

## Original Research Article

### **Cerebroplacental Doppler Ratio and Cerebrouterine Doppler Ratio in Predicting Neonatal Outcome in Preeclamptic Pregnant Women**

#### **Abstract**

**Background:** The umbilical artery and the middle cerebral artery are two of the most important arteries in the body. Doppler ultrasound vividly depicts placental resistance and changes in foetal hemodynamics in reaction to it. The aim of this work was to assess the role of cerebroplacental ratio and cerebrouterine ratio in prediction of neonatal outcome in preeclamptic women.

**Methods:** This prospective observational study was carried out on 110 pre eclamptic women. Patients were divided into two groups: Preeclampsia with severe (n=58) and preeclampsia (n=52). All patients were subjected to laboratory testing (complete blood count (CBC), coagulation profile, liver and kidney function tests, 24 hours urine sample collection) and ultrasonographic scanning trans - abdominal sonographic examinations.

**Results:** The cut off value of CP ratio was 1.09 with sensitivity 84%, specificity 89%, PPV (positive predictive value) 94%, NPV (negative predictive value) 73% and accuracy 85% while the study cut off value of CU ratio was 1.3 with sensitivity 97%, specificity 93%, PPV 95%, NPV 95% and accuracy 95%. UTA, UMBA, MCA, CP and CU were significantly higher in Preeclampsia with severe than Preeclampsia (P value <0.05).

**Conclusions:** Most of unfavourable neonatal outcome associated with abnormal cerebroplacental and cerebrouterine ratio so cerebroplacental and cerebrouterine ratio were complementary to each other in predicting the adverse neonatal outcomes.

**Keywords:** Cerebroplacental Doppler, Cerebrouterine Doppler, Neonatal Outcome, Preeclamptic, Pregnant

UNDER PEER REVIEW

**Introduction:**

Preeclampsia, which affects 2-5 percent of pregnancies and is one of the primary causes of maternal and foetal morbidity and mortality, is a disease defined by impaired organ perfusion due to vasospasm and endothelial dysfunction [1]. The presence of systolic blood pressure greater than 140 mm Hg or diastolic blood pressure greater than 90 mm Hg on two occasions at least 4 hours apart, as well as proteinuria or severe characteristics, is used to diagnose preeclampsia [2].

Preeclampsia causes IUGR (Intra Uterine Growth Restriction) and preterm labour by interfering with the development of uteroplacental and fetoplacental circulation, both of which are important for a healthy pregnancy [3]. Doppler velocimetry is an effective diagnostic test for foetal jeopardy that aids in the management of high-risk pregnancy [4]. It contains useful information about the fetus's hemodynamic status and is an effective diagnostic test for foetal jeopardy that aids in the management of high-risk pregnancy. Middle cerebral artery and umbilical artery Doppler ultrasound clearly shows information about placental resistance and changes in foetal hemodynamics as a result of it. The umbilical artery is a blood vessel that runs through the abdomen. Doppler reveals placental tertiary stem villi maldevelopment, which increases placental resistance [1]. Doppler imaging of blood flow through brain arteries can identify abnormalities in cerebral circulation long before significant foetal heart rate changes due to hypoxemia. The most accessible channel is the middle cerebral artery, which has been shown to show a drop in the Pulsatility index at the commencement of hypoxemia [5]. The cerebroplacental ratio (CPR) is a measure that quantifies the brain-sparing effect and provides information on how fetal cardiac output is distributed. It is calculated by dividing the middle cerebral artery (MCA) Doppler flow by the umbilical artery (UA) Doppler flow. Either the pulsatility index (PI), resistance index (RI) or

the systolic/diastolic ratio (S/D) can be used for the calculation. More recently the PI has been the computation of choice <sup>[6]</sup>.

Incomplete trophoblastic invasion to the uterine artery in the normal endovascular implantation process, whereby these cells replace the endothelium and muscular lining of blood vessels and produce an increase in vascular diameter, is one of the possibilities addressed in the aetiology of preeclampsia. This scenario results in a reduction in placental perfusion, which leads to an increase in uterine artery resistance [7]. Early detection and treatment of preeclampsia, on the other hand, can improve maternal and newborn outcomes. Given the high occurrence of preeclampsia, it is apparent that effective management is critical to improving health indices. Today, uterine arteries may be investigated using Doppler ultrasound, and high-risk pregnancies can be largely evaluated using existing indicators and existing markers [8]. The aim of this work was to assess the role of cerebroplacental ratio and cerebrouterine ratio in prediction of neonatal outcome in preeclamptic women.

### **Patients and Methods:**

This prospective observational study was carried out on 110 pre eclamptic women aged from 18 to 45 years old,  $\geq 32$  weeks of gestation, singleton viable pregnancy, BMI ( $\geq 18$ - $\leq 30$ ) kg/m<sup>2</sup> and primigravida or multigravida.

. There were adequate provisions to maintain privacy of participants and confidentiality of the data.

Exclusion criteria were women with chronic diseases like chronic hypertension, diabetes mellitus, renal diseases, liver diseases and vascular diseases, patients with fetal congenital anomalies, Intra Uterine Growth Restriction (IUGR) and Rh incompatibility, smokers, drug intake e.g., vasodilators, antiplatelet drugs, anticoagulant drugs, multiple gestations.

Patients were divided into two groups: Preeclampsia with severe (n=58) and preeclampsia (n=52)

All patients were subjected to: history (personal history, obstetric history, past history, history of present pregnancy), general examination, abdominal examination, laboratory testing (complete blood count (CBC), coagulation profile, liver and kidney function tests, 24 hours urine sample collection for proteinuria mainly) and ultrasonographic scanning.

ultrasonographic scanning: trans - abdominal sonographic examinations were performed with convex probe using Mindray DC-70 Exp and Mindray DC-30 to evaluate (fetal biometry, fetal weight).

Umbilical artery Doppler technique: with a pulsed wave Doppler system, an ultrasound scan was first carried out, a free-floating portion of the cord is identified, and the Doppler sample volume was placed over an artery and the vein, parallel to blood flow, using color-flow mapping with low-pass filter was set at 50Hz, the angle of insonation should be minimized and kept between  $150^\circ - 60^\circ$ <sup>[9]</sup>.

Middle cerebral artery technique: After that, the transducer was moved to the base of the skull, near the lesser wing of the sphenoid bone. The middle cerebral artery can be observed as a significant lateral branch of the circle of Willis using colour flow imaging. The pulsed Doppler sample gate was then installed in the vessel's centre section. The angle of insonation should be limited and kept  $< 150^\circ$  when using color-flow mapping with a low-pass filter set at 50Hz [9]. Uterine artery technique: The uterine artery's doppler velocity was measured at the location where it crossed over the external iliac artery cranial to the iliac artery crossing. The mean PI of both uterine arteries was calculated, and values of PI  $> 95$ th percentile were considered abnormal [9].

### **Statistical analysis**

IBM's SPSS statistics (Statistical Package for the Social Sciences) for windows (version 25, 2017) was used for statistical analysis of the collected data. Shapiro-Wilk test was used to check the normality of the data distribution. All tests were conducted with 95% confidence interval. P (probability) value < 0.05 was considered statistically significant. Charts were generated using SPSS' chart builder and Microsoft Excel for windows 2019. Descriptive Quantitative variables were expressed as mean and standard deviation, median, inter-quartile range, minimum and maximum as appropriate while categorical variables were expressed as frequency and percentage. ROC curve Receiver operating characteristic (ROC curve) analysis was used to find out the overall predictivity of parameter in and to find out the best cut-off value with detection of sensitivity and specificity at this cut-off value.

## Results:

Table 1 shows maternal demographics, parity and gestational age and hypertension grade among the studied cases.

**Table 1: Maternal demographics, parity and gestational age at presentation in the studied sample and hypertension grade among the studied cases**

<b>Age</b>	27.22 ± 5.97
<b>BMI (Kg/m2)</b>	26.16 ± 1.74
<b>Gestational age (weeks)</b>	34.18 ± 1.736
<b>Gravidity</b>	2.59 ± 1.599
<b>Parity</b>	1.24 ± 1.394
<b>Systolic</b>	154.32 ± 7.50
<b>Diastolic</b>	98.64 ± 7.66
<b>Preeclampsia with severe features</b>	58(52.7%)
<b>Preeclampsia</b>	52 (47.3%)

Data is expressed as mean ±SD or frequency (%), BMI: Body mass index

**Table 2** shows parity and degree of preeclampsia in the studied sample.

**Table 2: Parity and degree of preeclampsia in the studied sample**

<b>Parity</b>	<b>Primigravida</b>	<b>Multigravida</b>	<b>No. (%)</b>
<b>preeclampsia</b>	20	32	52 (47.3%)
<b>Preeclampsia with Severe features</b>	44	14	58 (52.7%)
<b>Total</b>	64 (58.18%)	46 (41.82%)	110(100%)

Table 3 shows umbilical artery and MCA and Doppler abnormality among studied cases.

**Table 3: Doppler assessment of uterine artery, umbilical artery and MCA and Doppler abnormality among studied cases**

			Range	Mean ± S. D
UTA	RI	Normal	0.43 – 0.66	0.56 ± 0.05
		Abnormal	0.7 – 0.89	0.78 ± 0.05
	PI	Normal	0.69 – 0.94	0.83 ± 0.06
		Abnormal	1.13 – 2.89	1.58 ± 0.42
UMBA	RI	Normal	0.52 – 0.74	0.66 ± 0.05
		Abnormal	0.7 – 0.76	0.73 ± 0.01
	PI	Normal	0.84 – 1.25	1.02 ± 0.11
		Abnormal	1.06 – 1.82	1.26 ± 0.17
MCA	RI	Normal	0.74 – 0.92	0.80 ± 0.04
		Abnormal	0.63 – 0.76	0.73 ± 0.02
	PI	Normal	1.11 – 2.89	1.56 ± 0.56
		Abnormal	0.99 – 1.54	1.27 ± 0.09
CP		Normal	1.03 – 2.78	1.31 ± 0.35
		Abnormal	0.68 – 1.11	0.98 ± 0.13
CU		Normal	1.21 – 3.26	1.74 ± 0.53
		Abnormal	0.43 – 1.4	0.86 ± 0.22
Doppler abnormality				
UTA		46(41.8%)		
UMBA		58(52.7%)		
MCA		78(70.9%)		
CP		36(32.7%)		
CU		48(43.6%)		

MCP: middle cerebral artery, UTA: uterine artery, PI: pulsatility index, RI: resistance index, UMBA: umbilical, middle cerebral artery, MCA: middle cerebral artery, CP: cerebroplacental ratio, CU: Cerebrouterine ratio

UTA, UMBA, MCA, CP and CU were significantly higher in Preeclampsia with severe than Preeclampsia (P value <0.05). There was insignificant difference between cerebroplacental, cerebrouterine ratio, and demographic characteristics. **Table 4**

**Table 4: Comparison between hypertension grades regarding abnormal indices, Comparison between cerebroplacental, cerebrouterine ratio and demographic characteristics**

Abnormal indices	Preeclampsia with severe (n=58)	Preeclampsia (n=52)	P value
	No. (%)	No.%	
UTA	33(56.9%)	13(25%)	0.001*
UMBA	40(69%)	18(34.6%)	0.001*
MCA	46(79.3%)	32(61.5%)	0.040*
CP	34(58.6%)	2(3.8%)	0.001*
CU	35(60.3%)	13(25%)	0.001*
	Normal CP (n=74)	Abnormal CP (n=36)	P value
Age	26.50 $\pm$ 5.82	28.69 $\pm$ 6.08	0.070
BMI (Kg/m <sup>2</sup> )	26.17 $\pm$ 1.75	26.13 $\pm$ 1.74	0.919
Systolic	153.45 $\pm$ 6.62	156.11 $\pm$ 8.87	0.080

<b>Diastolic</b>		98.85 ± 7.38	98.19 ± 8.29	0.675
<b>Parity</b>	<b>Primi</b>	43 (58.1%)	21 (58.3%)	0.982
	<b>Multi</b>	31 (41.9%)	15 (41.7%)	
		<b>Normal CU (n=62)</b>	<b>Abnormal CU (n=48)</b>	<b>P value</b>
<b>Age</b>		27.55 ± 6.09	26.79 ± 5.84	0.512
<b>BMI (Kg/m2)</b>		26.22 ± 1.54	26.07 ± 1.99	0.637
<b>Systolic</b>		154.76 ± 7.70	153.75 ± 7.26	0.487
<b>Diastolic</b>		97.58 ± 7.72	100.00 ± 7.44	0.101
<b>Parity</b>	<b>Primi</b>	35 (56.5%)	29 (60.4%)	0.676
	<b>Multi</b>	27 (43.5%)	19 (39.6%)	

MCP: middle cerebral artery, UTA: uterine artery, UMBA: umbilical, middle cerebral artery, MCA: middle cerebral artery, CP: cerebroplacental ratio, CU: Cerebrouterine ratio, Primi: Primigravida, Multi: Multigravida

Poor outcomes (except death) were significantly more frequent among cases with abnormal

CP and CU. **Table 5**

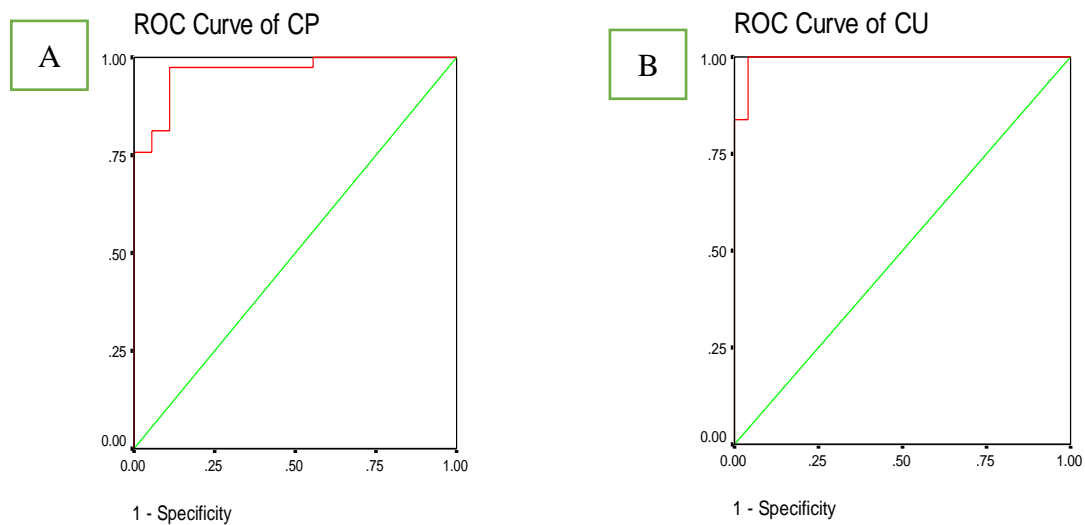
**Table 5: Neonatal outcomes among the studied cases and Comparison between cerebroplacental ratio, cerebrouterine ratio regarding outcomes**

<b>Outcomes</b>				
<b>MOD</b>	<b>NVD</b>	26(23.6%)		
	<b>CS</b>	84(76.4%)		
<b>Small GA</b>		9(8.2%)		
<b>APGAR 1 &lt; 7</b>		47(42.7%)		
<b>APGAR 5 &lt; 7</b>		34(30.9%)		
<b>NICU</b>		55(50%)		
<b>Death</b>		9(8.2%)		
	<b>Normal CP (n=74)</b>	<b>Abnormal CP (n=36)</b>	<b>P value</b>	<b>RR (95%CI)</b>
<b>Small GA</b>	3(4.1%)	6(16.7%)	0.024*	0.211 (0.050 – 0.901)
<b>APGAR 1 &lt; 7</b>	19(25.7%)	28(77.8%)	0.001*	0.099 (0.038 – 0.253)
<b>APGAR 5 &lt; 7</b>	18(24.3%)	16(44.4%)	0.032*	0.402 (0.173 – 0.935)
<b>NICU</b>	30(40.5%)	25(69.4%)	0.004*	0.300 (0.129 – 0.700)
<b>Death</b>	4(5.4%)	5(13.9%)	0.128	0.354 (0.089 – 1.410)
<b>CS</b>	54(73%)	30(83.3%)	0.230	1.852 (0.671 – 5.113)
	<b>Normal CU (n=62)</b>	<b>Abnormal CU (n=48)</b>	<b>P value</b>	<b>RR (95%CI)</b>
<b>Small GA</b>	2(3.2%)	7(14.6%)	0.031*	0.195 (0.039 – 0.987)
<b>APGAR 1 &lt; 7</b>	20(32.3%)	27(56.3%)	0.012*	0.370 (0.170 – 0.808)
<b>APGAR 5 &lt; 7</b>	9(14.5%)	25(52.1%)	0.001*	0.156 (0.063 – 0.386)
<b>NICU</b>	19(30.6%)	36(75%)	0.001*	0.147 (0.063 – 0.344)
<b>Death</b>	2(3.2%)	7(14.6%)	0.031*	0.195 (0.039 – 0.987)
<b>CS</b>	45(72.6%)	39(81.3%)	0.289	1.637 (0.656 – 4.086)

Data are represented by frequency (%), MOD: mode of delivery, NVD: normal vaginal delivery, CS: cesarean section, GA: gestational age, CP: cerebroplacental ratio, CU: Cerebrouterine ratio, NICU: neonatal intensive care unit, RR: Relative risk

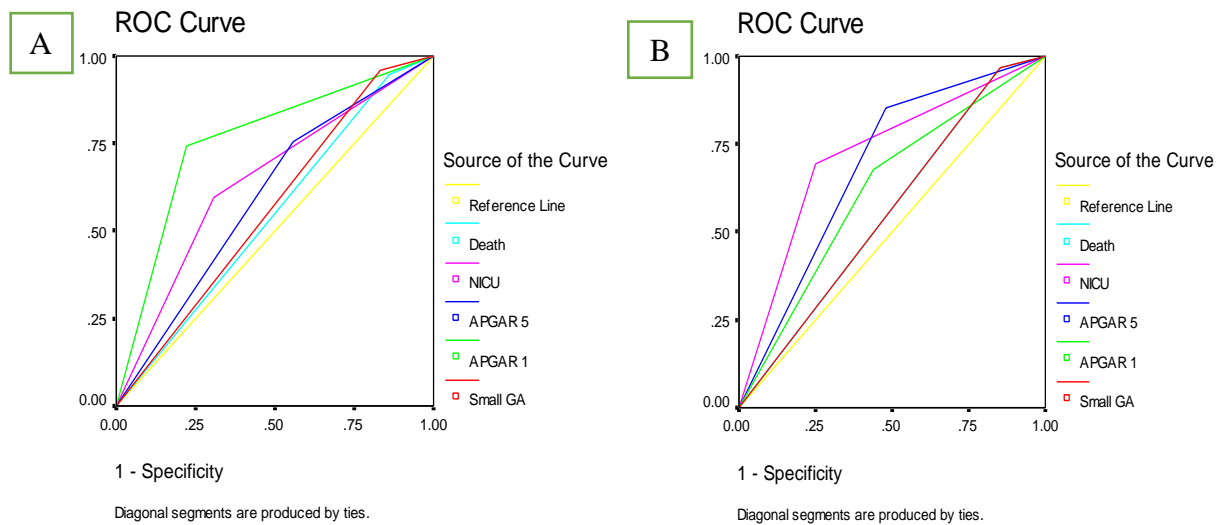
The cut off value of CP ratio was 1.09 with sensitivity 84%, specificity 89%, PPV (positive predictive value) 94%, NPV (negative predictive value) 73% and accuracy 85% while the

study cut off value of CU ratio was 1.3 with sensitivity 97%, specificity 93%, PPV 95%, NPV 95% and accuracy 95%. **Figure 1**



**Figure 1: (A) ROC Curve of cerebroplacental ratio and (B) cerebrouterine ratio**

CP ratio had higher sensitivity 96%, PPV 70%, NPV 67% and accuracy 70% in prediction of SGA while it had low specificity 17%, APGAR 1 < 7 its sensitivity was 74%, specificity 76%, PPV 87%, NPV 58% and accuracy 75%, APGAR 5 < 7 its sensitivity was 76%, specificity 56%, PPV 78%, NPV 53% and accuracy 69%, NICU its sensitivity was 59%, specificity 69%, PPV 80%, NPV 45% and accuracy 63% while in neonatal death its sensitivity was 95%, specificity 14%, PPV 69%, NPV 56% and accuracy 68%. SGA (small gestational age) CU ratio had higher sensitivity 97%, PPV 59%, NPV 78%, accuracy 61% and specificity 15%, APGAR 1 < 7 its sensitivity was 68%, specificity 44%, PPV 61%, NPV 51% and accuracy 57%, APGAR 5 < 7 its sensitivity was 85%, specificity 52%, PPV 70%, NPV 74% and accuracy 71%, NICU its sensitivity was 69%, specificity 75%, PPV 78%, NPV 65% and accuracy 72% while in neonatal death its sensitivity was 97%, specificity 15%, PPV 59%, NPV 78% and accuracy 61%. **Figure 2**



**Figure 2: (A) ROC Curve of cerebroplacental ratio and neonatal outcome and (B) cerebrouterine ratio and neonatal outcome.**

## Discussion

Preeclampsia is a pregnancy-related illness that can affect nearly every organ in the body [10]. The cerebroplacental ratio (CPR), the ratio of the pulsatility index (PI) of the MCA (middle cerebral artery) to that of the UA (umbilical artery), can detect fetal hypoxemia occurring via two different mechanisms: reduced resistance in the MCA (brain sparing effect) and increasing placental resistance [11].

In late pregnancy, the CU, a unique combination of Doppler characteristics, was the strongest Doppler predictor of delivery of an SGA or growth-restricted infant. In late pregnancy, CU performed better than did either of its constituent parameters in the prediction of birth weight <10th, <5th and <3rd centiles. Furthermore, CU demonstrated a strong biological gradient across all pregnancies with birth weight <50th centile, with an exponential increase in the rate of low CU in pregnancies with birth weight <10th centile [12].

Preeclampsia was more common in primigravida in our study, as it is in the general population. Nulliparous women are at a higher risk, which is linked to the mother's first contact with chorionic villi.

Maeda et al. [13] employed multiple logistic regression analysis to investigate the association between parity and preeclampsia in women with SLE in a single-center, retrospective record review study of 85 pregnant women. Multiparity was related with a lower risk of preeclampsia (adjusted odds ratio: 0.08; 95 percent confidence interval: 0.01–0.95), according to their findings. Das et al.,<sup>[14]</sup> made a retrospective study included 4820 pregnant women which aimed to determine the incidence of preeclampsia and distribution of risk factors of preeclampsia at Paropakar Maternity and Women's Hospital, Kathmandu, Nepal. The incidence rate of preeclampsia in the study population was 1.8% (n = 85). Higher incidence of preeclampsia was observed in primiparous women 64.7 % (n = 55) more than multipara 35.3 % (n = 30).

The current study showed that among patients with preeclampsia with severe features (n = 58), 33(56.9%) of cases had abnormal uterine artery Doppler, 40(69%) of cases had abnormal umbilical artery Doppler, 46(79.3%) of cases had abnormal middle cerebral artery Doppler. This shows that abnormal uterine (P value = 0.001), umbilical (P value = 0.001) and middle cerebral artery (P value = 0.040) ratio were significantly more frequent in patients with preeclampsia with severe features.

In agreement with our result, Adekanmi et al.,<sup>[15]</sup> conducted a prospective study was done among 98 high-risk singleton pregnant women, five were lost to follow-up, whereas 93 delivered at their institution. There was a statistically significant difference in the mean uterine artery PSV (peak systolic velocity) of pregnant women who had mild PE (preeclampsia) and that of women who developed severe PE (P = 0.024). The mean uterine S/D ratio of pregnant women with mild PE was statistically significantly lower than mean

uterine S/D (the ratio of peak systolic to end-diastolic velocity) ratio of pregnant women who had severe PE.

Deshmukh et al.,<sup>[16]</sup> divided their study into two groups-Group A “n=110” with MCA/UA PI>1.08 and Group B “n=40” with MCA/UA PI<1.08. In group B “abnormal CPR” 92.5% of cases were severe preeclampsia and 7.5% of cases were mild preeclampsia (P value <0.0001). Mean UA PI in study population was statistically significant higher in those with severe preeclampsia (p value<0.0001). On other hand mean MCA PI had no statistically significant difference (P value =0.4354) in mild and severe preeclampsia and this disagrees with our result

The present research revealed that abnormal CP (P value = 0.001) ratio and CU (P value = 0.001) ratio were significantly more common in patients with preeclampsia with severe features.

Eser et al.,<sup>[17]</sup>carried out their study on 64 preeclamptic and 131 normal pregnancies at or beyond 26 weeks of gestation and found that in 11(42.3%) of the preeclamptic pregnancies, the MCA/uterine artery PI was below the fifth percentile, related with fetal circulation redistribution. Four of these 11 cases had severe preeclampsia (36.3%) and seven had mild preeclampsia (63.6%), this disagrees with our study, while Simanaviciute and Gudmundsson<sup>[18]</sup>was conducted a cross sectional study on 231 normal pregnancies at or beyond 26 weeks of gestation to construct the reference range and a further 115 pregnancies with preeclampsia (50 mild and 65 severe) were assessed prospectively and the results were related to perinatal outcome. A low MCA/uterine artery PI ratio was seen in 30% of the mild (n = 15) and 46% of the severe (n = 30) preeclamptic cases and this is agreeing with our results.

The current study show that CP ratio had higher sensitivity in prediction of SGA while it had low specificity, NICU its sensitivity was 59%, specificity 69%, PPV 80%, NPV 45% and accuracy 63% while in neonatal death its sensitivity was 95%, specificity, while in SGA CU

ratio had higher sensitivity, NICU its sensitivity was 69%, specificity 75%, PPV 78%, NPV 65% and accuracy 72% while in neonatal death its sensitivity was 97%, specificity 15%, PPV 59%, NPV 78% and accuracy 61%.

Adiga et al.,<sup>[19]</sup> results showed that CU ratio had 54.5% sensitivity, 67.7% specificity, 47.4% PPV, 73.7% NPV and 63.2% accuracy in prediction of SGA, in poor Apgar score, CU ratio had 62.5% sensitivity, 64.6% specificity, 26.2% PPV, 89.5% NPV and 64.2% accuracy, while CP ratio had 33.3% sensitivity, 83.9% specificity, 52.4% PPV, 70.3% NPV and 66.3% accuracy in prediction of SGA, in poor Apgar score, CU ratio had 56.3% sensitivity, 84.8% specificity, 42.9% PPV, 90.5% NPV and 80.0% accuracy

In Eser et al.,<sup>[17]</sup> study, CU ratio sensitivity in prediction of NICU admission was 59.2%, specificity 73.8%, PPV 58.7% and NPV 73.4%. CP ratio sensitivity was 46.1%, specificity 87.8%, PPV 70.3% and NPV 71.9%. In Apgar 5 min < 7 prediction CU ratio had 27.2% sensitivity, 57.9% specificity, 3.8% PPV, and 89.8% NPV and CP ratio had 42.8% sensitivity, 74.3% specificity, 9.7% PPV, and 94.1% NPV. In SGA prediction CU ratio had 47.8% sensitivity, 63.9% specificity, 38% PPV, and 72.9% NPV and CP ratio had 31.1% sensitivity, 75.4% specificity, 36.2% PPV, and 70.4% NPV.

**Limitations:** One of the limitations of this study is that it was a single – center study and may not be representative of the general population. Second, small sample size selected which may make results of the study less generally applicable in a country as populous as Egypt.

### **Conclusions:**

Most of Cases with preeclampsia with severe features associated with abnormal uterine, umbilical and middle cerebral artery Doppler. Most of unfavourable neonatal outcome associated with abnormal cerebroplacental and cerebrouterine ratio so cerebroplacental and cerebrouterine ratio were complementary to each other in predicting the adverse neonatal

outcomes. Cut off value of the study cerebroplacental ratio was 1.09 and cerebrouterine ratio was 1.3.

#### **DISCLAIMER:**

Authors have declared that no competing interests exist. The products used for this research are commonly and predominantly use products in our area of research and country. There is absolutely no conflict of interest between the authors and producers of the products because we do not intend to use these products as an avenue for any litigation but for the advancement of knowledge. Also, the research was not funded by the producing company rather it was funded by personal efforts of the authors.

#### **Ethical Approval and Consent :**

Agreement for this study was obtained from the hospital's ethical committee. An informed consent was obtained from pregnant women

#### **References:**

1. Mohan S, Natarajan P, Madineni S, Rajasekhar K. Study of triple vessel wave pattern by Doppler studies in low risk and high risk pregnancies and perinatal outcome. IOSR J, Dental and Medical Sciences. 2017;16:14-23.
2. Abraham C, Kusheleva N. Management of pre-eclampsia and eclampsia: a simulation. MedEdPORTAL. 2019;15:108-12.
3. Smitha K, Sowmya K, Malathi T. Study of Doppler waveforms in pregnancy induced hypertension and its correlation with perinatal outcome. Int J Reprod Contracept Obstet Gynecol 2014;3:428-34.
4. Nagar T, Sharma D, Choudhary M, Khoiwal S, Nagar RP, Pandita A. The role of uterine and umbilical arterial doppler in high-risk pregnancy: a prospective observational study from India. Clin Med Insights Reprod Health. 2015;9:40-8.

5. PATIL V, GOWDA S, DAS S, Suma K, HIREMATH R, SHETTY S, et al. Cerebro-Placental Ratio in Women with Hypertensive Disorders of Pregnancy: A Reliable Predictor of Neonatal Outcome. *J Clin Diagn Res.* 2019;13:450-6.
6. Moreta D, Vo S, Eslick GD, Benzie R. Re-evaluating the role of cerebroplacental ratio in predicting adverse perinatal outcome. *Eur J Obstet Gynecol Reprod Biol* 2019;242:17-28.
7. Razavi M, Rashidi Fakari F, Jafari FS, Farzaneh F, Sargolzaei N. The role of uterine artery doppler ultrasound in the second trimester in predicting preeclampsia. *Int J Pediatr.* 2019;7:9405-11.
8. Tabatabaeian M, Kordi M, Dadgar S, Irani M. Evaluation of the role of serum markers and Doppler ultrasonography of uterine artery in predicting preeclampsia at the second trimester of pregnancy: A systematic review. *Iran J Obstet Gynecol Infertil.* 2018;21:76-85.
9. El Guindy AE, Nawara M, ElSanter O. Cerebro-placental ratio and cerebrouterine ratio in predicting neonatal outcome in pre-eclamptic pregnant women. *Int J Reprod Med Gynecol.* 2018;4:22-7.
10. Jeyabalan A. Epidemiology of preeclampsia: impact of obesity. *Nutr Rev.* 2013;71:18-25.
11. Vollgraff Heidweiller- Schreurs C, De Boer MA, Heymans M, Schoonmade L, Bossuyt P, Mol B, et al. Prognostic accuracy of cerebroplacental ratio and middle cerebral artery Doppler for adverse perinatal outcome: systematic review and meta- analysis. *Ultrasound Obstet Gynecol.* 2018;51:313-22.
12. MacDonald TM, Hui L, Robinson AJ, Dane KM, Middleton AL, Tong S, et al. Cerebral-placental-uterine ratio as novel predictor of late fetal growth restriction: prospective cohort study. *Ultrasound Obstet Gynecol.* 2019;54:367-75.
13. Maeda Y, Kaneko K, Ogawa K, Sago H, Murashima A. The effect of parity, history of preeclampsia, and pregnancy care on the incidence of subsequent preeclampsia in multiparous women with SLE. *Mod Rheumatol* 2021;31:843-8.

14. Das S, Das R, Bajracharya R, Baral G, Jabegu B, Odland JØ, et al. Incidence and risk factors of pre-eclampsia in the paropakar maternity and women's hospital, Nepal: A retrospective study. *Int J Environ Res Public Health*. 2019;16:357-60.
15. Adekanmi AJ, Roberts A, Akinmoladun JA, Adeyinka AO. Uterine and umbilical artery doppler in women with pre-eclampsia and their pregnancy outcomes. *Niger Postgrad Med J*. 2019;26:106-12.
16. Deshmukh V, Yelikas K, Deshmukh P. Cerebral-umbilical Doppler ratio as predictor of perinatal outcome in pregnancies with hypertension disorders. *J Evol Med Dent Sci*. 2013;2:7366-72.
17. Eser A, Zulfikaroglu E, Eserdag S, Kılıc S, Danisman N. Predictive value of middle cerebral artery to uterine artery pulsatility index ratio in preeclampsia. *Arch Gynecol Obstet*. 2011;284:307-11.
18. Simanaviciute D, Gudmundsson S. Fetal middle cerebral to uterine artery pulsatility index ratios in normal and pre-eclamptic pregnancies. *Ultrasound Obstet Gynecol*. 2006;28:794-801.
19. Adiga P, Kantharaja I, Hebbar S, Rai L, Guruvare S, Mundkur A. Predictive value of middle cerebral artery to uterine artery pulsatility index ratio in hypertensive disorders of pregnancy. *Int J Reprod Med*. 2015;2015:502-7.