2014

www.jmscr.igmpublication.org

Impact Factor 3.79 ISSN (e)-2347-176x

Journal Of Medical Science And Clinical Research

Effect of Coagulation Profile in Third Trimester of Pregnancy

Author

Dr Tazkira Begum¹, Dr Wasima Jahan², Dr Rwitusmita Bharali³, Dr Syed Md Ibrahim Kabir⁴

¹Assistant Professor, Department of Physiology, Assam Medical College, Dibrugarh
 ²Proffesor, Department of Physiology. Assam Medical College, Dibrugarh
 ³Demonstrator, Department of Physiology, Assam Medical College, Dibrugarh
 ⁴M & HO Obst & Gynaecology. Assam Medical College, Dibrugarh

ABSTRACT

Normal pregnancy associated with many changes in coagulation profile. These changes are important for intrapartum blood loss and also for thromboembolism during pregnancy and labour. The present study aimed at to find out the quantitative change in coagulation profile in early and later part of third trimester. This study was carried out in the Department of physiology, AMC, Dibrugarh, Assam. 50 cases of pregnant women in third trimester are enrolled for the study. Among which 29 cases are in 29th to 34th wks and 21 cases are within 35th to 39th wks of gestation. For control 50 ages matched healthy female of child bearing age group are taken. Platelet count , Bleeding time and coagulation time were done in each control and cases. The mean platelet count in pregnancy at 29th to 34th wks was more than that of non pregnant but the value was less in pregnancy at 35th to 39th wks. Similarly mean bleeding time and coagulation time was also increases in third trimester than the non pregnant. But both the values are decreases in 35th to 39th wks than the 29th to 34th wks. Platelet count was progressively decreased in later part of pregnancy women compared to non pregnant women. It is partly due to hemodilution and partly due to increased platelet activation and accelerated clearance (Shehlata et al., 1999 and McCrae, 2010).To conclude thromboembolism and bleeding disorder are two major complications of pregnancy. So to prevent the complication proper study of coagulation profile is necessary.

Key words: Platelet count, bleeding time, coagulation time, third trimester, thromboembolism.

INTRODUCTION

Normal pregnancy is associated with many hematological, biochemical and clinical changes. Among these. haemostatic changes of hematologic profile contribute to important maternal and fetal outcome. Most changes in coagulation profile create a state of hypercoagulability. Thrombocytopenia is the most common haemostatic abnormality observed in pregnancy.¹ In many healthy women (around10%) late pregnancy is associated with thrombocytopenia. At least in part this is due to haemodilution but the increase in mean platelet volume² suggests that a compensated state of progressive platelet destruction occurs. Additional evidence of *in vivo* platelet activation in late pregnancy is the increased concentration of \Box thromboglobulin³ of and thromboxane A2 derivatives⁴.

The normal human pregnancy lasts for about 280 days (40 weeks), and has a large impact on the well being of a woman without any underling medical disorder at the same time makes the foetus vulnerable to the changes in the mother's internal and external physiological status. Both mother and the foetus are major consideration in the management of pregnancy⁵.Third trimester of pregnancy starts from 29th to 40th wks of gestation. Fetus becomes mature at this stage.

During pregnancy the concentrations of coagulation factors VII, VIII, IX, X, XII and von Willebrand factor rise significantly, accompanied by a relevant increase in the concentration of plasma fibrinogen^{6,7,8,9}.Plasma fibrinogen often increases to over 600 mg/dL in late pregnancy¹⁰.

Factor VII may increase as much as tenfold in pregnancy^{1,11}. The von Willebrand factor and factor VIII are elevated in late pregnancy, when coagulation activity is about twice that in the non-pregnant state¹. The increase in factor IX concentrations during pregnancy is reported by several authors to be small¹¹, as is the decrease in factor XI¹². After an initial increase, factor XIII falls gradually, reaching 50% of the normal non pregnant value at term.

The important relationship of platelet to spontaneous haemostatic was first visualized by Hayem in 1891and it was strengthen by Duke in 1912, who clearly demonstrated a correlation between peripheral blood platelet count and bleeding time. Two important functions of platelet are a) Haemostasis by platelet plug formation and b) blood coagulation.

MATERIAL AND METHOD

The study was conducted in Department of Physiology, Assam Medical college, Dibrugarh, Assam. The 50 nos. of third trimester pregnant cases were randomly selected from the department of obst and gynae, Assam Medical college, Dibrugarh, Assam. For control 50 nos of healthy female of same age group were enrolled. For both cases and control the age group is 18 to 35 years. The cases were taken from both primipara and multipara women. The cases having high blood pressure, oedema, anaemia, albiminurea, and abnormalities in cardiovascular, respiratory and urinary system were excluded. In the study to look for the coagulation profile, quantitative estimation of platelet count, coagulation time and bleeding time were done.

Platelet count was done by Brecher and Cronkite method¹³. Normal platelet count was 1.5 to 3 lacs / cumm of blood. Coagulation time and bleeding time was carried out by Sabraze capillary tube method¹⁴ and Dukes method¹⁵ respectively. Normal counts were 2 to 6 minutes and 2.5 to 9.5 minutes respectively.

The data was analyzed by Microsoft excel and statistical package of social sciences (SPSS version 20.0). Mean and standard deviation were calculated and reported for quantitative variables. The statistical difference were tested by using one way ANOVA (Analysis of Variance). A P- value of < 0.05 was considered as statistically significant.

ETHICAL CONSIDERATION

The necessary approval to conduct the study was obtained from ethical committee of Assam Medical College, Dibrugarh, Assam. Consent form obtained from all participants to ensure their voluntary participation.

RESULTS

The study population consist of 50 control and 50 cases.

The distribution of cases and their results in third trimester together with the controls are summarized in Tables 1-3

Table 1 : Shows Platelet count in l	cs/ cumm in different stages
--	------------------------------

Sr.	Different stages	No. of	Range	Mean	SD	SE
no.		cases				
1	Non pregnant	50	1.5—3.35	2.396	0.408	0.057
2	Pregnant in third trimester	50	2.85	2.331	0.419	0.059
3	Pregnancy in 29 to 34 wks	29	2.62—5.48	2.432	0.421	0.014
4	Pregnancy in 35 to 40 wks	21	2.16-5.33	2.191	0.381	0.018

Histogram: 1 Shows Platelet count in lacs/ cumm in different stages



Sl no.	Different stages	No. of	Range	Mean	SD	SE
		cases				
1	Non pregnant	50	2.08-4.56	3.001	0.508	0.071
2	Pregnant in third trimester	50		3.464	0.760	0.010
3	Pregnancy in 29 to 34 wks	29	2.62-5.48	3.487	0.715	0.024
4	Pregnancy in 35 to 40 wks	21	2.16-5.33	3.439	0.826	0.028

Table 2: Shows Coagulation time in minute different stages





Table 3: Shows Bleeding time in minutes in different stages

Sl no	Different	No. of	Range	Mean	SD	SE
	stages	cases				
1	Non	50	1.5-2.66	1.783	0.203	0.027
	pregnant					
2	Pregnant in	50	2.85	1.883	0.350	0.049
	third					
	trimester					
3	Pregnancy	29	1.5-2.42	1.948	0.291	0.010
	in 29 to 34					
	wks					
4	Pregnancy	21	1.11-2.43	1.844	0.401	0.019
	in 35 to					
	40wks					

JMSCR Volume||2||Issue||10||Page 2559-2566||October-2014



Histogram : 3 Shows Bleeding time in minutes in different stages

Table :4 A, B, C, D Shows the results of statistical comparison of coagulation profile in the third trimesters to non pregnant

Table 4: A.

Parameter	Non	Pregnant	50	Pregnant	50	cases	Significance
	cases			Mean±SD			
	Mean±	SD					
Platelet count	2.396±	0.408		2.331±0.41	19		>0.05
Coagulation time	$1.783 \pm$	0.203		1.883±0.35	50		>0.05
Bleeding time	3.001±	0.508		3.464±0.76	50		>0.001

Table : 4 B

Parameter	Non Pregnant 50 cases	Cases During 29-34	Significance
	Mean±SD	wks of Pregnancy	
		29 cases	
		Mean ±SD	
Platelet count	2.396±0.408	2.432±0.421	>0.05
Coagulation time	1.783±0.203	1.948±0.291	< 0.001
Bleeding time	3.001±0.508	3.487±0.715	< 0.01

Table : 4 C

Parameter	Non Pregnant 50	Cases During 35-40 wks of	Significance
	cases	Pregnancy	
	Mean±SD	29 cases	
		Mean ±SD	
Platelet count	2.396±0.408	2.191±0.381	< 0.05
Coagulation time	1.783±0.203	1.844±0.401	>0.05
Bleeding time	3.001±0.508	3.439±0.826	< 0.01

Dr Tazkira Begumme et al JMSCR Volume 2 Issue 10 October 2014

Parameter Cases During 29-34		Cases during 35-	Significance
	wks of Pregnancy	40wks of Pregnancy	
	29 cases	ases 21 cases	
	Mean ±SD	Mean±SD	
Platelet count	2.432±0.421	2.191±0.381	< 0.001
Coagulation time	1.948±0.291	1.844±0.401	< 0.05
Bleeding time	3.487±0.715	3.439±0.826	>0.5

Table : 4 D

In table 1-3 and Histogram1,2,3 shows the mean \pm SD of platelet count, coagulation time and bleeding time in non pregnant , pregnant third trimester and in early and later part of third trimester. In table 1, shows the mean platelet count in entire third trimester but it increases in early third trimester of pregnancy than the non pregnant. But the value decreases in later part of third trimester or in 35 to 40 weeks of pregnancy. Similarly the mean \pm SD of coagulation time and bleeding time both increases in third trimester than the non pregnant, but the values are again decreases in later part of pregnancy than the early part of third trimester.

In table 4 -A, B, C, D shows the comparison of coagulation profile in control and study group. Here the comparison of mean \pm SD of platelet count between non pregnant and pregnant and non pregnant and early part of third trimester is statistically not significant (Table A and B). But comparison with non pregnant to later part of third trimester and comparison of early and later part of third trimester are statistically very significant (Table C and D).

Comparison of mean±SD coagulation time in non pregnant to early and later part of third trimester and early and later part of third trimester are statistically very significant. In bleeding time mean \pm SD comparison in non pregnant to early and later part of third trimester are statistically highly significant. But comparison between early and later part of third trimester is not significant.

DISCUSSION:

Quantitative changes in platelets occur in various thrombotic and hemorrhagic states are now a day in focused. In recent years considerable amount of works have been done regarding coagulation profile in pregnancy, labor and puerperium. But there are some lack of uniformity of findings by like socio-economic various factors and nutritional status of the woman and racial and other environmental condition. Now, the objective of the present study are to find out the quantitative change in platelet count , bleeding time and coagulation time in early and later part of pregnancy and also to compare the values with non pregnant and each other.

In the present study showed that the mean platelet count increases in early part of third trimester than the non pregnant but again decreases in later part of pregnancy. The increase in platelet count in early part of third trimester of pregnancy than the non pregnant is due to the hypervolumic status of pregnancy. But the mean platelet count decreases in later part of third trimester of pregnancy than the early part may be due to fall in plasma volume due to haemodilution and increase utilization or changes in stimulation of production during later part of third trimester of pregnancy^{16,17,18}. Present study correlates the findings of Celento et al, 1931: Kennan and Bell, 1957; Czernik et al 1963. The present study showed the mean value of coagulation time is increases in both early and later part of pregnancy than the non pregnant state. The present study findings regarding coagulation time is similar to that of M. Hellgren and M. Blomback. This may be due to increase no of clotting factors in the circulation, but at the same time there will be decrease no of anticoagulant in the circulation and reduces in fibrinolytic activity. So there will be increase coagulation time in pregnancy. But in later pregnancy there will be depletion of plasma volume leading to increase concentration of clotting factors in the circulation this imparts decreases in coagulation time in later part of third trimester of pregnancy than the early part of the same.

In the present study the bleeding time found to be prolonged in third trimester of pregnancy than the non pregnant state. Statistically the values are highly significant. But it does not correlate with the platelet count in the study. In the study we found that coagulation time correlated with the bleeding time. Because from the table 4 we found that when coagulation time increases in early third trimester ,the bleeding time decreases and in later part of third trimester coagulation time decreases than that of early third trimester ,so the bleeding time also decreases than the earlier. This may be due to decrease amount of anti coagulant in the circulation play a role in decreasing bleeding time.

CONCLUSION:

In summary, pregnancy is associated with major changes in haemostasis .It includes increases in the platelet count ,decreases in the quality of natural anticoagulants and a reduction in fibrinolytic activity leading to difference in bleeding and coagulation time. These changes are greatest at the time of delivery .Platelet counts may be lower in pregnancy most commonly due to gestational thrombocytopenia or ITP and it is more marked in later part of third trimester of pregnancy .

REFFERANCE

- Stirling Y, Woolf L, North WR, Seghatchian MJ, Meade TW. Haemostasis in normal pregnancy. Thromb Haemost 1984;52:176-82.
- 2. Wallenburg HC, van Kessel PH. Platelet lifespan in normal pregnancy as determined non-radio-isotopic by а technique. Br J Obstet Gynaecol 1978;85:33-6.
- Douglas JT, Shah M, Lowe GD, Belch JJ, Forbes CD, Prentice CR. Plasma fibrinopeptide A and □-thromboglobulin in pre-eclampsia and pregnancy hypertension. Thromb Haemost 1982;47:54-5.
- Fitzgerald DJ, Mayo G, Catella F, Entman SS, FitzGerald GA. Increased

JMSCR Volume||2||Issue||10||Page 2559-2566||October-2014

2014

thromboxane biosynthesis in normal pregnancy is mainly derived from platelets. Am J Obstet Gynecol 1987;157:325-30

- Loh FH, ArulkumaranS, Montan S,Ratnam SS. Maternal Mortility: Evolving Trends. Asia Oceania J.obstet & Gynaecol1994;20 (3): 301-4 Stirling Y, Woolf L, North WR, Seghatchian
- MJ, Meade TW. Haemostasis in normal pregnancy. Thromb Haemost 1984;52:176-82
- Bonnar J, Haemostasis and coagulation disorders in pregnancy. In: Bloom AL, Thomas DP, editors. Haemostasis and Thrombosis, Churchill Livingstone, Edinburgh (1987), p. 570-84.
- Letsky EA coagulation problems with pregnancy. Charchill and Livingstone. Edinbarg (1985)
- Greer IA, Haemostasis and thrombosis in pregnancy. In: pregnancy Bloom Al ,Forbes CD, Thomus DP, Tuddeham EGD, Editors Haemostasis and Thrombosis(3rd edition) Charchill and Livingstone, Edinbarg (1994) p. 987-1015
- Francalanci I, Comeglio P, Liotta AA, Celtai AP, Fedi S ,Parretti E et al. D-dimer concentrates during normal pregnancy, as measured by ELISA. Thomb Res 1995; 78: 399-405
- Beller FK, Ebert C. The coagulation and Fibrinolytic enzyme system in normal pregnancy and puerperium. Eur.J Obst Gynaecol Repord Biol 1982 ; 13: 177-97

- Hellgren M, Bloomback M. Studies on blood coagulation and fibrinolysis in pregnancy, during delivery and in puerperium. Gynaecol obst Invest 1981; 2: 37-46
- Brecher ,G; Cronkite, E.P ; Morphology and enumeration of human blood Platelets 1950; J. Apple. Physiol; 3: 365.
- 14. Sabarraze; lited by Kolmer, J.A; J
 Paulding E.H. Robinson, H.W; 1969
 Approved Laboratory Technique, 5th
 edition.
- 15. Duke, W.W; 1912 The pathogenesis of purpura haemorrhagica with special reference to the part played by blood Platelets; Amch, inter. Med.,10:445
- 16. Imoru M, Emeribe AO, Haemorrhagical profiles in apparently healthy pregnant woman in Calabar, Nigeria. African J Bio 2008; 7(24): 4354-8.
- Stuart C, Christoph L. Physiological changes in pregnancy. In obstetrics by Ten teachers. (Indian Edition) Ajanta offset and packagings Limited ; 2000.
- 18. Salawu L, Durosinmi MA. Erythrocye rate and plasma viscosity in health and disease . Niger J Med 2001; 10 (1) : 11-3.

Dr Tazkira Begumme et al JMSCR Volume 2 Issue 10 October 2014