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Original research article

Prevalence of thromboembolism in pregnancy and puerperium in a tertiary care center and risk factor assessment: A cross-sectional study

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Abstract

Background: Pregnancy and puerperium increases the risk of venous thromboembolism 4-to 5-fold over that in the non-pregnant state. Compared to non-pregnant, pregnant women have a two and a half times increased risk of developing pulmonary embolism. Venous thromboembolism is often fatal. Depending on reports estimate that 10%-30% of patients had mortality within 30 days. The purpose of this study is to estimate the prevalence of thromboembolism in pregnancy and puerperium, risk factors, need for anticoagulants, maternal and fetal outcome in a Govt. Tertiary care center in Chennai. **Materials and Methods:** A prospective Descriptive study was conducted in Dept of Obstetrics & Gynaecology, Govt. Kilpauk Medical college, Chennai during February 2019-september 2019 among 200 antenatal and postnatal mothers. Descriptive statistics, Chi Square test were used. A p value of <0.05 was considered statistically significant.

Results: In BMI up to 23 having thromboembolism of 0%, in BMI from 24-29, incidence was thromboembolism of 9.5% and in BMI of 33-36 having thromboembolism of 0%. In patients having BMI between 24-29, risk of thromboembolism was observed high. Hence BMI was considered significant risk factor. One patient developed pulmonary embolism in postnatal period, emergency CTPA done, given LMWH for 3 months & patient revived.

Conclusions: BMI more than 26, multiparity, previous history of surgery, immovability, varicose veins, gestational hypertension, anemia, APLA positive and Eclampsia were the significant risk factors. Since pregnancy is a hyper-coagulable state, by identifying these risk factors and initiating LMWH, we can prevent the complications of DVT. Hence maternal morbidity and mortality can be reduced.

Keywords: Thromboembolism, pregnancy, puerperium, risk factors

Introduction

Pregnancy and puerperium increases the risk of venous thromboembolism 4-to 5-fold over that in the non-pregnant state [1]. The manifestations of venous thromboembolism are deep venous thrombosis and pulmonary embolism. Compared to non-pregnant, pregnant women have a two and a half times increased risk of developing thrombotic event and 20 fold risk in puerperium. The incidence of pulmonary embolism is 7-8 times higher in puerperium [1]. Sequelae of DVT and PE include complications such as Pulmonary Hypertension, Post-thrombotic Syndrome and venous insufficiency. Incidence is estimated to be between 0.08 and 1.2 percent following vaginal delivery and 3.0 percent following caesarean section [2]. There is no surveillance for thromboembolism, so the number of people affected by it is not known. Based on analysis of clinical studies, the overall annual incidence is estimated to be between 1 and 2 per 1000 of the population [3]. These incidence rates differ by age, race, and gender. The incidence ranges from 1 per 100,000 in the young and increases to about 1 per 100 in people aged ≥80 years. The overall rate is higher among blacks and whites than among other races. Men have a slightly higher overall incidence rate than women, but women have a slight increase during the reproductive years [4]. Because of the difficulty in documenting DVT and PE, VTE may be under-reported. Venous thromboembolism is often fatal. Depending on reports estimate that 10%-30% of patients had mortality within 30 days; the majority of deaths occur among those with PE, as an estimated 20%-25% of all PE cases present as sudden death [3]. Other serious complications of DVT and PE include increased risks of recurrent

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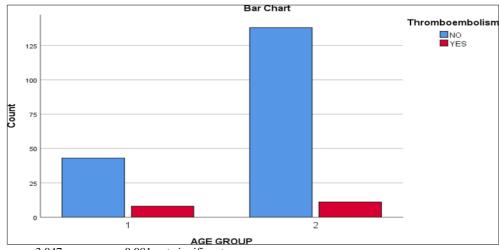
thromboembolism and chronic morbidity. Even after a standard course of anticoagulant therapy, $1/3^{rd}$ of previous thromboembolism patients have recurrence within 10 years of the primary event, with risk remaining in the rest of their life ^[5]. Half of DVT patients develop post-thrombotic syndrome and chronic venous insufficiency, conditions characterized by pain, swelling, skin necrosis, and ulceration. If on lifelong anticoagulation, they are at increased risk of bleeding tendencies. The purpose of this study is to estimate the prevalence of thromboembolism in pregnancy and puerperium, risk factors, need for anticoagulants, maternal and fetal outcome in a Govt. Tertiary care center in Chennai.

Methods

A prospective Descriptive study was conducted in Dept of Obstetrics & Gynaecology, Govt. Kilpauk Medical College, Chennai during February 2019-september 2019 among 200 antenatal and postnatal mothers. Patient who are having signs and symptoms of Venous Thromboembolism, antenatal risk factors like age > 35, Obesity, Multipara(>3), BMI more than 30, Intra Uterine Death, Abruption, Anemia, Pre-eclampsia, Gross Varicose veins, prolonged immobility, medical comorbidities, thrombophilias. Postnatal risk factors like ceasarean section, operative vaginal deliveries, prolonged surgeries were included. Patients with previous history of venous Thromboembolism Valvular heart diseases were excluded. The patients who are diagnosed as having signs and symptoms of Venous Thromboembolism, having risk factors are taken as study group. The following investigations were done in all like coagulation profile, Chest X ray in postnatal mothers, Color Doppler study of lower limb veins, CT and Pulmonary Angiography.

Results

The following figure describes the age group of the study participants.



Chi square = 3.047

p= 0.081 not significant.

Fig 1

In both the groups less than 10% observed in both age group, 8(4%) in 18-21 age group and 11(5.5%) in 22-25 age group. No statistically significant differences in age distribution of patients assigned to the 2 groups. The following table describes the relationship between BMI and the thromboembolism.

Table 1

| BMI Group * Thromboembolism | | | | | | |
|-----------------------------|----|--------------------------|--------|--------|--------|--|
| No | | | | | Total | |
| BMI Group | 1. | Count | 40 | 0 | 40 | |
| | | % within Thromboembolism | 22.1% | 0.0% | 20.0% | |
| | | % of Total | 20.0% | 0.0% | 20.0% | |
| | 2. | Count | 117 | 19 | 136 | |
| | | % within Thromboembolism | 64.6% | 100.0% | 68.0% | |
| | | % of Total | 58.5% | 9.5% | 68.0% | |
| | 3. | Count | 24 | 0 | 24 | |
| | | % within Thromboembolism | 13.3% | 0.0% | 12.0% | |
| | | % of Total | 12.0% | 0.0% | 12.0% | |
| | | Count | 181 | 19 | 200 | |
| Total | | % within Thromboembolism | 100.0% | 100.0% | 100.0% | |
| | | % of Total | 90.5% | 9.5% | 100.0% | |

p < 0.004

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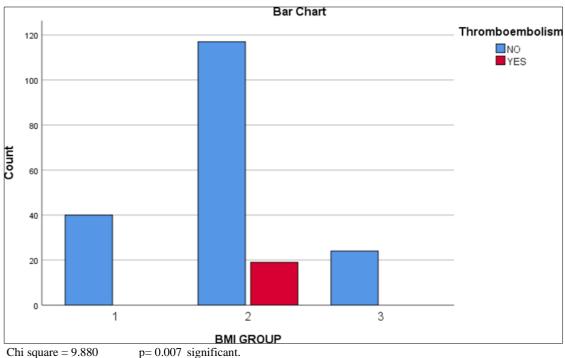


Fig 2

In 3 groups less than 10% observed in 3 BMI groups, 0(0%) in BMI group, 19(9.5%) in BMI group2 and 0(0%) in BMI group3. Statistically significant differences in in distribution of patients assigned to 2nd group with other groups.

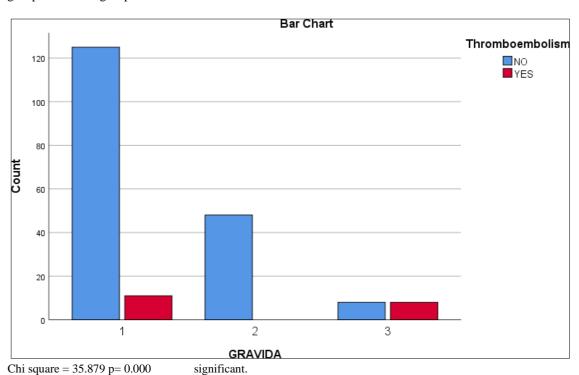
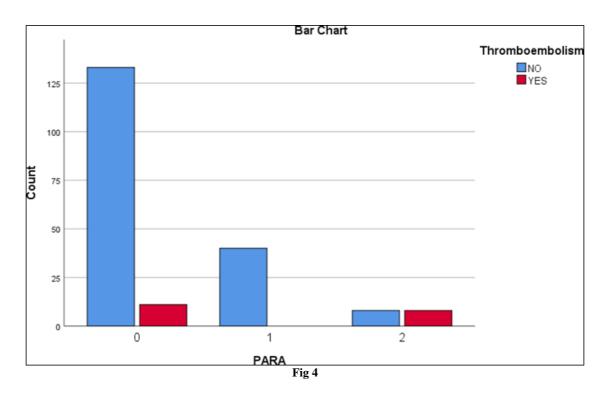


Fig 3

In 3 groups less than 10% observed in 1 and 3 GRAVIDA groups, 11(5.5%) in GRAVIDA group, 0(0%) in GRAVIDA group2 and 8(4%) in GRAVIDA group3. Statistically significant differences in distribution of patients assigned to the 1 and 3 groups

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In 3 groups less than 10% observed in 3 PARA groups, 11(5.5%) in GRAVIDA group, 0(0%) in PARA group 2 and 8(4%) in PARA group3. Statistically significant differences in distribution of patients assigned to the 1 and 3 Para groups

Table 2

| Thromboembolism | | | | | | |
|-----------------|---|--------------------------|--------|--------|--------|--|
| No | | | | | Total | |
| Immovability | 0 | Count | 178 | 11 | 189 | |
| | | % within Thromboembolism | 98.3% | 57.9% | 94.5% | |
| | | % of Total | 89.0% | 5.5% | 94.5% | |
| | 1 | Count | 3 | 8 | 11 | |
| | | % within Thromboembolism | 1.7% | 42.1% | 5.5% | |
| | | % of Total | 1.5% | 4.0% | 5.5% | |
| Total | | Count | 181 | 19 | 200 | |
| | | % within Thromboembolism | 100.0% | 100.0% | 100.0% | |
| | | % of Total | 90.5% | 9.5% | 100.0% | |

Chi square = 54.125 p = 0.000 significant.

In 2 groups less than 10% observed in 2 Immovability groups, 11(5.5%) in group and 8(4%) in group2. Statistically significant differences in distribution of patients assigned to the two groups.

Other factors like varicose veins, presence of Anemia, GHT were significantly associated with thromoboembolism. Factors like RD, NICU, FET INFEC, Death were not statistically associated with thromboembolism. Thrombophilia was associated with antenatal or puerperal thromboembolism like as APLA, Eclampsia, Heparin usage all had statistically significant association with Thromboembolism.

Discussion

In age group of 18-22 prevalence of Thromboembolism is 4% and in age group of 22-25 prevalence of Thromboembolism is 5.5%. It is not statistically significant. Hence age group is not considered as risk factor in Ante Natal and Puerperium. In BMI up to 23 having Thromboembolism of 0% in BMI from 24-29 having Thromboembolism of 9.5% and in BMI 33-36 having Thromboembolism of 0%. In patients having BMI between 24-29 risk of Thromboembolism present. Hence BMI considered significant Risk factor. The risk of Thromboembolism is high in multi Gravida (multiple pregnancy), hence Gravida considered significant Risk factor ^[8, 9].

Abortion is not considered significant risk factor ^[10], since prevalence of Thromboembolism in abortion is 9.5%. Previous History of surgery is considered as a significant risk factor. History of varicose veins is considered as a significant risk factor. History of immovability for prolonged period is considered as a significant risk factor since Prevalence of Thromboembolism is 5.5%. History of Gestational Hypertension for is considered as a significant risk factor since Prevalence of Thromboembolism is 4%. There is no Significant Correlation between mild and severe GHTN ^[11]. The NICU admission for babies in mothers having Thromboembolism not considered as significant risk factor. In babies having

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respiratory distress no statistical significant persist. Hence it is not considered as risk factor. In babies having fetal infection no statistical significant persist. Hence it is not considered as risk factor. There is no significant correlation between death and Thromboembolism.

The prevalence of Thromboembolism in Antenatal patients is 60% and in Post Natal patient is 40%. Hence there is no statistical significance between both groups. Prevalence is nearly the same in both groups. Hence it is not statistically significant. There is statistical significance between Eclampsia and Thromboembolism. Hence it is considered as Risk factor. Four cases developed cortical venous thrombosis (have not received prophylactic LMWH. Low Molecular weight Heparin is given in patients having risk factors and in patients having Thromboembolism. Duration and dose of prophylactic LMWH has to Increased. One patient developed pulmonary embolism in postnatal Period, emergency CTPA done, given LMWH for 3 months & patient revived. There is statistical significance between APLA and thromboembolism. There is statistical significance between Anemia and Thromboembolism. No correlation between mild and severe anemia.

Conclusions

Thus patients with BMI more than 26, multiparity, Previous History of surgery, Immovability, Varicose veins, Gestational Hypertension, Anemia, APLA positive and Eclampsia as significant risk factors ^[12]. Since pregnancy is a hypercoagulable state, by identifying these risk factors and initiating LMWH, we can prevent the complications of DVT and hence maternal mortality and morbidity can be reduced ^[13].

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Competing Interest: Nil.

References

- 1. Pomp ER, Lenselink AM, Rosendaal FR, Doggen CJ. Pregnancy, the postpartum period and prothrombotic defects: risk of venous thrombosis in the MEGA study. J Thromb Haemost. 2008 Apr;6(4):632-7.
- 2. Heit JA, Kobbervig CE, James AH, Petterson TM, Bailey KR, Melton LJ 3rd. Trends in the incidence of venous thromboembolism during pregnancy or postpartum: a 30-year population-based study. Ann Intern Med. 2005 Nov;143(10):697-706.
- 3. Goldhaber SZ, Visani L, De Rosa M. Acute pulmonary embolism: clinical outcomes in the International Cooperative Pulmonary Embolism Registry (ICOPER). Lancet. 1999 Apr;353(9162):1386-9.
- 4. Polak JF, Wilkinson DL. Ultra-sonographic diagnosis of symptomatic deep venous thrombosis in pregnancy. Am J Obstet Gynecol. 1991 Sep;165(3):625-9.
- 5. Pabinger I, Grafenhofer H, Kyrle PA, *et al.* Temporary increase in the risk for recurrence during pregnancy in women with a history of venous thromboembolism. Blood. 2002 Aug; 100(3):1060-2.
- 6. Macklon NS, Greer IA, Bowman AW. An ultrasound study of gestational and postural changes in the deep venous system of the leg in pregnancy. Br J Obstet Gynaecol. 1997 Feb;104(2):191-7.
- 7. American College of Obstetricians and Gynecologists Practice Bulletin No 123: Thromboembolism in pregnancy; c2011 Sep.
- 8. Nijkeuter M, Ginsberg JS, Huisman MV. Diagnosis of deep vein thrombosis and pulmonary embolism in pregnancy: a systematic review. J Thromb Haemost. 2006 Mar;4(3):496-500.
- 9. Perrier A, Roy PM, Sanchez O, Le Gal G, Meyer G, Gourdier AL. Multidetector-row computed tomography in suspected pulmonary embolism. N Engl. J Med. 2005 Apr 28;352(17):1760-8.
- 10. Bonnar J. Venous thrombosis and pulmonary embolism. In Turnbull's Obstetrics, ed. G Chamberlain, PJ Steer, 671-80. Churchill Livingstone: London; c2001.
- 11. Greer IA. Epidemiology, risk factors and prophylaxis of venous thromboembolism in Obstetrics and Gynaecology. Balliere's Clin Obstet Gynaecol. 11:403-30.
- 12. Macklon NS Greer. The deep venous system in the puerperium: An ultrasound study. Br J Obstet Gynaecol. 1997;104:198-200.
- 13. Royal College of Obstetricians and Gynaecologists. Thromboembolic disease in pregnancy and the puerperium: Acute management. Guidelines No 28. RCOG Press: London; c2007.