Introduction

During pregnancy, there are physiological changes that favor the formation of thrombus, this being the prompting non obstetric cause of maternal mortality and morbidity.

These changes are:

A. Increase of coagulation factors: FI, FVII, FVIII, FIX, FXII
B. Increase of platelet aggregation
C. Changes in the fibrinolytic system, caused basically.

These alterations are transitory, and disappear eight weeks after birth. But a history of thrombosis and the presence of thrombophilia, increases the risk of venous or lung thromboembolism [1]. The recurrent spontaneous abortions, are two or more consecutive miscarriages before the 12th week, and are more frequent in women with thrombophilia, this risk is also associated with thromboembolism and implantation failure [2]. Here are three polymorphisms of the gen of the inhibitor of the activator of plasminogen (PAI-1): 4G/4G, 4G/5G and 5G/5G. The polymorphism 4G/5G has a higher incidence than the homozygous 4G/4G, 5G/5G. PAI-1 is the principal inhibitor physiological of the fibrinolytic process. This process prevents the deposition of excess of fibrin in the placenta, fetal vasculature and stabilizes the base of the placenta [3]. The decrease of the fibrinolytic process can be the cause of infertility and recurrent abortions.

Metodology

A bibliographic review was made, using platforms pubmed and medline. Languages: English, Spanish and Portugues.

Results and Discussion

Due to inconsistencies in studies of PAI-1 and its association with recurrent abortion, Yuejiao Li et al, made a meta-analysis based on two inclusion criteria: 1-case-control studies, 2-studies with enough data to calculate the frequency of allele-genotype. Twenty two studies were included. A positive association was found in Caucasian population, but no association was found for Asian and Latino population. It shows us that different ethnic groups have different frequencies, and they lead to different results, besides that only one study in the Latino population was included.

Another meta-analysis, was made by [9], they used new and more rigorous inclusion criteria: 1-evaluate the association between PAI-1 4G/5G with the risk of recurrent abortion, 2-studies case-control, our projected with cohort 3-Studies with enough information to infer results. This analysis showed that there is an association of PAI-1 4G/5G with recurrent abortion in Caucasians, Asians and Africans, but not in Latinos. Also with three spontaneous abortions, there was no difference between them, the 4G/5G polymorphism, conferring an increased risk to both groups. In summary, this meta-analysis found a positive association between the 4G/5G PAI-1 polymorphism and the increased risk of recurrent miscarriage in Caucasian, African and Asian populations. In addition, effective therapeutic prevention can theoretically be developed by focusing on anticoagulation and improving genetic polymorphism studies in recurrent miscarriages.

Only one study of all the analysis showed no association of PAI-1 and recurrent abortion. It was done by Arabi et al. [8].
wanted to demonstrate this association of PAI-1 and angiotensin I converting enzyme (ACE). Due to heterogeneity in the groups included, it was not possible to demonstrate the relationship of PAI-1 with recurrent abortions, but there was a relationship with ACE. Khosravi [4] study focused in detail on the comparison of 4G/5G polymorphism with recurrent failure and implantation failure.

The study with women with recurrent miscarriage plus two implantation failure. One hundred women with at least two normal live births were included in the control group. The results showed an increase in the PAI-1 4G allele in the three subgroups, significantly higher than in the control group. In several studies therapeutic perspectives are suggested for women with PAI polymorphism, which sometimes is accompanied by other thrombophilias and/or other pathologies, for example the application of metformin in case of polycystic, heparin and aspirin [4,5].

According to a study by [5] on congenital thrombophilia, its reproductive impact and treatment prospects, PAI-1 polymorphism has an uncertain risk, together with protein S deficiency, factor XIII, factor VIII increase, protein C receptor polymorphism. Thrombophilias within this group, 100 grams of aspirin daily and follow up with Doppler of uterine arteries and fetal growth are recommended. Evidence indicates that an increase in PAI-1 levels increases the risk of recurrent abortion and complications during pregnancy, such as pre-eclampsia among others [5].

Conclusion

In conclusion, there may be association with recurrent abortions and PAI-1 (4G/5G). However, large and prospective studies are needed to validate this result. According to the literature evaluated, in the present study, it was demonstrated that the homozygous heterozygote 4G/4G PAI-1 and 4G/5G should be considered as a risk factor and its investigation should be considered in cases of implantation failure and/or recurrent abortions.

References