Maternal Outcome in Pregnanncies with Thrombocytopenia

Authors
Dr Rinku G¹, Dr. M I Geetha²
¹Assistant Professor, Dept. of O & G, SATH, Govt. Medical College, Trivandrum.
²Associate Professor, Dept. of O & G, SATH, Govt. Medical College, Trivandrum.

Abstract

**Background:** Thrombocytopenia is a common occurrence in pregnancy. Although pregnancy is associated with physiological changes in platelet count, several pathological conditions cause thrombocytopenia, which can have a significant impact on the mother and the baby. The present study aims at the maternal outcome of pregnancies in patients with thrombocytopenia during pregnancy.

**Materials and Methods:** Comparative cross sectional survey conducted in SAT Hospital for a period of one year.

**Results:** In our study gestational thrombocytopenia constitutes 40% of cases and majority are mild type. Moderate and severe types were constituted by Pre eclampsia 10.8%, HELLP syndrome 17.4%, ITP 11.4% SLE 9% and DIC especially following grade III abruption. Out of moderate to severe cases 13.8% developed post-partum hemorrhage requiring blood and blood products transfusion. Maternal mortality (1 case) occurred in SLE group due to intractable pulmonary edema. Cesarean section rate was more in thrombocytopenia group.

**Conclusion:** Gestational Thrombocytopenia group had a favorable outcome in our study. The other group constituted by pre eclampsia SLE, ITP, DIC had complications like PPH, abruptions, DIC requiring multi-disciplinary approach.

**Keywords:** Thrombocytopenia, ITP (immune thrombocytopenic purpura), SLE (systemic lupus erythematosus).

Introduction
A mild thrombocytopenia is relatively frequent during pregnancy and has no consequences for either the mother or the fetus. But it may result from a range of pathologic conditions requiring closer monitoring and therapy. It is defined as a platelet count <1.5 L per mm³ and is second only to anemia.

Incidental thrombocytopenia of pregnancy – gestational thrombocytopenia- accounts for 70-80% of cases and it occurs in the mid –second, third trimester of pregnancy¹. It may result from various mechanisms including hemodilution, decreased platelet production and accelerated clearance⁶ and diagnosis is only of exclusion. Counts from 1 L to 1.5 L per mm³ are mildly depressed, 50000 to 1L per mm³ moderately depressed and >50000 are severely depressed³. The most common cause is gestational thrombocytopenia. Hypertensive disorders
account for 21%. Thrombocytopenia occurs more commonly in patients with eclampsia (30%) than in patients with both mild and severe forms of pre-eclampsia (15-18%) of the patients who have severe pre-eclampsia 4-12% will manifest criteria of HELLP Syndrome.

Immune mediated thrombocytopenia including idiopathic thrombocytopenic purpurs is responsible for 4.1% of cases. Other less common causes include rheumatological diseases (SLE), DIC, TTP, APLA, HIV infection and medications. ITP is the most common cause of thrombocytopenia at less than 20 weeks gestation and aim of treatment is to increase platelet count and to ensure safe delivery. Maternal anti platelet antibodies can cross the placenta and studies indicate that 12-15% of infants born to these mothers develop severe thrombocytopenia.

TTP is a rare but life threatening disease that should be suspected with thrombocytopenia and microangiopathic hemolytic anemia and treatment with IVIG should begin immediately to prevent maternal death. Occasionally a previously undiagnosed congenital platelet disorder maybe recognized for the first time during pregnancy. At delivery placental separation occurs at a time when normal blood flow is approximately 700 ml per minute. This flow is dampened by uterine contraction leading to myometriyal compression and occlusion by physiologic thrombosis of the open maternal vessels. Defect in either mechanism to arrest uterine bleeding leads to potentially lethal haemorrhage. Therefore when thrombocytopenia is diagnosed in pregnancy the women should undergo further clinical and laboratory assessment to determine the cause.

**Objective**

**Primary Objective**
To study the maternal outcome in patients with thrombocytopenia and comparing with those without thrombocytopenia.

**Secondary Objective**
To compare the maternal outcome of different etiologies of thrombocytopenia.

**Materials and Method**

**Study Design-** Comparative cross sectional survey

Study population include all pregnant women with thrombocytopenia delivering in SAT Hospital, Govt. Medical College, Trivandrum for one year and comparing with equal number of pregnant patients without thrombocytopenia and hypertensive disorders who deliver during the same period.

Sample size is calculated by the formula:

\[ N = \frac{2pq(Z_\alpha + Z_\beta)^2}{(P_1 - P_0)^2} \]

where:
- \( N \) is the sample size
- \( P \) is the proportion
- \( Z_\alpha \) is the standard normal deviate for a given confidence level
- \( Z_\beta \) is the standard normal deviate for a given power
- \( P_0 \) is the control group response
- \( P_1 \) is the study group response

A= Type 1 error (fixed at 5% level)
1-\( \beta \)= Power (fixed at 80% level)
RR= Relative Risk

200 cases of thrombocytopenia patients and 200 consecutive pregnancies without thrombocytopenia and hypertension

The clinical details of all women were collected by reviewing their hospital records, labour records, and the cases will be analyzed according to the history and the medical records. Following clinical characteristics – maternal age, previous gestations, parity, gestational age, birth weight, cause of thrombocytopenia were evaluated.

Obstetrics risk factors- GDM, overtdiabetis, previous CS, maternal anemia, hydramnious, ologamnious, multiple pregnancies, thyroid disease were examined including the following.

- Requirement of Blood and platelet transfusion
- Placental abruption, Intra uterine death
- Labour induction, Mode of delivery
- PPH, episiotomy haematoma, anaesthetic complications

Statistical analysis done by Chi square test, odds ratio and logistical regression.
Ethical issues
Being comparative cross sectional survey, permission from hospital authorities is required for collecting information.

Results
Observation and Results
I. Age of the patients
Maximum patients (85 %) in the thrombocytopenia group belong to the age group 20-29 years. The controls too had maximum number in the same age group (89.8 %) but in the age group of 30-34 years, there were 20 patients in the case against 6 in the control.

| Age group | Case | Control |
|-----------|------|---------|
|           | Number | Percentage | Number | Percentage |
| <19       | 0      | 0        | 7      | 4.2        |
| 20-24     | 75     | 44.9     | 86     | 51.5       |
| 25-29     | 67     | 40.1     | 64     | 38.3       |
| 30-34     | 20     | 12       | 6      | 3.6        |
| ≥35       | 5      | 3        | 4      | 2.4        |
| Total     | 167    | 100      | 167    | 100        |

II. Gestational age of diagnosis of thrombocytopenia
76% of the cases of thrombocytopenia were diagnosed in the third trimester. ITP and SLE cases had thrombocytopenia presenting in the first and second trimester. Abruption which was an important cause of acute onset of thrombocytopenia also occurred more towards the third trimester.

III. Booked cases vs un booked cases
One patient among the control group was unbooked and all cases among the thrombocytopenia group were booked cases.

IV. Referred cases
59.9% of patients among the cases were referred to our hospital while only 18.6% of the control group were referred.

V. Parity
No significant difference noted in the case and control group (p value= 0.219). however, pre-eclampsia is more common in Primigravida.

| Referred | Case | Control | Total |
|----------|------|---------|-------|
|          | N    | %      | N     | %    | N     | %    |
| Primi    | 95   | 56.9   | 106   | 63.5 | 201   | 60.2 |
| Multi    | 72   | 43.1   | 61    | 36.5 | 133   | 39.8 |
| Total    | 167  | 100    | 167   | 100  | 334   | 100  |

VI. Gestational age at delivery
There was no significant difference among the two groups in terms of gestational age at delivery, majority of patients in both the arms delivered at 37-40 weeks.

VII. Severity of thrombocytopenia
Majority were mild cases of thrombocytopenia (55%). The moderate (33%) and (12 %) severe forms were constituted by ITP, SLE and DIC especially following cases of grade 3 abruption.
VIII. Causes of thrombocytopenia

There was overlap between cases of SLE, ITP and pre-eclampsia as some of these patients had all these disorders together. Abruption was the most important cause of DIC and associated thrombocytopenia. Some patients with pre-eclampsia also showed evidence of DIC.

IX. Gestational hypertension and its association with thrombocytopenia

27.1% of patients in the case group had gestational hypertension.

X. Thrombocytopenia and maternal risk factors

No significant difference was noted between the cases and the control group with respect to maternal risk factors like diabetes mellitus, maternal hypothyroidism, anaemia, oligohydramnios, polyhydramnios, multiple pregnancy and preterm labour.

XI. Abruption as a cause of thrombocytopenia

There were 20 cases of grade III abruption, majority of which required blood transfusion and ICU management for correction of hypotension, coagulation failure and renal compromise.

XII. Mode of delivery

There were a greater number of caesarean sections in the thrombocytopenia group 54.5% vs 25.1% in the control group.

XIII. PPH and types of PPH

There were 23 cases (13.8%) of PPH in the thrombocytopenia group vs 4 cases (2.4%) in the control group. The various factors contributing to PPH were pre-eclampsia, HELLP, DIC and coagulation failure especially, following abruption which resulted in both atonic PPH and PPH due to DIC.
Other complications noted in the case group was seizures 6 cases (3.2%), renal complications and oliguria 5 cases (2.8%), bleeding and coagulation abnormalities 15 cases (9%), sepsis cases 1 (0.57%), pulmonary oedema 1 cases (0.57%) and 1 cases (0.57%) case of maternal death while no such maternal complications were observed in the control group.

The case of maternal mortality was an SLE patient who went into DKA and thrombocytopenia and died due to intractable pulmonary oedema during LSCS done for failed induction. She had received 3 doses of methyl prednisolone prior to her LSCS to improve her platelet count. There were 38 cases of abruption in the case group (Grade I-3 cases, Grade II-15 cases, Grade III-20 cases) compared to only 2 cases (Grade II) in the control group.

Discussions
Our study aims at the maternal outcome of pregnancies complicated by thrombocytopenia. All the causes of thrombocytopenia in pregnant ladies admitted to our labour room were studied including acute onset of thrombocytopenia in patients presenting with acute causes like abruption and DIC.

55.1% of cases had only mild thrombocytopenia and 76% of the cases of thrombocytopenia were diagnosed or had their first presentation in the third trimester. The severe form of thrombocytopenia (count < 50000/ cmm) were mostly constituted by ITP patients (11%) and SLE (9%) patients, pre eclampsia with HELLP (17%) (similar to the study by Vyas. Retal – GT 44.6%, HELLP 22%). Also causes of abruptio placentae with DIC had a severe thrombocytopenia with associated coagulation abnormalities. There were 2 cases of viral infections (dengue fever) with thrombocytopenia managed with supportive measures. Hence the importance of routine examination of platelet counts in all the trimesters to diagnose the condition and reduce maternal and fetal complications.

In our study, gestational thrombocytopenia is the most common cause of thrombocytopenia (40%). Around 10.8% cases were due to pre-eclampsia and 17.4% constituted by HELLP syndrome. Many patients had co-existing multiple conditions like SLE, ITP and pre-eclampsia.

Autoimmune causes like ITP (11.4%), SLE(9%) and SLE with 2 degree ALPA(2.4%) and 1 degree ALPA(2.4%) also constituted a sizeable portion. Being a tertiary case centre with an efficient haematology department in our medical college, we had many such patients referred to us, which resulted in a multidisciplinary and efficient management system for these patients.

Regarding gestational age at delivery, 44% of the cases delivered before 36 weeks of gestation compared to only 8.4% among controls. This was because the pre-eclampsia, HELLP, APLA syndrome and abruptio placentae cases had to be terminated before term for both maternal and fetal sake. These patients had placental insufficiency and FGR which also demands earlier termination of pregnancies.

There was more caesarean section in the above case group, 54.5% vs 25.1% in the control group. There were a greater number of previous CS cases among the thrombocytopenia patients. Postpartum haemorrhage (13.8%) among case group vs (2.4%) among controls was significant. Factors contributing were pre-eclampsia, HELLP, DIC, coagulation failure, especially following...
abruption which resulted in both above PPH and PPH due to DIC.

There were other complications noted in the case group- seizures (6 cases), oliguria and renal dysfunction- 5 cases, bleeding and coagulation abnormality- 15 cases, sepsis-1, pulmonary oedema-1 and one case of maternal death due to intractable pulmonary oedema in an SLE patient. No such maternal complications were observed in the control group.

Blood and blood products transfusion like FFP, cryoprecipitate, platelet rich plasma, platelet concentrate were given in Abruption Grade III (12%), Grade II (9%), patients with DIC, HELLP syndrome (17.4%) and also in ITP patients intractable to medical management.

Conclusion
In general, the gestational thrombocytopenia (GT) and ITP groups had a favorable outcome in our study. Gestational thrombocytopenia is not associated with an increased incidence of pregnancy related complications or with the delivery of a thrombocytopenic offspring. ITP and SLE tend to occur in younger women. The rarer and more serious group of causes of thrombocytopenia including DIC, familial TTP, APLA syndrome etc are associated with placental abruption, low Apgar scores and stillbirths.

Higher rates of preterm deliveries (< 34 weeks 22.15%, 34 -37 weeks 21.55%) were observed among patients with moderate to severe thrombocytopenia. This is due to labour induction, abruption, IUeGR and for maternal sake in severe forms of pre-eclampsia and HELLP syndrome. The management of SLE, ITP patients etc were done in collaboration with the Haematology Department Multidisciplinary approach of management was undertaken whenever needed. Adverse perinatal outcome was mostly associated with abruption, DIC, APLA syndrome and HELLP syndrome.

Careful surveillance is required for all such high-risk pregnancies for early detection and treatment of possible complications, in order to reduce to maternal and neonatal morbidities.

Recommendation
Since majority of cases of thrombocytopenia are diagnosed during pregnancy, it should be made a routine to check the platelet count in the first trimester of pregnancy itself. The etiology of thrombocytopenia is to be determined and treatment to be initiated in collaboration with other departments like Haematology, Transfusion medicine, Nephrology, Medical Gastroenterology etc.

Reference
1. Sainio S, Kekomaki R, Riikonen S, Teramo K. Maternal thrombocytopenia at term- a population based study. Actaobstgynecolscand 2000; 79(9);744-749.
2. Magann IT et al. Twelve steps to optimal management of HELLP syndrome. Clin Obstet Gyncol 1999; 42:532-50.
3. Provan D et al (2010) international consensus report on the investigation and management of primary immune thrombocytopenia. Blood I 15:168-186.
4. Adams J0T et al. Maternal thrombocytopenia in pregnancy. Cliniscs in laboratory medicine, 2013-06-01, Volume 25. Issue 2, Pages 327-341.
5. Sullivan CA et al. Management of the obstetric patient with thrombocytopenia. ClinObstetGynecol 1995;38:521-34.
6. Rodger M et al. Hematological problems in obstetrics. Best practice & Research: Clinical Obstetrics &Gynaecology, 20d15-07-01, Volume 2.9. Issue 5. Pages 671-684.
7. Ying-Hsuan Lin et al. Perinatal outcome in normal pregnant women with incidental thrombocytopenia at delivery. Taiwanese Journal of Obstetrics and Gynaecology, 2013-19’01, Volume 52, Issue 3, Pages 341-350.
8. Meyer o, (2003) Lupus et autresconnectivites et vie hormonale, Gynecologie Obstetrique & Fertilite.. 31, 746-756. Doi: 10.1016/S 1297-9589(03)00203-0.

9. Errarhay, s et al. Antiphospholipid antibody syndrome and pregnancy. Open Journal of Obstetrics and Gynecology Vol. 3 No. 4 (2013), Article ID: 31566, 3 pages DOI :10.4236/ojog.2013.34071.

10. Arvieux, J. and Machulla..E.(2002) Le syndrome des antiphospholipids. Annales de Cardiologie et d Ange-iologie, 51J 46-151. Doi: 10.1016/S0003-3928(02)00087-2.

11. Cleary KL et al. (2009) Pre-eclampsia and the kidney. SeminPerinatol 33:173-178.

12. Barton JR et al (2004) Diagnosis and management of hemolysis, elevated liver enzymes and low platelets syndrome. Clin Pcrinaiol 31:807-833.

13. Vyas. Retal study of mild versus moderate to severe thrombocytopenia in 3\textsuperscript{rd} trimester of pregnancy in a tertiary care hospital, NHL journal of Medical Sciences/Jan 2014/vol 3/issue.

14. Mamta S et al. Thrombocytopenia during pregnancy journal of evalution of medical anda dental sciences 2010 month, October Volume; 3 issue:57 page;12956-12960.