The effect of Preeclampsia on some biochemical parameter

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الخلاصة

دراسة ميدانية تهدف إلى التعرف على تسمم الحمل وعلاقته في وظائف الكلى والايض للماء والأملاح **والهدف من الدراسة**: الاستفادة من نسبة الكالسيوم والفسفور كعلاقة في تحديد شدة تسمم الدم، تم إجراء الدراسة في مستشفى الولادة والأطفال في مدينة الديوانية للفترة من1/1ولغاية 2011/6/30. كان عدد الحالات 96 حالة وقد كانت 20 حالة حمل طبيعي و25 حالة بوجود ارتفاع ضغط الدم و20 حالة بوجود تسمم دم بسيط و31 حالة بوجود تسمم دم شديد وقد تم اخذ عينات من إدرار كل مريضة لغرض إيجاد الكالسيوم والفسفور في الإدرار.

وبعد إجراء العمليات إلاحصائية وجد ما يلي: قلة في إفراز الكالسيوم والفسفور في الحالات الشديدة مع عدم وجود تغييرات ملموسة في الحالات البسيطة وارتفاع ضغط الدم اي يمكن الاعتماد على نسبة الكالسيوم والفسفور في تحديد شدة تسمم الدم في الحوامل.

ونستنتج من ذلك إن انخفاض مستوى الكالسيوم والفسفور بالإدرار يمكن أن يستخدم كعلامة لتسمم الحمل

Abstract

This survey research to identify the effect of preeclampsia, alterations in renal function, electrolyte and water metabolism are common findings.

To evaluate the usefulness of urine calcium and phosphorus levels as a marker of severity in preeclampsia.

Our study was carried at the Hospital of Maternity and pediatrics in Aldiwanyia city from 1/1-30/6 -2011.

The total number of all cases 96, among them 20 women with normal pregnancy, 25 cases with hypertension, 20 cases with mild preeclampsia, 31 cases with sever preeclampsia. Every patient was send for urinary calcium and phosphate were determined by the Kramer-Tisdall and phosphomolibidic acid method, respectively.

In the Results There was significant decrease in excretion of calcium and phosphorus in sever preeclampsia[p < 0.001, p < 0.01] respectively as a result of decrease glomerular filtration rate. While in mild cases and pregnancy induced hypertension cases there's no significant change.

Conclusion: Urine calcium and phosphorus level is significant determinant of severity of preeclampsia and may be considered as usefulness marker for predicting the level of renal impairment.

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Introduction

Preeclampsia(toxemia of pregnancy) is a multisystem disorder of unknown cause that is unique to human pregnancy. The incidence is reported to be between 2% and 7%, depending on the population[1]. Preeclampsia occurs more frequently in primigravidas. The reported rate ranges from 6% to 7% in primigravidas and from 3% to 4% in multiparous patients[2].

Criteria for the Diagnosis of Preeclampsia

SBP >140 mm Hg and/or DBP >90 mm Hg on two occasions at least 6 hours apart, typically occurring after 20 weeks gestation and protein urea of 300 mg in a 24-hours urine collection or >1+ on two random sample urine dipsticks at least 6 hours apart (no more than 1 week apart)[3].

Preeclampsia affects approximately 3% of all pregnancies worldwide, with onset of symptoms in the late second or third trimester most commonly after the 32nd week. Some women will experience preeclampsia as early as 20 weeks ,preeclampsia may also occur in the immediate post-partum period. This is referred to as postpartum preeclampsia[4].. The most dangerous time for the mother is the 24–48 hours postpartum and careful attention should be paid to preeclampsia signs and symptoms[5].

Risk Factors for Preeclampsia

Null parity, Family history of preeclampsia and eclampsia, Extremes of maternal age, Prior pregnancy complicated by preeclampsia, - Chronic or vascular disease[pregestational diabetes, renal disease chronic hypertension, rheumatic disease, connective tissue disease][6]., Molar pregnancy, Fetal hydrops, Multifetal gestation, Obesity and insulin resistance, Antiphospholipid antibody syndrome and thrombophilia[7]., Fetal aneuploidy, Maternal infection, Maternal susceptibility genes, Partner related factor [limited sperm exposure, doner insemination[Oocyte and embryo donation][8].

Classification of preeclampsia[9&10]

Preeclampsia can be classified as mild and sever. * Sever preeclampsia can be characterized:

1-Asystolic blood pressure greater than 160mmHg or diastolic blood pressure of 110 mmHg on bed rest.

2-The presence of significance proteinurea. Marked protein urea is defined as 5g or more of protein in 24 hours urine collection.

3-Sever preeclampsia may be associated with oliguria.

4-Cerbral or visual disturbance .

5-Pulmonary oedema or cyanosis.

6-Epigastric or right upper quadrant abdominal pain .

7-Impaired liver function.

8-Thrombocytopenea.

9-Intrauterine growth retardation.

*Mild preeclampsia the hypertension and protein urea are present but not to that extreme levels and the patient has no evidence of other organ dysfunction. shallowly implanted placenta which becomes hypoxic leading to an immune reaction characterized by secretion of unregulated inflammatory mediators from the placenta, and acting on the vascular endothelium[11].. In many cases of the preeclampsia syndrome, however, the maternal response to the placenta appears to have allowed for normal implantation. It is possible that women with higher baseline levels of inflammation stemming from underlying conditions such as chronic hypertension or autoimmune disease may have less tolerance for the inflammatory burden of pregnancy[12].

Material and method

This prospective study was carried at the Hospital of Maternity and Pediatrics in AL-diwanyia city from 1/1- 30/6 -2011.

The100 cases were divided into four groups:21 of normal pregnancy, 26 with hypertension and 20 with mild preeclampsia and 33pregnant women with severe preeclampsia.

The following inclusion criterias were followed

1-preeclampsia was diagnosed by blood pressure elevation equal or more than 140/90 ml in combination with proteinuria++ and /or oedma after 20 weeks gestation in previously normotensive non protein uric patient.

2-Age group from 14-44 years . 3-Parity less than 10.

4-Gesstation from 27 -40 week. 5-Singleton pregnancy.

Blood pressure was recorded in the study in the sitting position with cuff that is large enough for the subjects arm on at least two occasions 6 hours apart korotkoff phase 5[k5] disappearance of sound was used to detect the diastolic blood pressure. Protein urea was diagnosed by collecting clean catch midstream urine sampled in clean dry container then urine protein was determined using the reagent strip [Albustix] reading at 2+[1 gm./l] or more was considered to be positive result for protein urea [significant protein urea] or equal to 1+[0.3gm/l] if the specific gravity is less than 10.30

Results

Table (1) Comparison of Age, GA, Blood Pressure.

• No significant differences were observed in the age of different groups.

•Lower gestational age (P < 0.01) was recorded in the sever preeclamptic patients.

• Systolic and diastolic blood related significantly (P<0.01) with severity of preeclampsia.

	PARAMETERS			
Patient groups	AGE	GA	SYS. P	DIST. P
	Mean ±SD	Mean ±SD	Mean ±SD	Mean ±SD
NORMAL (20)	27.28±6.14	37.93±1.18	117.4±9.95	76.7±9.13
HYPERTENSIVE (25)	27.93±6.387	37.09±1.94	155.0±15.56	96.9±9.70
MILD PRECLAMPSIA (20)	26.20±6.67	35.45±2.71	160.2±11.71	103.5±6.71
SEVER PRECLAMPSIA (31)	24.64±5.968	33.42±3.01	179.7±16.29	114.5±7.65
P value	0.067NS	0.000**	0.000**	0.000**

Table (2) Renal function test.

All studied biochemical parameters showed significant relation with severity of preeclampsia. Blood urea , Serum creatinine and Serum uric acid were higher (P<0.01) in the sever preeclamptic women in comparison with other groups.

	BLOOD UREA	S. CREAT	S.URIC Acid
Patient groups	Mean ±SD	Mean ±SD	Mean ±SD
NORMAL	27.238	0.695	4.774
(20)	(3.419)	(0.172)	(0.812)
HYPERTENSIVE	31.375	0.873	4.442
(25)	(4.282)	(0.295)	(1.203)
MILD PRECLAMPSIA	32.000	0.868	5.409
(20)	(6.410)	(0.293)	(1.441)
SEVER PRECLAMPSIA	39.929	1.111	6.474
(31)	(7.468)	(0.499)	(1.445)
P value	0.000**	0.001**	0.000**

Table (3) Liver function test of studied groups.

Sever preclampsia women showed a higher value of GPT and GOT than other studied groups (P<0.01) and(P<0.05) respectively. The higher value GPT and GOT strongly significant in sever preeclampsia and Serum Alkaline Phosphatase (S.A.P.) showed significant differences between studied groups.

Patient groups	Liver function Test			
	GPT	GOT	S.A.P.	
NORMAL (20)	5.762±1.972	14.333±9.947	175.143±23.277	
HYPERTENSIVE (25)	7.346±3.249	15.462±8.363	186.192±15.998	
MILD PRECLAMPSIA (20)	9.200±3.764	23.400±11.325	187.200±23.896	
SEVER PRECLAMPSIA (31)	9.945±3.152	33.009±13.610	205.091±92.503	
P value	0.000**	0.000**	0.014*	

*significant at P \leq 0.05, **significant at P \leq 0.01, using ANOVA, values are presented as mean \pm SD ; NS, not significant.

Table (4) Urine analysis of studied groups.

- Significant differences were observed in the urine calcium of different studied groups.
- Urine phosphorus and creatinine showed significant differences between studied groups. Mild and sever preclamptic womens showed lowerr value (P<0.01) of urine phosphorus and creatinine than other groups.

Patient groups		U. calcium mg/day	U. phosphor g/day	U. creatinine mg/day
NORMAL	MEAN	218.20	1.03	1.705
(20)	SD	32.045	0.152	0.784
HYPERTENSIVE	MEAN	198.92	0.892	1.431
(25)	SD	39.912	0.256	0.178
MILD PRECLAMPSIA	MEAN	110.28	0.439	0.972
(20)	SD	20.625	0.109	0.113
SEVER PRECLAMPSIA	MEAN	98.35	0.394	0.906
(31)	SD	15.597	0.147	0.125
P value		0.000**	0.00**	0.000**

Discussion

Preeclampsia is often referred to as the disease of theories, and there is more questions than answers. Preeclampsia, avascular disorder of pregnancy is a leading cause of maternal morbidity as well as perinatal morbidity and mortality. In our study we found lower gestational age was recorded in the sever preeclamptic patients these finding similar to study done by Gerther et al [13]. In normal pregnancy renal blood flow and filtration rate increase[8] while in Preeclampsia the renal blood flow and glomerular filteration rate decrease with development of toxemia [2],in our study the blood urea, serum creatinine and serum uric acid were higher in sever preeclamptic women in comparison with other group which result in the decrease of urea and creatinine excretion .These finding similar to study done by Hayachi T[5] he found the serum level of urea, creatinine and uric acid are elevated with sever preeclampsia [p<0.001, p<0.01] respectively.

In our study the sever preeclamptic women showed a higher value of GPT and GOT than other group [p<0.01] and [p<0.05] respectively. The higher value of GPT and GOT are strongly significant with the severity of preeclampsia, also serum Alkaline phosphatase [S.A.P.] showed significant differences between studied groups. these finding similar to study done by Taylor et al [3].

Renal execration of calcium and phosphate increases during pregnancy due to increase in renal filtration rate[6].Excretion usually increases during each trimester, with maximum level reached during the third trimester. Alteration



of phosphate and most notably calcium excretion are commen finding of hypertension and some renal disorder in general. There is a decrease in urinary calcium in preeclampsia [7]. Our finding in preeclampsia confirm the result of other studies [7&14]they found decrease level of calcium and phosphorus [p<0.001, p<0.01] respectively. The reason for hypercalciuria in pregnancy is probably the increased glomerular filteration rate [14]. Pedersen et al. reported that fractional excretion of calcium in preeclampsia pregnant women was lower in the third trimester than it was in normotensive pregnant women [15]. Because parathyroid hormone and calcitonon level were not altered in the patient with preeclampsia. It was concluded that decrease renal filteration rate and increase tubular reabsorption of calcium and phosphate may result in hypocalciuuria and hypophosphateuria in toxemia. In conclusion strong relation was observed between decreasing of urine calcium and severity of preeclampsia.

Lower calcium excretion may result from dietary variation. All participants in our study were on a free range diet. Because we did not advise any of our patient to alter their diets, however, we believe it is unlikely that dietary calcium intake played an important role in our findings. As a conclusion, hypocalciuria and hypophosphateuria are important features of severe preeclampsia and probably are indirectly related to the altered renal function seen in toxemia of pregnancy.

Conclusion

Urine calcium and phosphorus levels are significant determinant of severity of preeclampsia and may be considered as usefulness marker for predicting the level of renal impairment and time of delivery.

References

1. Wang Y, Walsh SW, Kay HH. (2010): Placental lipid peroxides and thromboxane are increased and prostacyclin is decreased in women with preeclampsia. Am J Obstet Gynecol 167:946-9.

2. Seligman SP. (2010). The role of nitric oxide in the pathogenesis of preeclampsia. Am J Obstet Gynecol 17(4): 944-8.

3. Taylor RN, Musci TJ, Kuhn RM Roberts JM.(2011): Partial characterization of a novel growth factor from the blood of women with preeclampsia. J Clin Endocrinol Metab 70: 1285-91.

4. Walsh SW. (2009): Preeclampsia: an imbalancein placental prostacyclin and thromboxane production. Am J ObstetGynecol, 152 : 335-340.

5. Hayachi T.(2010): Uric acid and endogenous creatinine clearance studies in normal pregnancy and toxemia of pregnancy. Am J Obstet Gynecol 71 (4) .

6. Mozdzien G, Schinninger M, Zazgornik J.(2011): Kidney function and electrolyte metabolism in healthy pregnant women. Wien. Med. Wochensch

 Taufield PA, Ales KL, Resnick LM Druzin ML, Gerther JM, Laragh JH. (2011): Hypocalciuria in preeclampsia. N Engl J Med 316 (12): 715-8 (IVSL).
Yoshida A, Morozumi K, Suganuma T,Sato K, Aoki J, Oikawa T, Fujinami T.rinary.(2011): calcium excretion in toxemia of pregnancy. Nippon Jinzo Gakkai Shi. 31: 327-34.

9. Institute of Medicine, Food and Nutrition Board. Dietary Reference Intakes.(2011): Calcium, Phosphorus, Magnesium, Vitamin D, and Fluoride. Washington, DC: National Academy Press,: pages 146-189.

10. Bringhurst, F.R. et al.(2011): Hormones and disorders of mineral metabolism. In Wilson, J.D. et al Eds. Williams Textbook of Endocrinology, 9th Edition. Philadelphia: W.B. Saunders Company,: pages 1155-1210.

11. Calvo, M.S. & Park, Y.K.(2010): Changing phosphorus content of the U.S. diet: potential for adverse effects on bone. Journal of Nutrition.; volume 126: pages 1168S-1180S (IVSL).

12.Calvo, M.S.(2000): Dietary considerations to prevent loss of bone and renal function. Nutrition; volume 16: pages 564-566.

13. Gerther JM, Coustan DR, Kliger AS, Mallette LE, Ravin N. (2011): regnancy as state of physiologic absorptive hypercalciuria. Am J Med 81: 451-5, (IVSL).

14. Sanchez-Ramos L, Sandroni S, AndresFJ, Kaunitz AM.(2010): Calcium excretion in preeclampsia. Obstet. Gynecol 77:510-3.

15. Pedersen EB, Johannesen P, KristensenS. (2009): Calcium, parathyroid hormone and calcitonin in normal pregnancy and preeclampsia. Gynecol Obstet Invest18: 156-64.