placenta accrete syndrome.18

CONCLUSION

The current study shows that the risk factors of placenta accreta and percreta to be advanced maternal age, previous delivery by C/S, bleeding during pregnancy and multiparity. history of uterine surgeries and previous caesarean are some important risk factors for accreta in placenta previa patients. A positive correlation was obtained between history of induced abortion/spontaneous abortion without check curettage and incidence of placenta accrete/percreta in this study. However, results of this study are not statistically conclusive because of small sample size and further studies are required.

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REFERENCES

1. Irving C, Hertig AT. A study of placenta accrete. Surg Gynecol Obstet. 1937;38(6):1088-200.
2. Jauniaux E, Chantraine F, Silver RM, Langhoff-Roos J. For the FIGO Placenta Accreta Diagnosis and Management Expert Consensus Panel. FIGO consensus guidelines on placenta accreta spectrum disorders: epidemiology. Int J Gynecol Obstet. 2018;140(3):265-73.
3. Chuong E. Evolutionary perspectives into placental biology and disease. Applied Translat Genom. 2013;2:64-9.
4. Higgins M, Monteith C, Foley M, and O’Herlihy C. Real increasing incidence of hysterectomy for placenta accreta following previous caesarean section. Eur J Obstet Gynecol Repro Biol. 2013;171(1):54-6.
5. Silver RM, Barbour KD. Placenta accreta spectrum: accreta, increta, and percreta. Obstet Gynecol Clin North Am. 2015;42(2):381-402.
6. O’Brien JM, Barton JR, Donaldson ES. The management of placenta percreta: conservative and operative strategies. Am J Obstet Gynecol. 1996;175(6):1632-8.
7. Knight M, Nair M, Tuffnell D, Shakespeare J, Kenyon S, Kurinczuk JJ. Saving lives, improving mothers’ care: lessons learned to inform maternity care from the UK and Ireland confidential enquiries into maternal deaths and morbidity 2013-15. National Perinat Epidemiol Unit. University of Oxford: Oxford. 2017:1-10.
8. Placenta Praevia, Placenta Praevia Accreta and Vasa Praevia: Diagnosis and Management (Green-top Guideline No. 27), Royal College of Obstetricians and Gynaecologists, 2011:1-26.
9. Bailit JL, Grobman W, Rice MM, Reddy UM, Wapner RJ, Varner MW, et al. Morbidly adherent placenta treatments and outcomes. Obstet Gynecol. 2015;125(3):683.
10. Fitzpatrick KE, Sellers S, Spark P, Kurinczuk JJ, Brocklehurst P, Knight M. The management and outcomes of placenta accreta, increta, and percreta in the UK: a population-based descriptive study. BJOG: Int J Obstet Gynaecol. 2014;121(1):62-71.
11. Shobeiri F, Jenabi E, Karami M, Karimi S. Determinants of placenta previa: a case-control study. Biomed Res Therapy. 2017;4:1411-9.
12. Kashanian M, Akbarian A, Baradaran H, Shabandoust S. Pregnancy outcome following a previous spontaneous abortion (miscarriage). Gynecol Obstet Invest. 2006;61:167-70.
13. Eniola AO, Bako AU, Selo-Ojeme DO. Risk factors for placenta previa in Southern Nigeria. East Afr Med J. 2002;79:535-8.
14. Wax JR, Seiler A, Horowitz S, Ingardia CJ. Interpregnancy interval as a risk factor for placenta accreta. Connecticut Med. 2000;64(11):659-61.
15. Mehboob R, Ahmed N. Fetal outcome in major placenta previa. J Med Res. 2003;42:372-38.
16. Francois K, Johnsen JM. Is placenta previa more common in multiple pregnancy? Am J Obstet Gynecol. 2002;79:535-8.
17. Mortensen JT, Thulstrup AM, Larsen H, Møller M, Sørensen HT. Smoking, fetal sex, and risk of placental abruption, placenta previa, and preeclampsia: a population-based cohort study. Acta Obstet Gynecol Scand. 2001;10:894-8.
18. Yang Q, Wen SW, Oppenheimer L, Chen XK, Black D, Gao J, et al. Association of caesarean delivery for first birth with placenta praevia and placental abruption in second pregnancy. BJOG: An Int J Obstet Gynaecol. 2007;114(5):609-13.

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