A Comparative Study: Gene Expression VEGF, FLT-1 and KDR And level of PLGF In Preeclampsia And Normal Pregnancy.

Ibrahim M. EL-Deen¹, Amal K. Seleem², A. A. Abdel Aziz³, E. H. EL-Sayd⁴, Ateya G. Amin⁵,*

¹Department of Chemistry, Faculty of Science, Port Said University, Port Said, Egypt.
²Department of Clinical Biochemistry, Faculty of Medicine, Mansoura University, Mansoura, Egypt.
³Department of Gynecology, Faculty of Medicine, Mansoura University, Mansoura, Egypt.
⁴Department of Chemistry, Faculty of Science, Port Said University, Port Said, Egypt.
⁵Insurance of Health, Central Laboratories, Mansoura, Egypt.

*Corresponding author: attiagomaa72@yahoo.com

ABSTRACT

Involved The vascular endothelial growth factor (VEGF) family of angiogenic growth factors in the pathophysiology of these inconveniences. aims This study to examine Maternal serum levels of placental growth factor (PLGF) and Placental mRNA expression of VEGF, fms-like tyrosine kinase 1(FLT-1) and kinase insert domain receptor (KDR) in normotensive control (NC) women (n = 24) and women with preeclampsia (PE) (n = 50) [25 severe PE (SPE) and 25 Mild PE (MPE)]. We conducted a cross-sectional study in 50 preeclamptic women: 25 women with mild PE, 25 patients with severe PE. While 24 normotensive pregnant women matched for age as a control group. we measured by real time PCR (RT-PCR) Placental mRNA expression of VEGF FLT-1 and KDR, and we measured Maternal serum levels of PLGF by ELISA. there was statistically significant difference between the three study groups as regards all 4 studied laboratory variables. KDR, FLT-1 and PLGF were all statistically significantly higher in control group as compared to either of the two PE groups. The mean VFGF expression was statistically higher in cases of severe PE than Mild PE, both were higher than control group (p<0.0005).

Keywords

Preeclampsia; vascular endothelial growth factor; fms-like tyrosine kinase 1; kinase insert domain receptor

1. INTRODUCTION

Pre-eclampsia affects about 7% of all pregnancies, worldwide. It is a major cause of maternal morbidity and mortality and neonatal (1). Normally, during pregnancy, spiral arteries of uterus undergo a renovating by the intrusion of cytotrophoblast in their walls and in this way diminish an opposition in blood stream to intervillous spaces (2). Lacking invasion of trophoblast brings about placental ischemia and advancement of
hypothesis(3). FLT1 is expressed VEGFA and PIGF and KDR and in villous and extravillous trophoblasts, the vascular endothelium excess as well as increased natural killer cells known to change the level of expression in PE (4). It is suggested that disturbed placental angiogenesis is the main characteristic of causing preeclampsia. Angiogenesis formation process begins with various angiogenic factors such as vascular endothelial growth factor (VEGF) and placental growth factor (PIGF) along with receptors associated with the membrane (5). Mechanisms that promote formation angiogenesis and vascular regeneration in uteroplacental unit is not fully understood. As seen many of the molecular pathways as the placenta linked to defects such as PE; it is known that the angiogenic mediated VEGF family plays a key role path (6). Current data indicates that placental ischemia& trophoblastic hypoprefusion with endothelial dysfunction as the most consistent change. So far, There are many theories pathophysiological of gene expression in PE except for the decrption complete with many Muslim available to explain the different results obtained. In this way, we consider that it is useful to carry out our study using a control typical in situations in a different society to add a guide to the existing literature; Placental VEGF,KDR andFlt-1 mRNA levels have been reported to be decreased, increased or unchanged in PE cases versus uncomplicated controls. So, we undertook this study to determine the level of PLGF in maternal serum and expressions of VEGF,KDRandFlt-1 mRNA in placentae of PE in order to clarify the various discrepancies and gain further evidence for its role in Preeclampsia.

2. PATIENT AND METHODS

2.1 Patients
Fifty patients determined to have Preeclampsia were gathered from the Obstetrics and Gynecology division, Mansoura University, Egypt (Their mean age (years) ± SD were 28.15 ± 5.25) during the period from July 2015 and May 2017. Preeclamptic women were partitioned into 25 women analyzed as mild preeclampsia (MPE), diagnosed as blood pressure ≥140/90 mmHg and 25 women were analyzed as severe preeclampsia (SPE), diagnosed as blood pressure ≥160/110 mmHg. The control group included 24 normotensive pregnant women. The blood tests were taken from all participants during pre-birth visits at second trimester between 20 to 24 gestational weeks. Maternal ages, weights, and height, gravidity, systolic and diastolic blood pressure, were recorded. Exclusion criteria included twin pregnancy, Women with a background marked by diabetes; thyroid, liver, or chronic renal disease; or preexisting chronic hypertension (defined as blood pressure >140/90 mmHg or need for antihypertensive medications before pregnancy or before 20 wk gestation) also if they had a pregnancy termination, a major anomaly all these cases were excluded or if the pregnancy outcome was unknown (i.e. if they did not deliver at our hospital). Inclusion Criteria: Normotensive pregnant women had blood pressure less than 120/80 mm Hg and there was no history of high blood pressure or proteinuria. All pregnant women showed gestational age from 20 to 40 weeks. Control group were taken from women who entered Mansoura University Hospital, Egypt, Obstetric department, Maternal Study cohort within 2wk of each case. Ethical approval: A composed educated assent was taken before test withdrawal. The examination was acted as per the moral guidelines set down in Mansoura Faculty of Medicine.

2.2 Methods
2.2.1 PlGF ELISA
It was withdrawn 3 ml of venous blood samples and left to clot between 15 °C and 22 °C for 30 to 90 minutes. Then were centrifuged for them for 15 minutes at 300 g. Serum was isolated, collected and stored at -20 °C until the time of measure for PlGF (human placenta growth factor, PIGF ELISA Kit / SAB Catalog No: EK1302) Sun Red (Biotechnology Company). Shanghai.
2.2.2 Urine Collection:
For pregnant ladies. Was required to ten ml of urine to detect the protein by dipstick measurement in two samples random at least were randomized to get ahead of them at least 4 hours. Protein means the protein ≥3 g. In 24 hours (medi-test combi & Co. KG, Germany).

2.2.3 RNA extraction from placental tissue
Were performed Fresh samples of 74 placentas humanity within 30 minutes after birth of the placenta, and after washing the samples with normal saline, removed layers amniotic and decidua motherly, then freeze samples in liquid nitrogen and stored at -70 ° C until the displacement of the RNA using miRNeasy mini kit (Qiagen, cat No. 217 004, Germany). RNA was quantified by spectrophotometry(7).

we measured the expression of VEGF, FLT-1 and KDR genes in placenta by used TaqMan real-time PCR (provided by Thermo Scientific, USA, cat No. # K1641).

2.2.3.1 Reverse transcription of RNA extracted for the production of DNA
For each sample, was used 1 μg of total RNA Adjust by reverse transcription using the Maxima® First Strand cDNA synthess kit provided by Thermo Scientific, USA, cat. No.K1641. Following the manufacturer’s instructions, Reverse transcription and the quantitative PCR were performed as previously described (8,9) . Genetic prefixes own purchased from Invitrogen by Thermo Fisher Scientific. It was selected primer sets to amplify the gene PCR after three of the gene sequence database NCBI (10). It included the following assay targeting specific mRNAs in the study.:

VEGF:
Forward: 5'-GGGGGCAGAATCATCAGGAA-3', Reverse: 5'CCAGGGTCTCGATTTGATG-3'.
Ftl1:
Forward: 5'-CTGCAAGATTCAGGCACCTATG-3', Reverse: 5'-GTTTCGCAGGAGGTATGGTGCT-3'.
KDR:
Forward: 5'-GTAGGACTGCTCAGTTCAAACAT-3', Reverse 5'-ACAGTTACTACACCCGTAAG-3'.

2.2.3.2 Quantitative PCR Analysis (11)
PCR was performed in real-time serial to estimate the quantification of genes VEGF and Ftl1 and KDR using SensiFAST SYBR® Lo-ROX (purchased from Bioline, London, United Kingdom, Catalog Number: BIO-94005) following the manufacturer’s instructions. Analysis was performed fusion curve after amplification to confirm the product's privacy and exclude the presence Altmhedy- dimmers. Genetic expression was performed using the relative delta delta cycle threshold (DDCT)by using method 2−ΔΔct . The relative gene expression analysis was done by Delta Delta cycle threshold (DDCT) method, and the average DCT of the healthy volunteers for each target gene was used as the calibrator sample (14,15). The amount of target, normalized to an endogenous reference and relative to a calibrator, was calculated. The fold change is obtained by 2−DDCT. This method assigns a value of 0.7 to the calibrator sample, and all other quantities are expressed as an n-fold difference relative to the calibrator.

3. STATISTICAL ANALYSES
Data analysis was performed by Statistical package for social science software (SPSS) version 25. Data are presented as mean ± SD and compared by One-Way ANOVA and they are presented as median (IQR) and compared by Kruskal-Wallis (K-W) test. Significance was considered at p value less than 0.05.*Data are presented as frequency (percentage) and compared by Chi-Square test (Monte Carlo significance).

**Data are presented as median (IQR) and compared by Kruskal-Wallis (K-W) test. Pairwise comparisons are presented as capital letters (similar letters = no significant difference, while different letters = significant difference).
4. DISCUSSION AND RESULTS

This study involved 74 subjects divided into 3 groups: The first is control group: 24 pregnant ladies with no current or previous pre-eclampsia. Their mean age (years) ± SD = 27.8 ± 4.7 years. The second is mild pre-eclampsia group: 25 pregnant ladies fulfilling the criteria of mild pre-eclampsia. Their mean age±SD were 27.5 ± 4.3 years. The third is severe pre-eclampsia group: 25 pregnant women fulfilling the criteria of severe pre-eclampsia. Their mean age (years) ± SD = 28.8 ± 6.2 years. (figure 1 and Table 1); It showed that a statistically significant difference existed for the other variables; systolic blood pressure (SBP), Diastolic blood pressure (DBP) and Mean average pressure (MAP), they all were statistically significantly higher in severe PE > Mild PE > Control. There are statistically significant differences in the level of protein in the urine with trace and + in mild cases, but ++ and +++ in severe cases (P<0.0005) (figure 2).

This study contains many interesting results and new that will help in the understanding of the expression pattern of angiogenic and anti angiogenic factors in the placenta. figure (3) that there are statistically significant differences between the study groups with respect to the three variable studied laboratory. VFGF was statistically significantly higher in severe PE > Mild PE > Control.

VEGF expression of receptors in the discussion of the placenta, there is a contradiction in the large literature on levels of VEGF placenta in cases of pregnancy PE (14, 15). In 2011, Paiva and colleagues worked on both maternal blood and placental tissue in preeclampsia and found correlation of gene expression between the two (16). mRNA expressions were, highly expressed in preeclampsia, however it have been studied in maternal blood samples by a few others (17,18,19).

Table (1): Comparison of the clinical variables between the three study groups

<table>
<thead>
<tr>
<th>Variable</th>
<th>Group</th>
<th>Statistic</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Control (n=24)</td>
<td>Mild PE (n=25)</td>
<td>Severe PE (n=25)</td>
</tr>
<tr>
<td>Age (years)</td>
<td>27.8 ± 4.7</td>
<td>27.5 ± 4.3</td>
<td>28.8 ± 6.2</td>
</tr>
<tr>
<td>SBP (mmHg)</td>
<td>115 (110-120) A</td>
<td>140 (135-140) B</td>
<td>170 (165-172.5) C</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>DBP (mmHg)</td>
<td>75 (75-80) A</td>
<td>85 (80-90) B</td>
<td>110 (107.5-115) C</td>
</tr>
<tr>
<td>MAP (mmHg)</td>
<td>90 (86.6-91.7) A</td>
<td>103 (99.1-105.8) B</td>
<td>130 (126.6-134) C</td>
</tr>
</tbody>
</table>

*Data are presented as mean ± SD and compared by One-Way ANOVA. **Data are presented as frequency (percentage) and compared by Chi-Square test (Monte Carlo significance). ***Data are presented as median (IQR) and compared by Kruskal-Wallis (K-W) test. Pairwise comparisons are presented as capital letters (similar letters = no significant difference, while different letters = significant difference).

It supports the current results the results of a number of studies that have announced increased expression of VEGF in pre-eclampsia (20-21). On the contrary, some studies have shown reduced expression of VEGF expression in placental tissues of preeclampsia(22). Kim et al. and Bates, revealed that , expression of the low expressions of VEGF in both the level of mRNA and protein in the tissues of the placenta in the third trimester of pregnancy for patients with PE (n = 20) compared with an equal number of standard controls (p<0.05) (22-23). In addition, some studies revealed an expression of VEGF is variable in the tissues of the placenta in women PE (6,7).
The results of the discrepancy may be observed between these studies because of the phenotypic classification of PE chosen, the variations in sample size, the timing of taking placenta tissue samples, the method of measuring the amount of VEGF, Imaging results and references taken for comparison. There are many genetic and environmental factors that may change for pre-eclampsia gene expression. Also, the pathophysiology of the expression of VEGF in the remote PE to decode the full encryption code with many of the assumptions available to explain the different results obtained (21). The present study showed that, PLGF maternal serum level was statistically significantly higher in control group as compared to either of the two PE groups (p<0.0005) and there is no statistically significantly difference between sPE and mPE (figure 4). Low level of PI GF than normal due to the lack of re-formation of blood vessels of the arteries of the spiral ultimately leads to increased ischemia of the placenta, which may be involved in the following adverse pregnancy outcomes: SGA, PE, HELLP (hemolysis, increased liver enzymes, low platelets) (syndrome) 15,24,25). Our study correspond with many of the studies that have shown that the level of PI GF low in complex pregnancies due to PE (26, 27, 28).
no distinction in VEGFR2 (37,29,31). This study presented a significant reduction in both the expression of VEGFR-1 and VEGFR-2 (P <0.01) in PE contrasted to the control group (figures 5 and 6), which is consistent with a previous study (6,32,21,33), and in serum concentration VEGFR2 (34,35).

Inadequate placentation in PE prompts hypoxia, one of the variables known to decrease VEGFR-2 expression in the endothelial cells. Besides, the low oxygen likewise prompts expanded circling sFlt-1, a splice variant of VEGFR-1 which does not have the transmembrane and cytoplasmic areas. sFlt-1 antagonizes the proangiogenic effects of both VEGF and PlGF by binding and in this manner forestalling collaboration with their receptors (18). Moreover, a negative connection with an expansion in sFlt-1 and decrease in VEGFR-2 expression in the PE bunch is as per a past report by Nevo et al., sFlt-1 is known to tie VEGF prompting diminished accessibility of VEGF (38). Diminished accessibility of free VEGF may have brought about diminished VEGFR-2 mRNA expression in PE as free VEGF stimulates VEGFR-2 synthesis alongside its dealing at the outside of endothelial cells (36).

**Fig. 5:** Compared FLT-1 gene expression between the three study groups

**Fig. 6:** Compared KDR gene expression between the three study groups

*Data are presented as mean ± SD and compared by One-Way ANOVA. **Data are presented as frequency (percentage) and compared by Chi-Square test (Monte Carlo significance).

***Data are represented as median (IQR) and compared by Kruskal-Wallis (K-W) test. Pairwise comparisons are presented as capital letters (similar letters = no significant difference, while different letters = significant difference).

5. **CONCLUSION**

Vascular and angiogenesis factors are helpful in reducing pre-eclampsia in women with risk factors and early discovery of pre-eclampsia for diagnosis and anticipating the improvement of complications accurately. There is a need for more studies to understand the molecular defect in the pre-eclampsia and the role of the vital signs of the composition of these angiogenic biomarkers to diagnose the condition of obstetric and predictable treatment.
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