

Original Research Paper

# ROLE OF CEREBROPLACENTAL RATIO OF DOPPLER VELOCIMETRY TO DETERMINE IMMEDIATE PERINATAL OUTCOME IN HYPERTENSIVE DISORDERS OF PREGNANCY IN A TERTIARY CARE HOSPITAL

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## ABSTRACT:

**BACKGROUND:** Hypertensive disorders are the most common pregnancy related complications with an incidence of 5-10% of all pregnancies and has a substantial impact on mother and perinatal morbidity and mortality. Doppler evaluation of blood flow through the cerebral vessels will allow for the detection of altered cerebral circulation even before significant fetal heart rate changes due to hypoxemia. In this study, we aimed to To study determine immediate perinatal outcome in hypertensive disorders of pregnancy using cerebroplacental ratio by combining vessels like umbilical artery and middle cerebral artery. **METHODOLOGY:** A total of 200 participants are selected sure of their date of the last menstrual period and had regular menstrual cycles previously. Their gestational age is further corroborated by first trimester ultrasound. All the antenatal women planned on induction or presented in labor are examined by routine ultrasound scan and Doppler subsequently. Blood flow velocity waveforms of UA and MCA are obtained with pulsed Doppler ultrasound equipment. The peak systolic, end diastolic and mean velocity are recorded from these vessels and Umbilical artery S/D ratio, PI, RI and Middle Cerebral Artery PI, RI, S/D and CPR are calculated. **RESULTS:** In this study, most of the women were in the age group 20 – 29 yrs. Most of them were gestational hypertension 55%, followed by preeclampsia and eclampsia contributing to 40.5% and the remaining were chronic hypertensives and with superimposed preeclampsia. 16% of the babies had NICU admissions for more than 24hrs for complications like low birth weight with foetal distress, birth asphyxia, meconium aspiration syndrome and-hypothermia. The overall diagnostic accuracy of predicting adverse perinatal outcome is 79.50% with Cerebroplacental ratio and 70% with UA PI which proves the former to be a better predictor. **CONCLUSIONS:** The study states that cerebroplacental ratio is a better tool for assessing immediate perinatal outcome. It has good accuracy for determining low birth weight and APGAR <7. Compared to other methods of foetal monitoring, the use of Doppler has proved to be more sensitive in detecting foetal compromises early and helps in aiding appropriate and timely delivery.

**Keywords:** Hypertensive disorders of pregnancy, middle cerebral artery, umbilical artery, cerebroplacental ratio, perinatal outcome

## INTRODUCTION:

Hypertensive disorders are the most common pregnancy related complications with an incidence of 5-10% of all pregnancies and has a substantial impact on mother and perinatal morbidity and mortality.<sup>(1)</sup> It is characterized by a syndrome stemming from decreased

organ perfusion which is secondary to vasospasm and endothelial pathology.<sup>(2)</sup>

The earliest pathology is scarce infiltration of spiral arteries by trophoblasts, resulting in their impaired conversion to utero-placental arteries. These changes are considered as a necessary part of placental

development and helps in establishing chorio-decidual blood flow in normal pregnancies.<sup>(3) (4)</sup> Hypertensive disorders of pregnancy are categorised as Gestational hypertension, Pre-eclampsia, Eclampsia, Chronic hypertension, Chronic hypertension with Superimposed Pre-eclampsia.<sup>(5)</sup> Intrauterine growth restriction (IUGR) is most common foetal complication of Hypertensive disorders of pregnancy and is due to a lack of normal placental invasion and growth, as well as decreased foetal perfusion.<sup>(1)</sup> Doppler velocimetry offers a non-invasive, objective and reliable method for assessing placental and foetal cardiovascular functions on the basis of blood flow dynamics measured in uterine, umbilical and foetal arteries. It has become an efficient diagnostic tool for detecting foetal compromise that helps in the management of high risk pregnancies.<sup>(6)</sup> The utero-placental insufficiency leads to reduced blood supply to the foetus and thus results in foetal growth restriction. The utilization of umbilical artery Doppler shows to improve foetal outcome in foetal growth restriction, in which case there is increased impedance to blood flow in placenta reflected by abnormal umbilical artery velocimetry findings.<sup>(7)</sup> Inadequate placental perfusion leads to foetal hypoxia and signs of circulation redistribution occurs to the foetal brain, adrenal glands and myocardium with a decrease in blood flow to the foetal kidneys, intestines and lower extremities which is considered as a compensatory mechanism.<sup>(8)</sup> Doppler evaluation of blood flow through the cerebral vessels will allow for the detection of altered cerebral circulation even before significant fetal heart rate changes due to hypoxemia. Middle cerebral artery is the most accessible vessel and is reported to demonstrate reduction in the Pulsatility index at the onset of hypoxemia.<sup>(9)</sup> Studies have shown that brain sparing effect reaches its

maximum by 2-3 weeks before late decelerations appear on cardio-tocography. This implies that the middle cerebral artery blood flow alterations in the group of patients with a high risk for unfavourable pregnancy outcome can be identified 2-3 weeks prior to delivery.<sup>(2) (8)</sup> Doppler ultrasound indices such as cerebroplacental ratio (CPR) evaluation are commonly used where CPR is calculated as the absolute ratio between the Doppler pulsatility indices (PIs) of foetal middle cerebral artery (MCA) and umbilical artery (UA) or as the ratio between the corresponding multiples of the median (MoM) for the respective gestational age.<sup>(10,11)</sup> It enables us to assess the foetal response to hypoxia by detecting blood flow distribution pattern in placento-umbilical and fetocerebral circulations. With progression of gestational age, the resistance in foetal circulation gradually decreases. The presence of one or more of the following conditions is defined as an adverse perinatal outcome - foetal distress as evidenced by foetal bradycardia or persistent tachycardia requiring caesarean section or instrumental delivery, meconium stained liquor, APGAR score of < 7 at 5 minutes, neonatal complications such as meconium aspiration syndrome or respiratory distress syndrome, admission to neonatal intensive care unit (NICU) and perinatal mortality.<sup>(12)</sup> By combining multiple vessels, like umbilical artery, middle cerebral artery and uterine artery the sensitivity of the Doppler studies can be significantly increased. Doppler in the fetoplacental circulation helps in not only monitoring of compromised fetus but also can help in predicting perinatal morbidity.<sup>(13)</sup> The present study was conducted to determine immediate perinatal outcome in hypertensive disorders of pregnancy using Cerebroplacental ratio.

**Table: 1. REFERENCE RANGE FOR UMBILICAL ARTERY PULSATILITY INDEX (PI) SAMPLED AT FREE LOOP FROM 32 TO 40 WEEKS<sup>(14)</sup>**

| Gestational Age(weeks) | 95 <sup>th</sup> centile |
|------------------------|--------------------------|
| 32                     | 1.25                     |
| 33                     | 1.22                     |
| 34                     | 1.20                     |
| 35                     | 1.18                     |
| 36                     | 1.16                     |
| 37                     | 1.14                     |

|    |      |
|----|------|
| 38 | 1.12 |
| 39 | 1.10 |
| 40 | 1.09 |

**Table 2 REFERENCE RANGE FOR MIDDLE CEREBRAL ARTERY PULSATILITY INDEX FROM 32 TO 39 WEEKS <sup>(14)</sup>**

| Gestational Age(weeks) | 5 <sup>th</sup> centile |
|------------------------|-------------------------|
| 32                     | 1.61                    |
| 33                     | 1.58                    |
| 34                     | 1.53                    |
| 35                     | 1.47                    |
| 36                     | 1.39                    |
| 37                     | 1.30                    |
| 38                     | 1.20                    |
| 39                     | 1.10                    |

**AIMS AND OBJECTIVES:**

**AIM:**

To determine the immediate perinatal outcome in hypertensive disorders of pregnancy using cerebroplacental ratio.

**OBJECTIVES:**

1. To evaluate the immediate perinatal outcome using cerebroplacental ratio which is the ratio of pulsatility index of fetal middle cerebral artery and umbilical artery in hypertensive disorders of pregnancy.
2. To compare the predictive value of umbilical artery pulsatility index with cerebroplacental ratio.

**METHODOLOGY:**

After proper counselling and obtaining informed consent from each antenatal mother selected in the study population, detailed history taking and clinical examinations are performed. All the antenatal women are examined by routine ultrasound scan and Doppler subsequently. Doppler indices are calculated using the specified software supplied within the Doppler equipment. The average values of at least three consecutive waveforms are calculated. All participants selected are sure of their date of the last menstrual period and had regular menstrual cycles previously. Their gestational age is further corroborated by first trimester ultrasound. When a discrepancy of more than seven days is detected, first trimester ultrasound is used to calculate gestational age. Women selected for

the study are subjected to complete general, physical, obstetric and pelvic examinations. Regular obstetric scan, foetal parameters obtained, AFI assessed, biweekly NST done. Blood flow velocity waveforms of UA and MCA are obtained with pulsed Doppler ultrasound equipment. The peak systolic, end diastolic and mean velocity are recorded from these vessels and Umbilical artery S/D ratio, PI, RI and Middle Cerebral Artery PI, RI, S/D and CPR are calculated.

**Umbilical artery Doppler technique:** The transducer is positioned on the pregnant mother's abdomen overlying the foetus and the waveforms from the umbilical artery and vein are adjusted to acquire the distinctive waveforms. An ultrasound scan is performed with a pulsed wave Doppler device, a free-floating section of the cord is located, and the Doppler sample volume is placed over the artery and vein, parallel to the blood flow. Colour-flow mapping with low-pass filter was set at 50Hz. The angle of the foetal Doppler insonation should be kept to < 45°. This gives an optimal umbilical artery Doppler recording.<sup>(15)</sup> With foetal inspiration (during which the foetal abdominal wall draws inward), umbilical venous blood flow increases, and with expiration, it decreases (during which the wall moves outward). There is also breathing-related modulation of arterial pulsatility and umbilical artery Doppler studies should be avoided during foetal breathing. Following the acquisition of waveforms, at least three waveforms are averaged and

impedance indices are computed.. PI values above the 95th percentile were assumed abnormal.<sup>(16)</sup>

#### **Middle Cerebral Artery (MCA) Doppler technique:**

At the level of the biparietal diameter, a transverse view of the foetal brain is acquired (BPD). The transducer is advanced to the level of the lesser wing of the sphenoid bone to locate the base of the skull. Using color flow imaging, the middle cerebral artery can be seen as a major lateral branch of the circle of Willis. It runs anterolaterally at the borderline between the anterior and the middle cerebral fossae. The pulsed Doppler gate is placed over the middle portion of the MCA to obtain flow velocity waveforms. Using color-flow mapping with low-pass filter was set at 50Hz, the angle of insonation should be restricted at a minimum of 15 degrees.<sup>(17)</sup> During the studies, the transducer should be used with caution to apply modest pressure to the maternal abdomen, because cerebral arterial waveforms are altered when the foetal head is compressed. Doppler wave forms were acquired. PI levels below the 5th percentile are assumed abnormal.<sup>(16)</sup>

- The cerebroplacental ratio (CP) (PI ratio of the middle cerebral artery to the umbilical artery) is calculated.
- Immediate perinatal outcomes are analyzed based on
  - a. Colour of the liquor
  - b. APGAR at 5 mins
  - c. Weight of the baby
  - d. Admission to neonatal intensive care unit(NICU)
  - e. Perinatal mortality if any.

#### **MATERIALS AND METHODS:**

Study Design: Prospective Observational Study

Study Period: Dec 2019 to Aug 2021

Study Setup: Pregnant women with hypertension attending Dept of Obstetrics and Gynaecology, KGH, Visakhapatnam

Sample Size: A sample of 200 cases.

Inclusion Criteria:

- Pregnant women of age >18yrs with hypertension who are eligible to give consent for the study.
- Pregnant women  $\geq 32$  and  $\leq 40$  weeks period of gestation with hypertension.
- Singleton pregnancies
- Cephalic presentation
- Pregnancies with gestational age calculated according to Last Menstrual Period with early scans corresponding.
- Pregnant women who are willing to participate in the study are included.

Exclusion criteria:

- Gestational age uncertain.
- Polyhydramnios.
- Women with multiple pregnancies.
- Congenital foetal anomalies.
- Antepartum haemorrhage.
- Previous caesarean section.
- Fetal malpresentation.

#### **PARAMETERS STUDIED:**

Maternal:

1. Age
  - <19yrs
  - 20-29yrs
  - >30yrs
2. Parity
  - Primipara
  - Multipara
3. Gestational Age at delivery
  - $\geq 32$  - <34weeks
  - $\geq 34$  - <37weeks
  - $\geq 37$ weeks
4. Pregnancy complications
  - Gestational hypertension
  - Preeclampsia/Eclampsia
  - Chronic hypertension
  - Chronic hypertension with superimposed preeclampsia
5. Mode of delivery
  - NVD
  - LSCS

Fetal:

1. Weight of the baby
  - $\geq 2.5$ kg
  - $\geq 1.5$  - <2.5kg
  - <1.5kg
2. Duration of NICU stay
  - <24hrs
  - >24hrs
3. APGAR @5mins:
  - 8-10
  - 4-7
  - <4
4. CTG
  - Reassuring
  - Non reassuring
5. Colour of liquor:
  - Clear
  - Meconium stained
  -

#### **ETHICAL CONSIDERATIONS:**

The Institutional Ethical Committee clearance was obtained. Written and informed consent was obtained from every individual of the study. Confidentiality of every patient is maintained.

#### **STATISTICAL METHODS:**

The data obtained were entered in Microsoft Excel spreadsheet and analysis were done using Statistical Package for Social Sciences (SPSS) version 26.0. Descriptive analysis of data was done calculating mean, median and standard deviation. Categorical variables were expressed as frequencies and percentages.  $P < 0.005$  were considered statistically significant. Sensitivity, specificity, positive predictive

value, negative predictive value and diagnostic accuracy were calculated.

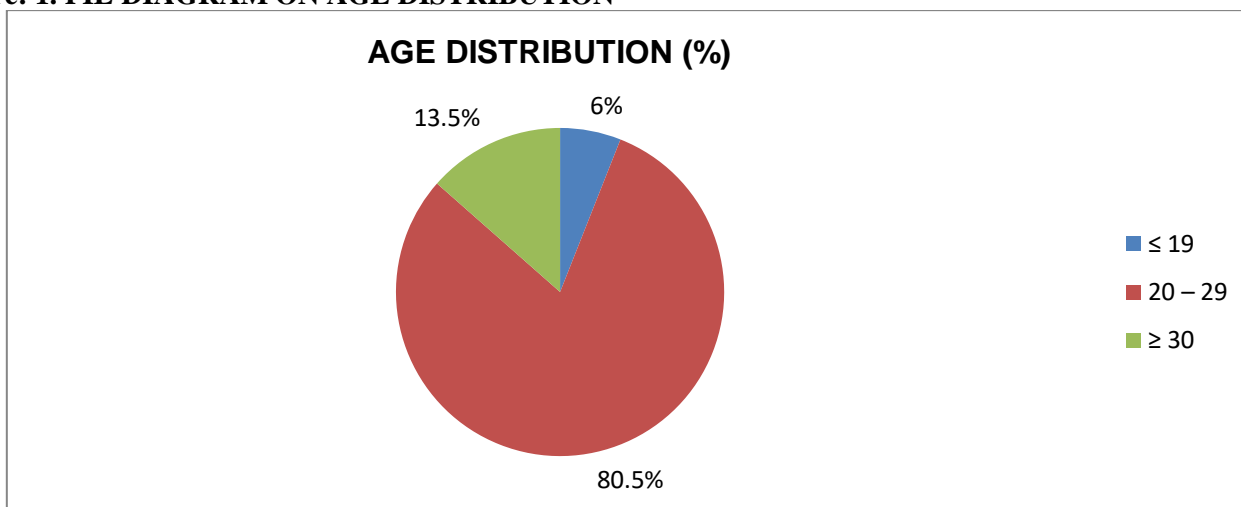
**OBSERVATIONS AND RESULTS:**

**AGE DISTRIBUTION:**

**Table: 3. AGE DISTRIBUTION**

| AGE (YEARS) | n                  | %    |
|-------------|--------------------|------|
| ≤ 19        | 12                 | 6    |
| 20 – 29     | 161                | 80.5 |
| ≥ 30        | 27                 | 13.5 |
| Total       | 200                | 100  |
| Mean ± SD   | 25.46 ± 3.48 years |      |

**Figure: 1. PIE DIAGRAM ON AGE DISTRIBUTION**



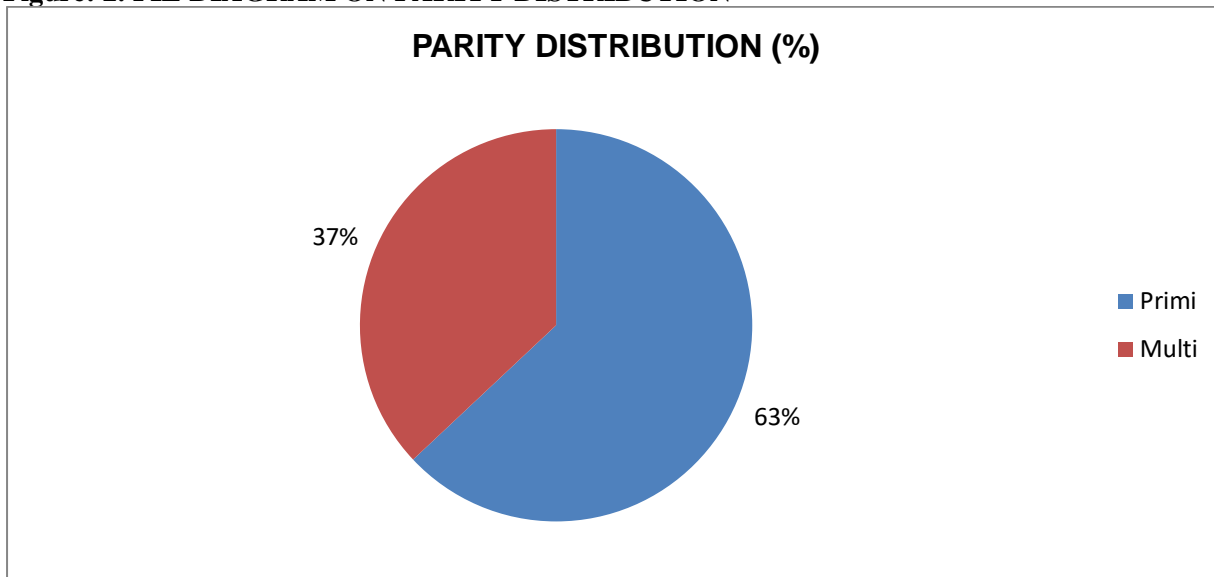
Out of 200 patients included in the study 12 were below 19 yrs ,161 were between 20 – 29 yrs , 27 above 30 yrs. The mean age of the study group was 25.4 with a standard deviation of 3.48.

**PARITY:**

**Table: 4. PARITY DISTRIBUTION**

| PARITY | n   | %   |
|--------|-----|-----|
| Primi  | 126 | 63  |
| Multi  | 74  | 37  |
| Total  | 200 | 100 |

**Figure: 2. PIE DIAGRAM ON PARITY DISTRIBUTION**



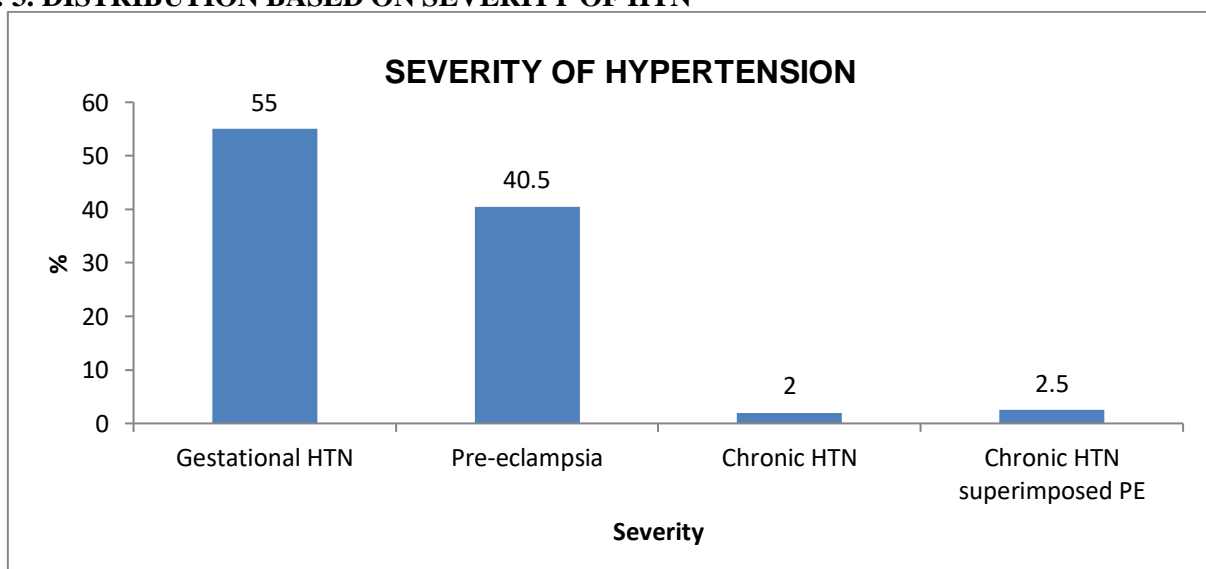
Out of 200 antenatal women, 126 were primi gravida and 74 were multigravida.

**BASED ON CATEGORY OF HTN:**

**Table: 5. DISTRIBUTION BASED ON SEVERITY OF HTN**

| CATEGORY OF HTN             | n   | %    |
|-----------------------------|-----|------|
| Gestational HTN             | 110 | 55   |
| Pre-eclampsia/Eclampsia     | 81  | 40.5 |
| Chronic HTN                 | 4   | 2    |
| Chronic HTN superimposed PE | 5   | 2.5  |
| Total                       | 200 | 100  |

**Figure: 3. DISTRIBUTION BASED ON SEVERITY OF HTN**



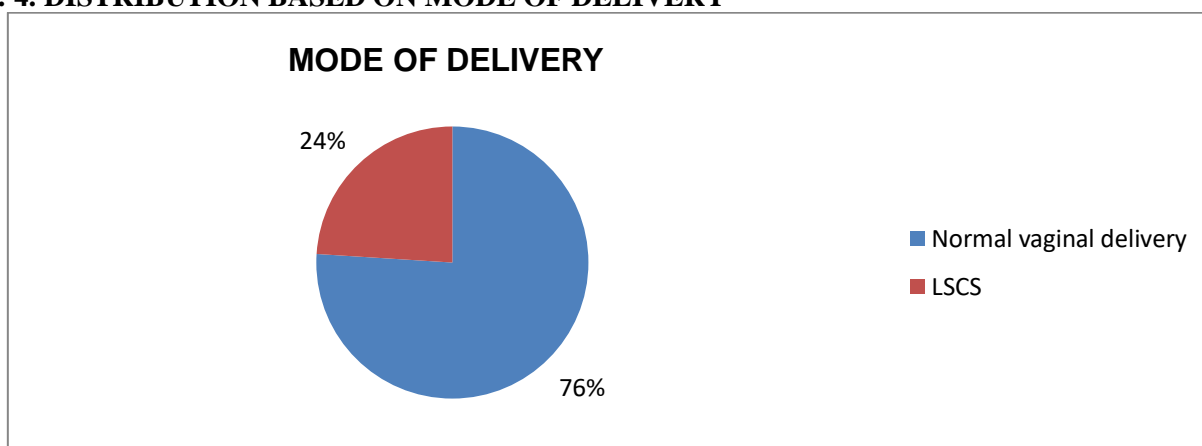
Out of 200 antenatal women, most of them had gestational hypertension 55%, followed by preeclampsia and eclampsia contributing to 40.5% and the remaining were chronic hypertensives and with superimposed preeclampsia.

**BASED ON THE MODE OF DELIVERY:**

**Table: 6. DISTRIBUTION BASED ON MODE OF DELIVERY**

| MODE OF DELIVERY        | n   | %   |
|-------------------------|-----|-----|
| Normal vaginal delivery | 152 | 76  |
| LSCS                    | 48  | 24  |
| Total                   | 200 | 100 |

**Figure: 4. DISTRIBUTION BASED ON MODE OF DELIVERY**



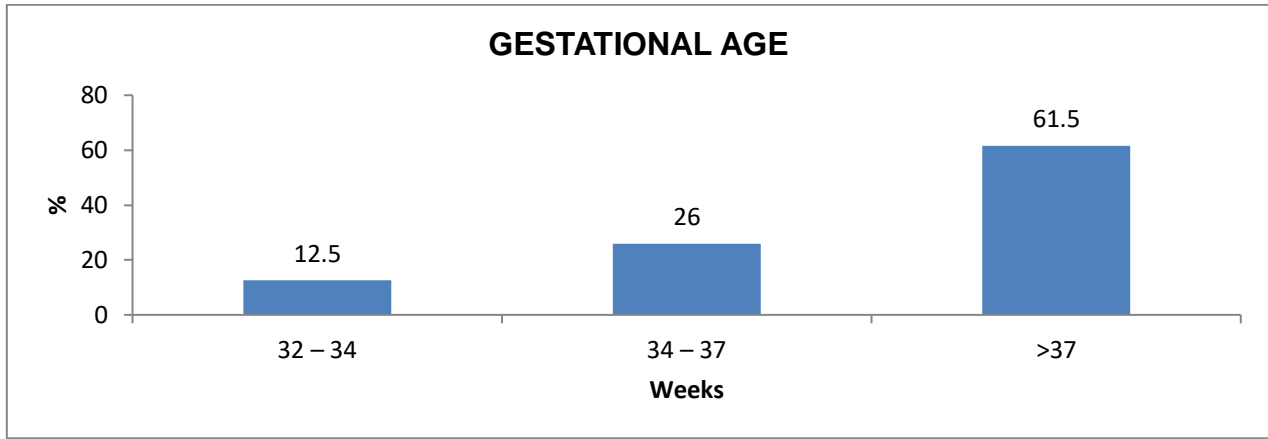
Out of 200 antenatal women, 152 delivered by vaginal route and the remaining 48 required caesarean section. Most common indication for LSCS was failed induction, foetal distress, meconium stained liquor and abnormal Doppler.

**BASED ON GESTATIONAL AGE:**

**Table: 7. DISTRIBUTION BASED ON GESTATIONAL AGE**

| GESTATIONAL AGE (WEEKS) | n                 | %    |
|-------------------------|-------------------|------|
| ≥32 – <34               | 25                | 12.5 |
| ≥34 – <37               | 52                | 26   |
| ≥37                     | 123               | 61.5 |
| Total                   | 200               | 100  |
| Mean ± SD               | 36.9 ± 2.24 weeks |      |

**Figure: 5. DISTRIBUTION BASED ON GESTATIONAL AGE**



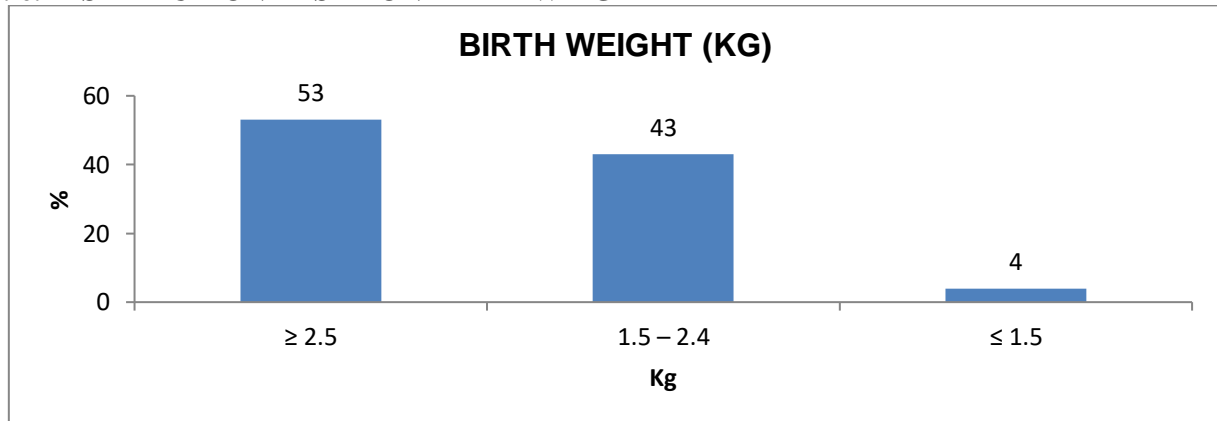
Out of 200 antenatal women, 12.5 % delivered between  $\geq 32 - < 34$  weeks, 26% delivered between  $\geq 34 - < 37$  weeks and 61.5% delivered after 37 weeks. The mean gestational age at delivery was 36.9 weeks with a standard deviation of 2.24.

**BASED ON BIRTH WEIGHT:**

**Table: 8. DISTRIBUTION BASED ON BIRTH WEIGHT**

| BIRTH WEIGHT (KG) | n                 | %   |
|-------------------|-------------------|-----|
| $\geq 2.5$        | 106               | 53  |
| 1.5 – 2.4         | 86                | 43  |
| $\leq 1.5$        | 8                 | 4   |
| Total             | 200               | 100 |
| Mean $\pm$ SD     | 2.46 $\pm$ 0.45kg |     |

**Figure: 6. DISTRIBUTION BASED ON BIRTH WEIGHT**



In the present study, out of 200 babies, 106 babies weighed more than 2.5kg (53%), 86 babies weighed between 1.5 – 2.4 and 8 babies weighed  $\leq 1.5$ kg. The mean birth weight was 2.46kg with a standard deviation of 0.45kg.

**BASED ON APGAR**

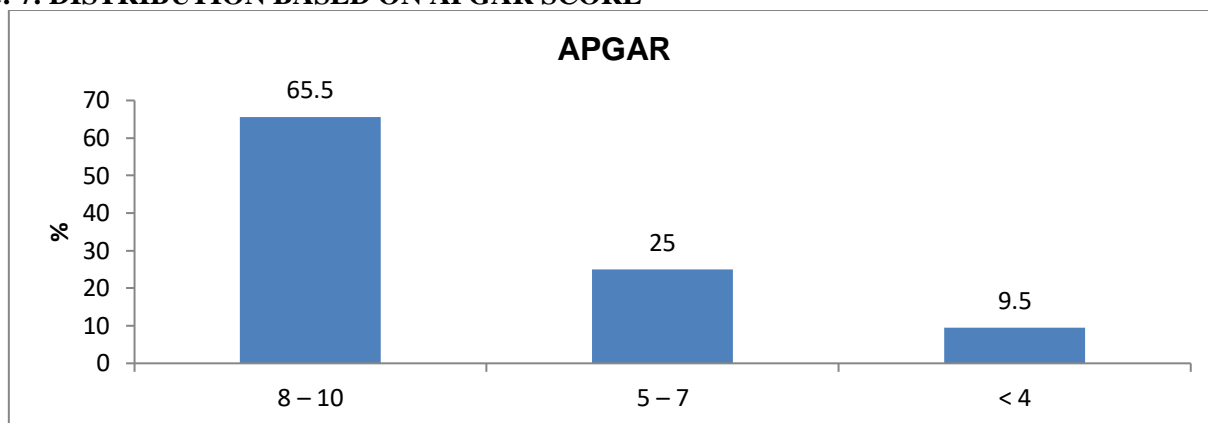
**Table: 9. DISTRIBUTION BASED ON APGAR**

| APGAR          | n          | %    |
|----------------|------------|------|
| 8 – 10         | 131        | 65.5 |
| 5 – 7          | 50         | 25   |
| < 4            | 19         | 9.5  |
| Total          | 200        | 100  |
| Median (range) | 8 (2 – 10) |      |



**Table: 10. DISTRIBUTION BASED ON CTG**

**Figure: 7. DISTRIBUTION BASED ON APGAR SCORE**

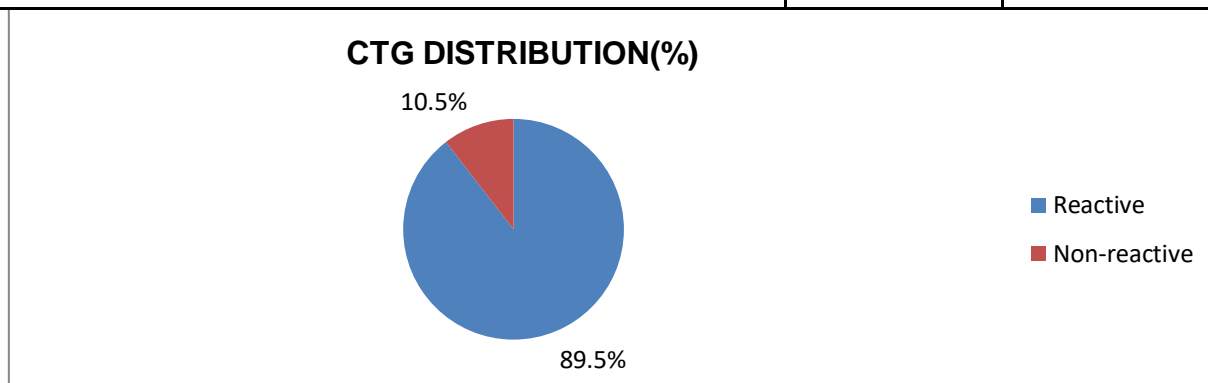


Out of 200 babies, 131 babies had APGAR >7 and 69 babies had APGAR <7. Median APGAR is 8.

**BASED ON CTG:**

**Figure: 8. PIE DIAGRAM ON CTG REACTIVITY**

| CTG            | n   | %    |
|----------------|-----|------|
| Reassuring     | 179 | 89.5 |
| Non-reassuring | 21  | 10.5 |
| Total          | 200 | 100  |



In the present study, CTG was found Reactive/Reassuring in 179 (89.5%) cases and Non reactive/Non reassuring in 21(10.5%) cases.

**BASED ON NICU STAY:**

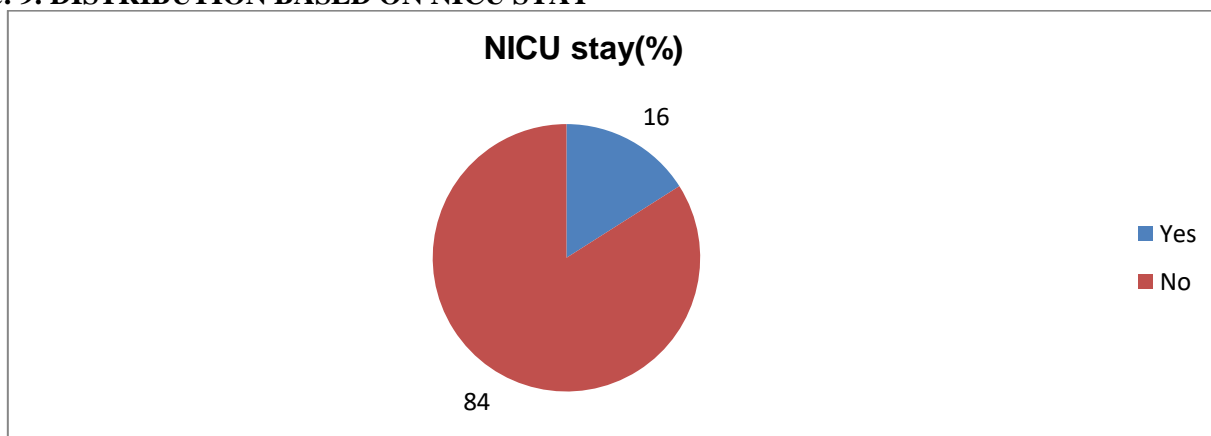
**Table: 11. DISTRIBUTION BASED ON NICU ADMISSIONS**

| NICU stay | n | % |
|-----------|---|---|
|-----------|---|---|

|     |     |    |
|-----|-----|----|
| Yes | 32  | 16 |
| No  | 168 | 84 |

| COLOUR OF LIQUOR        |     | n   | %   |
|-------------------------|-----|-----|-----|
| Clear                   |     | 158 | 79  |
| Meconium stained liquor |     | 42  | 21  |
| Total                   |     | 200 | 100 |
| Total                   | 200 | 100 |     |

**Figure: 9. DISTRIBUTION BASED ON NICU STAY**

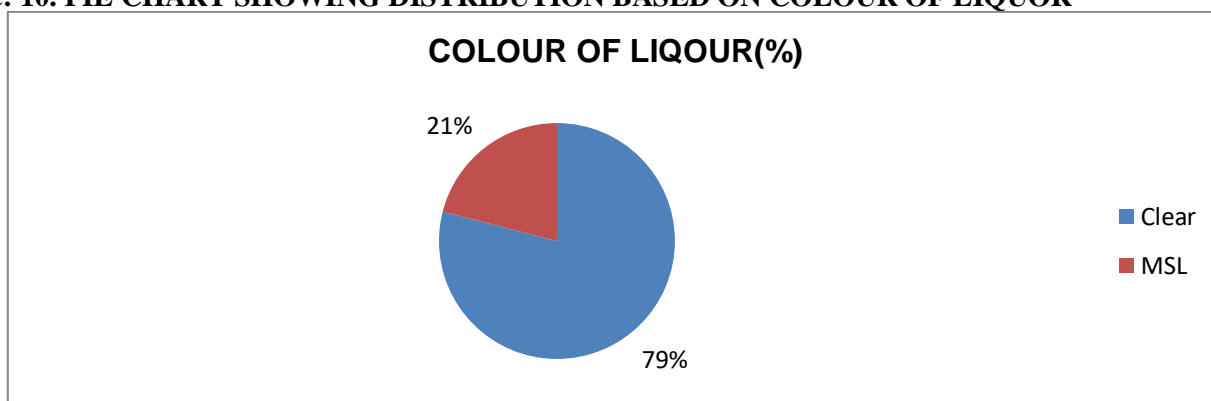


Out of 200 babies delivered, 32 babies required NICU admission for more than 24hrs. Most common complications were low birth weight with foetal distress, birth asphyxia, meconium aspiration syndrome and hypothermia.

#### **BASED ON THE COLOR OF LIQUOR:**

**Table: 12. DISTRIBUTION BASED ON COLOR OF LIQUOR**

**Figure: 10. PIE CHART SHOWING DISTRIBUTION BASED ON COLOUR OF LIQUOR**



Out of 200 deliveries, 42 babies had meconium stained liquor and 158 babies had clear liquor.

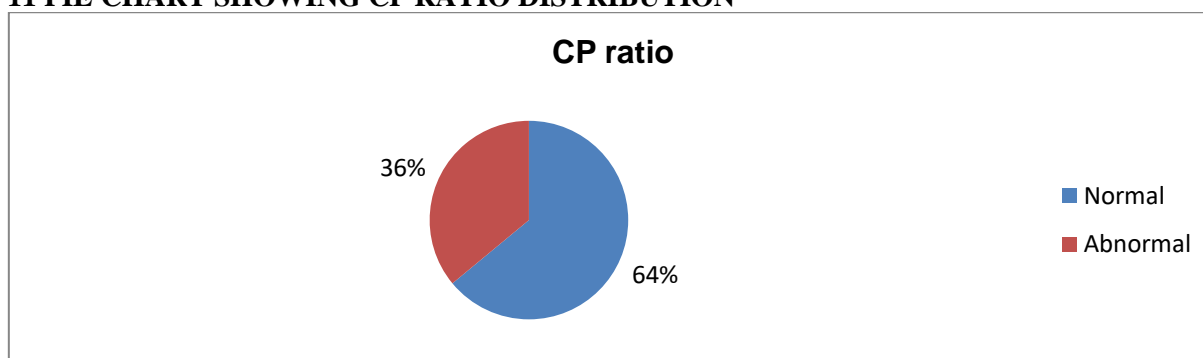
## CEREBROPLACENTAL RATIO AND PERINATAL OUTCOME:

**Table: 13. CP RATIO AND PERINATAL OUTCOME**

| CPR             | Birth weight |             |         |      | Total |     |
|-----------------|--------------|-------------|---------|------|-------|-----|
|                 | <2.5kg       |             | ≥ 2.5kg |      |       |     |
|                 | n            | %           | n       | %    | N     | %   |
| Abnormal(<1.08) | 58           | 80.6        | 14      | 19.4 | 72    | 100 |
| Normal (≥1.08)  | 36           | 28.1        | 92      | 71.9 | 128   | 100 |
| Total           | 94           | 47          | 106     | 53   | 200   | 100 |
| P value         | <0.0001      |             |         |      |       |     |
| CP RATIO        |              | n           |         | %    |       |     |
| Normal          |              | 128         |         | 64   |       |     |
| Abnormal        |              | 72          |         | 36   |       |     |
| Total           |              | 200         |         | 100  |       |     |
| Mean ± SD       |              | 1.57 ± 0.56 |         |      |       |     |

The efficacy of employing gestational age-specific reference levels of CPRs in predicting adverse outcomes was compared by using a CPR value of less than 1.08. Mean CPR in the present study is 1.57 with a standard deviation 0.56.

**Figure 11 PIE CHART SHOWING CP RATIO DISTRIBUTION**



## CEREBROPLACENTAL RATIO AND BIRTH WEIGHT:

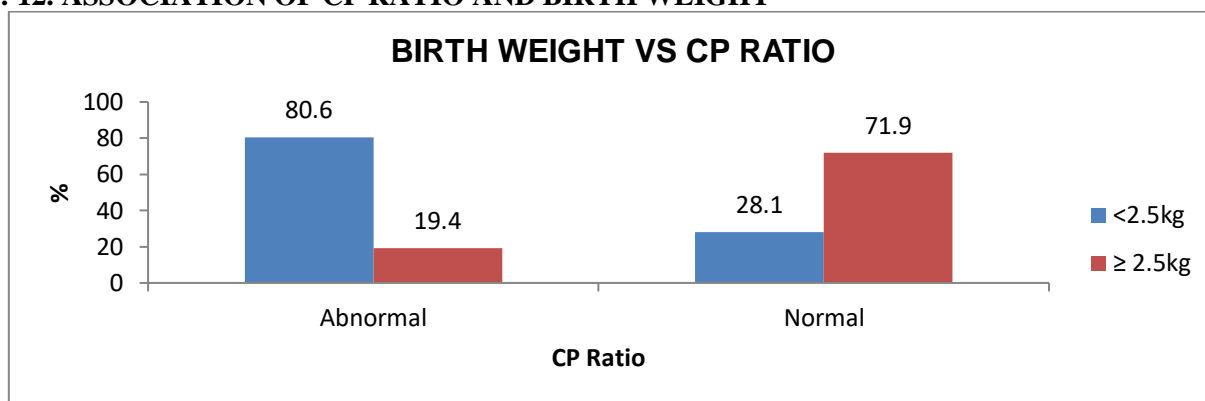
**Table: 14. CORRELATION OF CP RATIO AND BIRTH WEIGHT**

**Table: 15. STATISTICAL ANALYSIS OF CP RATIO AND BIRTH WEIGHT**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 61.70% |
| Specificity               | 86.79% |
| Positive predictive value | 80.56% |
| Negative predictive value | 71.88% |
| Accuracy rate             | 75%    |

Cerebroplacental ratio  $\leq 1.08$  has a sensitivity of 61.7% and specificity Of 86.79 % in predicting low birth weight <2.5kg which was statistically significant with a p value < 0.0001 and a diagnostic accuracy rate of 75%.

**Figure: 12. ASSOCIATION OF CP RATIO AND BIRTH WEIGHT**



**CEREBROPLACENTAL RATIO AND APGAR:**

**Table: 16. CORRELATION BETWEEN CP RATIO AND APGAR**

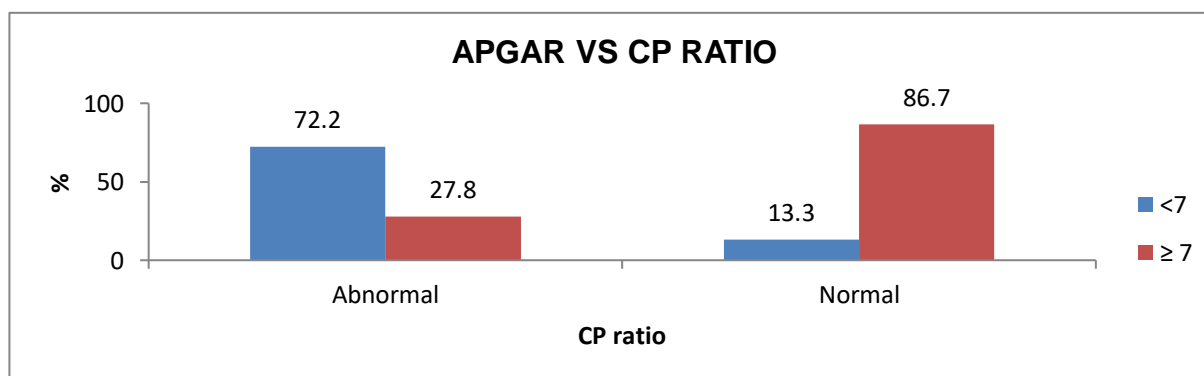
| CPR             | APGAR   |      |     |      | Total |     |
|-----------------|---------|------|-----|------|-------|-----|
|                 | <7      |      | ≥ 7 |      | N     | %   |
|                 | N       | %    | n   | %    |       |     |
| Abnormal(<1.08) | 52      | 72.2 | 20  | 27.8 | 72    | 100 |
| Normal (≥1.08)  | 17      | 13.3 | 111 | 86.7 | 128   | 100 |
| Total           | 69      | 34.5 | 131 | 65.5 | 200   | 100 |
| P value         | <0.0001 |      |     |      |       |     |

**Table: 17. STATISTICAL ANALYSIS OF CP RATIO AND APGAR**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 75.36% |
| Specificity               | 84.73% |
| Positive predictive value | 72.22% |
| Negative predictive value | 86.72% |
| Accuracy rate             | 81.50% |

Cerebroplacental ratio < 1.08 has a sensitivity of 75.36% and specificity of 84.73% in predicting APGAR <7 which was statistically significant with a p value < 0.0001 and a diagnostic accuracy rate of 81.50%

**Figure: 13. ASSOCIATION OF CP RATIO AND APGAR**



**CEREBROPLACENTAL RATIO AND COLOR OF LIQUOR:**

**Table: 18. CORRELATION OF CP RATIO AND COLOUR OF LIQUOR**

| CPR | COLOUR OF LIQUOR |   |     |   | TOTAL |   |
|-----|------------------|---|-----|---|-------|---|
|     | CLEAR            |   | MSL |   | N     | % |
|     | N                | % | n   | % |       |   |

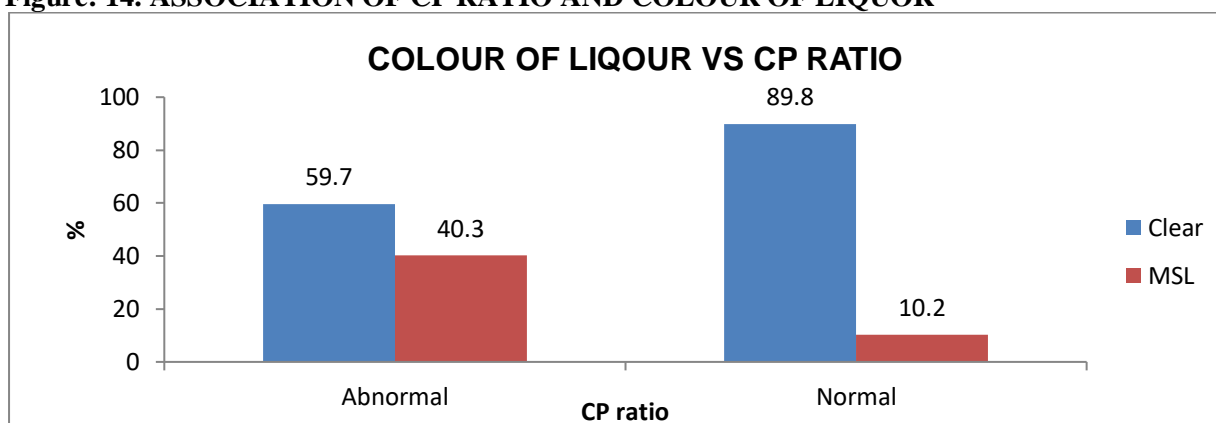
|                 |         |      |    |      |     |     |
|-----------------|---------|------|----|------|-----|-----|
| Abnormal(<1.08) | 43      | 59.7 | 29 | 40.3 | 72  | 100 |
| Normal (≥1.08)  | 115     | 89.8 | 13 | 10.2 | 128 | 100 |
| Total           | 158     | 79   | 42 | 21   | 200 | 100 |
| P value         | <0.0001 |      |    |      |     |     |

**Table: 19. STATISTICAL ANALYSIS OF CP RATIO AND COLOUR OF LIQUOR**

|                                  |        |
|----------------------------------|--------|
| <b>Sensitivity</b>               | 27.22% |
| <b>Specificity</b>               | 30.95% |
| <b>Positive predictive value</b> | 79.00% |
| <b>Negative predictive value</b> | 59.72% |
| <b>Accuracy rate</b>             | 28.00% |

Cerebroplacental ratio < 1.08 has a very low accuracy rate of predicting colour of liquor with an accuracy rate of 28% and hence is not statistically significant.

**Figure: 14. ASSOCIATION OF CP RATIO AND COLOUR OF LIQUOR**



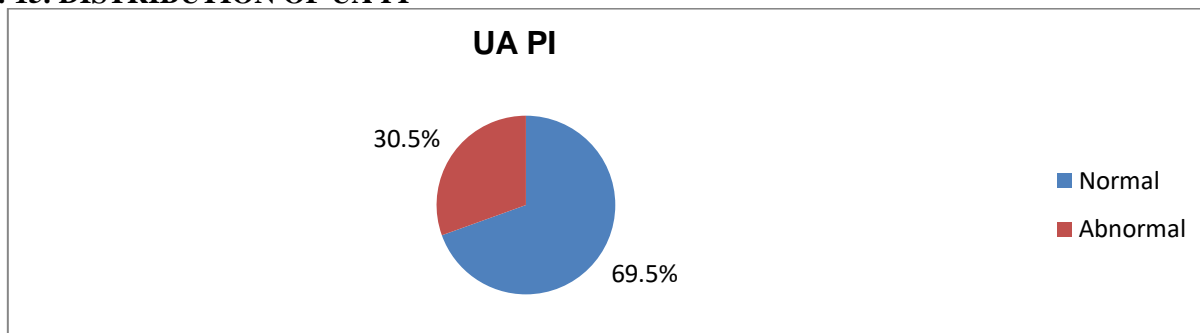
## UA PI AND PERINATAL OUTCOME:

**Table: 20. CORRELATION OF UA PI AND PERINATAL OUTCOME**

| UA PI         | n               | %    |
|---------------|-----------------|------|
| Normal        | 139             | 69.5 |
| Abnormal      | 61              | 30.5 |
| Total         | 200             | 100  |
| Mean $\pm$ SD | 1.01 $\pm$ 0.21 |      |

UA PI value were considered based on gestational age specific reference values of which 69.5% individuals showed normal value and 30.5% showed values beyond 95<sup>th</sup> percentile which is considered abnormal for the corresponding gestational age. Mean UA PI in the present study is 1.01 with a standard deviation of 0.21

**Figure: 15. DISTRIBUTION OF UA PI**



## UA PI AND BIRTH WEIGHT:

**Table: 21. CORRELATION OF UA PI AND BIRTH WEIGHT**

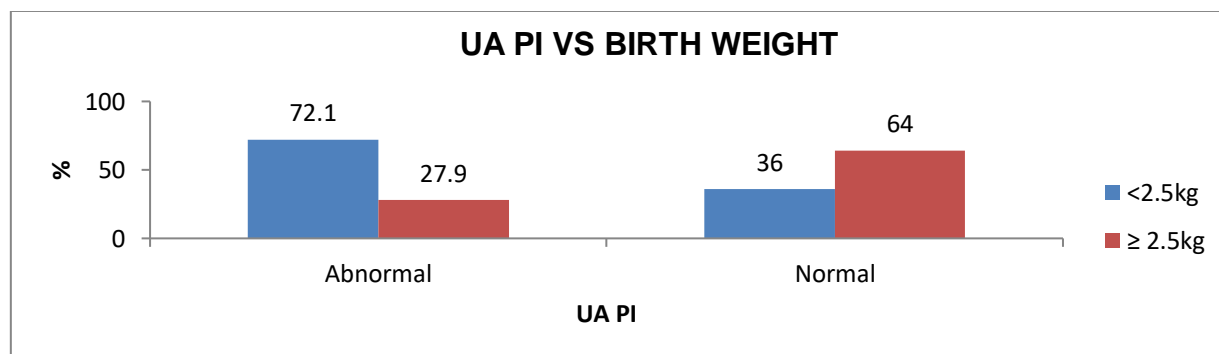
| UA PI    | BIRTH WEIGHT |      |              |      | TOTAL |     |
|----------|--------------|------|--------------|------|-------|-----|
|          | <2.5kg       |      | $\geq$ 2.5kg |      |       |     |
|          | n            | %    | N            | %    | N     | %   |
| Abnormal | 44           | 72.1 | 17           | 27.9 | 61    | 100 |
| Normal   | 50           | 36   | 89           | 64   | 139   | 100 |
| Total    | 94           | 47   | 106          | 53   | 200   | 100 |
| P value  | <0.0001      |      |              |      |       |     |

**Table: 22. STATISTICAL ANALYSIS OF UA PI AND BIRTH WEIGHT**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 46.81% |
| Specificity               | 83.96% |
| Positive predictive value | 72.13% |
| Negative predictive value | 64.03% |
| Accuracy rate             | 66.50% |

Abnormal UA PI value has a sensitivity of 46.81% in predicting low birth wt <2.5kg and specificity of >83.96% with a diagnostic accuracy rate of 66.50% for a p value<0.0001.

**Figure 16 ASSOCIATION OF UA PI WITH BIRTH WEIGHT**



### UA PI AND APGAR:

**Table: 23. CORRELATION BETWEEN UA PI AND APGAR**

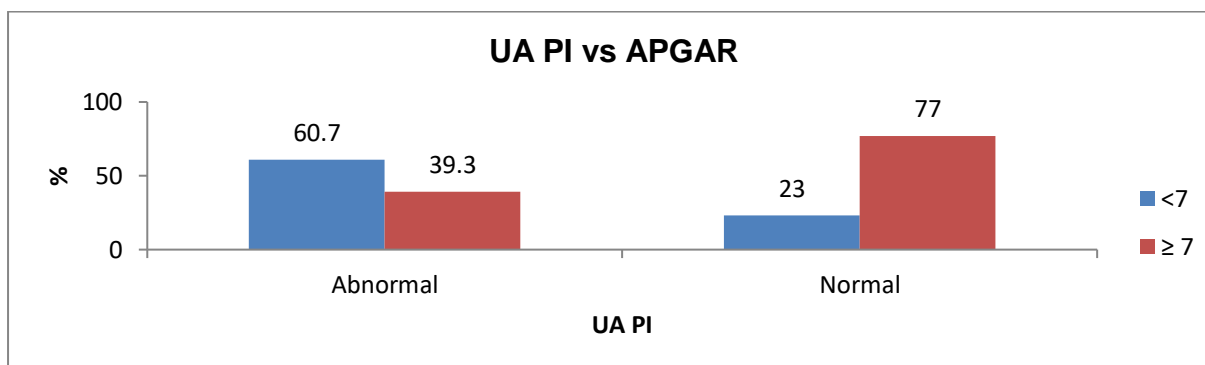
| UA PI    | APGAR   |      |     |      | TOTAL |     |
|----------|---------|------|-----|------|-------|-----|
|          | <7      |      | ≥ 7 |      | N     | %   |
|          | N       | %    | N   | %    |       |     |
| Abnormal | 37      | 60.7 | 24  | 39.3 | 61    | 100 |
| Normal   | 32      | 23   | 107 | 77   | 139   | 100 |
| Total    | 69      | 34.5 | 131 | 65.5 | 200   | 100 |
| P value  | <0.0001 |      |     |      |       |     |

**Table: 24. STATISTICAL ANALYSIS OF UA PI AND APGAR**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 53.62% |
| Specificity               | 81.68% |
| Positive predictive value | 60.66% |
| Negative predictive value | 76.98% |
| Accuracy rate             | 72.00% |

Abnormal UA PI value has a sensitivity of 53.62% and specificity of 81.68% in determining APGAR<7 with an accuracy rate of 72% for p value <0.0001.

**Figure 17 ASSOCIATION OF UA PI WITH APGAR**



### UA PI AND COLOR OF LIQUOR:

**Table: 25. CORRELATION OF UA PI AND COLOUR OF LIQUOR**

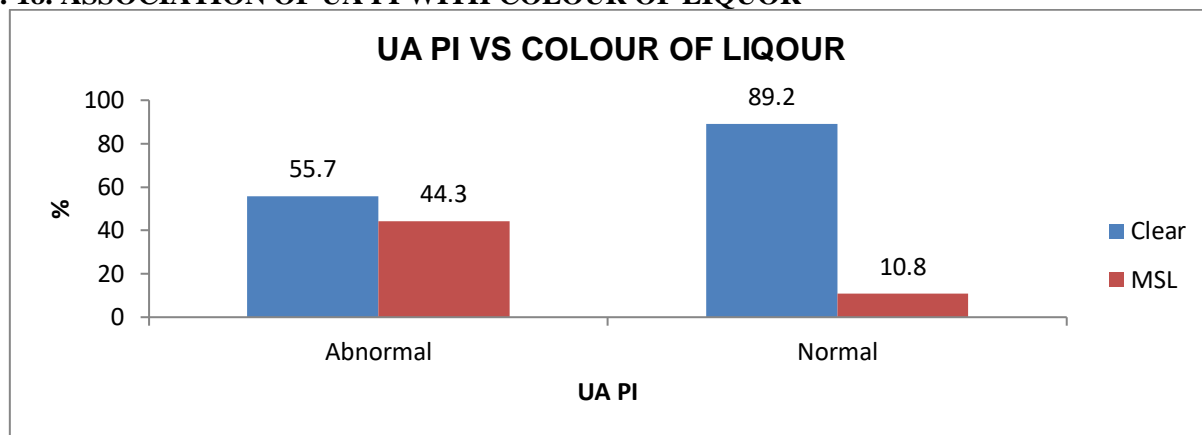
| UA PI    | COLOUR OF LIQUOR |      |     |      | TOTAL |     |
|----------|------------------|------|-----|------|-------|-----|
|          | CLEAR            |      | MSL |      | N     | %   |
|          | n                | %    | N   | %    |       |     |
| Abnormal | 34               | 55.7 | 27  | 44.3 | 61    | 100 |
| Normal   | 124              | 89.2 | 15  | 10.8 | 139   | 100 |
| Total    | 158              | 79   | 42  | 21   | 200   | 100 |
| P value  | <0.0001          |      |     |      |       |     |

**Table: 26. STATISTICAL ANALYSIS OF UA PI AND COLOUR OF LIQUOR**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 21.52% |
| Specificity               | 35.71% |
| Positive predictive value | 55.74% |
| Negative predictive value | 10.79% |
| Accuracy rate             | 24.50% |

Abnormal UA PI value has no significance in relation to color of liquor as sensitivity and specificity is 21.52% and 35.71% with very low positive and negative predictive values.

**Figure: 18. ASSOCIATION OF UA PI WITH COLOUR OF LIQUOR**



**ADVERSE PERINATAL OUTCOME:**

**Table: 27. ADVERSE PERINATAL OUTCOME**

|                 | ADVERSE PERINATAL OUTCOME | %      |
|-----------------|---------------------------|--------|
| Perinatal death | 4                         | 2%     |
| Low APGAR       | 69                        | 34.50% |
| NICU admissions | 32                        | 16%    |
| Total           | 75                        | 37.50% |

Out of 200 cases, adverse perinatal outcome is seen in 75 cases which is 37.5% of the total cases. In the present study, low APGAR, NICU admissions and perinatal death were all seen cumulatively in some cases.

**Table: 28. UA PI AND ADVERSE PERINATAL OUTCOME**

| UA PI    | GOOD OUTCOME | ADVERSE PERINATAL OUTCOME | TOTAL |     |
|----------|--------------|---------------------------|-------|-----|
|          |              |                           | N     | %   |
| Abnormal | 23           | 38                        | 61    | 100 |
| Normal   | 102          | 37                        | 139   | 100 |
| Total    | 125          | 75                        | 200   | 100 |

**Table: 29. STATISTICAL ANALYSIS OF UA PI AND ADVERSE PERINATAL OUTCOME**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 50.66% |
| Specificity               | 81.66% |
| Positive predictive value | 62.29% |
| Negative predictive value | 73.38% |
| Accuracy rate             | 70%    |

**Table: 30. CPR AND PERINATAL OUTCOME**

| CPR                  | GOOD OUTCOME | ADVERSE PERINATAL OUTCOME | TOTAL |     |
|----------------------|--------------|---------------------------|-------|-----|
|                      |              |                           | N     | %   |
| Abnormal(<1.08)      | 19           | 53                        | 72    | 100 |
| Normal( $\geq$ 1.08) | 106          | 22                        | 128   | 100 |
| Total                | 125          | 75                        | 200   | 100 |

**Table: 31. STATISTICAL ANALYSIS OF CP RATIO AND ADVERSE PERINATAL OUTCOME**

|                           |        |
|---------------------------|--------|
| Sensitivity               | 70.66% |
| Specificity               | 84.8%  |
| Positive predictive value | 73.61% |
| Negative predictive value | 82.8%  |
| Accuracy rate             | 79.5%  |



In the present study, for the 37.5% cases with adverse perinatal outcome, specificity of CP Ratio is 84.8% in predicting adverse perinatal outcome while that of UA PI is 81.6%. overall diagnostic accuracy rate is also high for CP ratio(79.5%) compared to UA PI alone (70%).

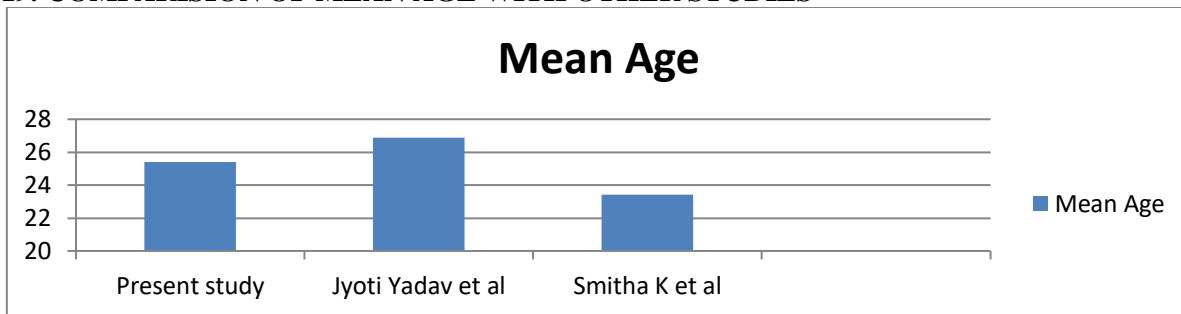
**DISCUSSION:**

**COMPARISION WITH RESPECT TO AGE:**

Out of 200 antenatal women included in the study 12 were below 19 yrs ,161 were between 20 – 29 yrs , 27

above 30 yrs. The mean age of the study group was 25.4years with a standard deviation of 3.48. The highest no.of cases were reported in the age group 20 – 29yrs.In a study conducted by Jyoti Yadav et al., 200 antenatal women with hypertensive disorder of pregnancy were recruited, maximum women 96 (48%) were within 25-30 yrs of age group and mean  $\pm$  SD age was  $26.9 \pm 4.140$ yrs.<sup>(18)</sup> In a study conducted by Smitha k et al., out of 100 cases with PIH, 58% women were between the age group 21-25 yrs. The mean age in the study is 23.41yrs with a standard deviation of 3.1yrs. <sup>(19)</sup>

**Figure: 19. COMPARISION OF MEAN AGE WITH OTHER STUDIES**

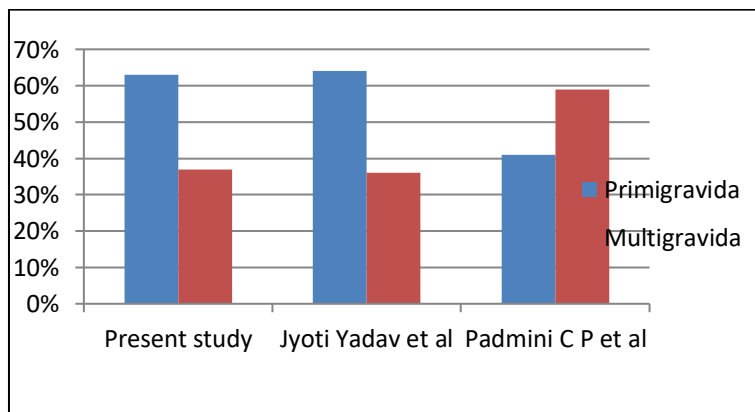


**COMPARISION BASED ON PARITY:**

In the present study, 126 were primi gravida and 74 were multigravida. In the present study, hypertensive disorders are observed twice as common in primigravida compared to multigravida which is also

seen in a study conducted by Andersch B et al. <sup>(18)</sup> In a study conducted by Jyoti Yadav et al., most of the women i.e., 128 out of 200 (64%) women were primigravida.<sup>(18)</sup> In a study by Padmini C P et al., 41%primigravida and 59% were multigravida.<sup>(20)</sup>

**Figure: 20. COMPARISION BASED ON PARITY WITH VARIOUS STUDIES**



**COMPARISION BASED ON SEVERITY OF HTN:**

In the present study, out of 200 antenatal women, most of them had gestational hypertension 55%, followed by preeclampsia and eclampsia contributing to 40.5% and the remaining were chronic hypertensives and with superimposed preeclampsia. Smitha K et al., categorised the subjects into severe PIH and mild PIH which is 62% and 38% respectively.<sup>(19)</sup> Padmini C P et

al categorised HTN into mild PIH and severe PIH which was 51.25% and 48.75% respectively.<sup>(20)</sup>

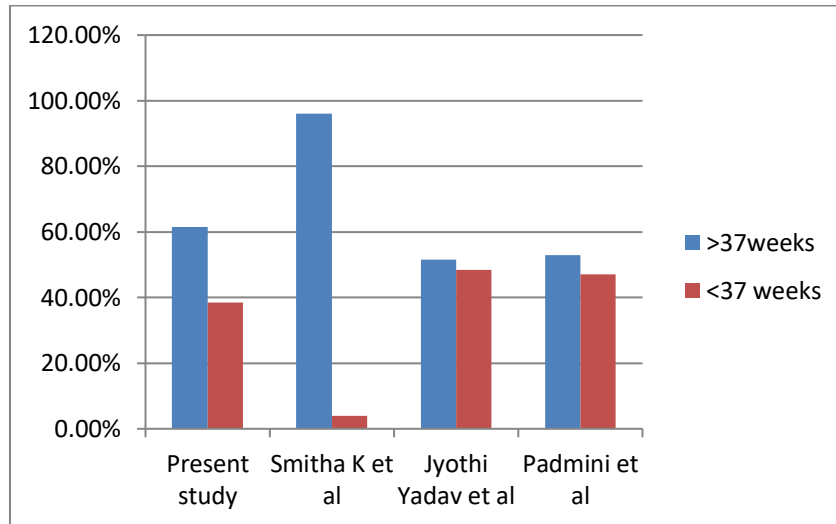
**COMPARISION BASED ON GESTATIONAL AGE:**

In the present study, the highest no of cases delivered beyond 37weeks of age. 12.5 % delivered between  $\geq 32$  – <34weeks, 26% delivered between  $\geq 34$ -<37 weeks and 61.5% delivered at 37 weeks and later. The mean gestational age at delivery was 36.9 weeks with a

standard deviation of 2.24. In a study conducted by Smitha k et al., 96% women delivered before 37 weeks and 4% delivered beyond 37weeks.<sup>(19)</sup> In a study conducted by Jyoti Yadav et al., mean gestational age at delivery was 34.2±1.95 weeks in women with abnormal Doppler while it was 37.46±1.28 weeks in normal Doppler group. 37.5% in study group were late

preterm delivery and 28% delivered <34 weeks while in control group 28% delivered were late preterm and only 11% delivered before 34 weeks.<sup>(18)</sup> In a study conducted by Padmini c.p et al, 52.9% showed gestational age >37weeks, 40% were between 33-37weeks gestation and 7.1% <32 weeks gestation.<sup>(20)</sup>

**Figure: 21. COMPARISION BASED ON GEST AGE WITH VARIOUS STUDIES**



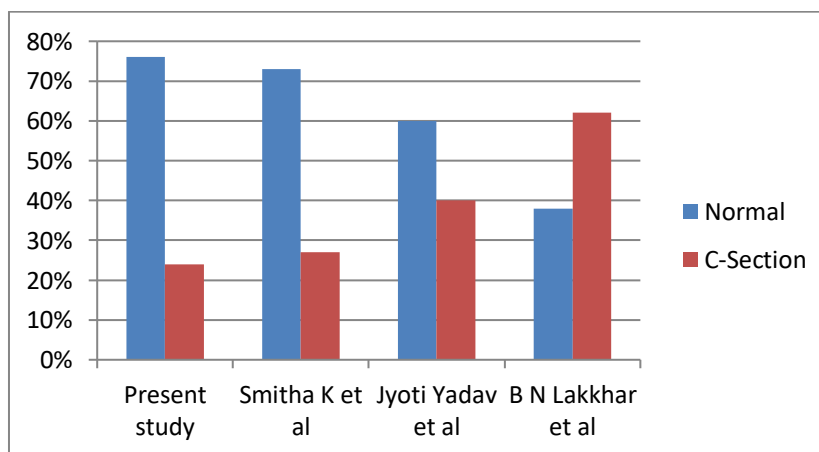
**COMPARISION BASED ON MODE OF DELIVERY:**

After routine investigations and follow up of the patient with serial ultrasound and Doppler, induction was planned based on the severity of the hypertension. Abnormal CP ratio was found in 72 women, and induction was planned in 48 cases of which 33 women delivered by vaginally with 15 women requiring Emergency LSCS for failed induction. 33 women required LSCS for indications causing foetal distress, meconium stained liquor, abnormal Doppler indices.

In the study by Smitha K et al., 73% delivered vaginally and 27% delivered by c-section.<sup>(19)</sup> In a study conducted by Jyoti Yadav et al., 60% delivered vaginally and 40% delivered by c section. It was found that 52.77% women in abnormal doppler group delivered by caesarean where as caeseran rate in normal Doppler group was 33%.<sup>(18)</sup>

In a study conducted by Bn Lakkhar et al., 37.9% delivered vaginally and 62.06% delivered by c-section.<sup>(21)</sup>

**Figure 22 COMPARISION BASED ON MODE OF DELIVERY WITH VARIOUS STUDIES**



## COMPARISON BASED ON BIRTH WEIGHT:

In the present study, out of 200 babies, 106 babies weighed more than 2.5kg (53%), 86 babies weighed between 1.5 – 2.4 and 8 babies weighed  $\leq$  1.5kg. The mean birth weight was 2.46kg with a standard deviation of 0.45kg. In a study conducted by S.Kanmani et al., mean birthweight was 1.9 kg with a standard deviation of 0.44.<sup>(22)</sup> In a study conducted by Smitha k et al., 36% babies are  $>$ 2.5kg and 64%  $<$ 2.5kg. UA-PI is significantly related to the low Birth weight with  $P < 0.0001$  in their study.<sup>(19)</sup>

## COMPARISON OF ADVERSE PERINATAL OUTCOME:

In our study incidence of perinatal death, NICU admission and low APGAR score at 5 min were the criteria for adverse perinatal outcome. Out of 200 cases, adverse perinatal outcome is seen in 75 cases which is 37.5% of the total cases. In the present study, low APGAR, NICU admissions and perinatal death were all seen cumulatively in some cases ( $n=3$ ). Pregnancy outcome was abnormal in 75 patients. More than one adverse outcome observed in some cases ( $n=26$ ). In the study by Smitha k et al, variables that are considered for adverse perinatal outcome was Low APGAR, NICU admission, Caesarean section, IUD, Perinatal death. Pregnancy outcome was abnormal in 46 patients. More than one adverse outcome observed in some cases ( $n=23$ ).<sup>(19)</sup> The present study demonstrates the cerebroplacental ratio and UA PI efficacy for determining the immediate perinatal outcome. Several studies conducted by Gramellini et al, Berkowitz et al and Fairlie et al have demonstrated the correlation between abnormal Doppler indices of fetal vessels and adverse perinatal outcome and fetal distress.<sup>(23-25)</sup>

## COMPARISON OF UA PI AND PERINATAL OUTCOME WITH OTHER STUDIES:

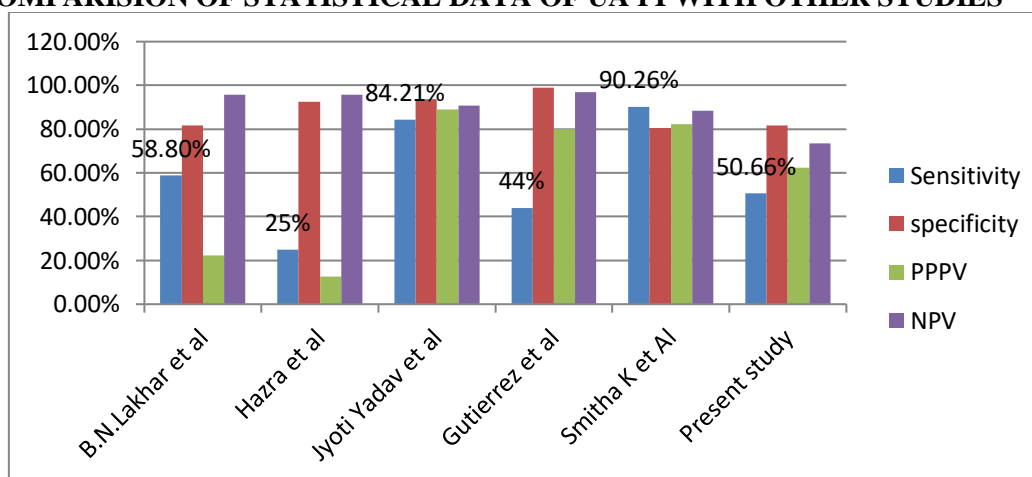
In the present study we studied umbilical artery PI and MCA/UA PI ratio, a Doppler index that reflects both umbilical-placental and cerebral vascular beds for identifying adverse perinatal outcome. Out of the two, Cerebroplacental ratio(MCA PI/UA PI) serves to be a better predictor compared to UA PI for determining adverse perinatal outcome. In the present study, UA PI value were considered based on gestational age specific reference values of which 69.5% individuals showed normal value and 30.5% showed values beyond 95<sup>th</sup> percentile which is considered abnormal for the corresponding gestational age. UA PI has better predictability at 32 weeks gestation in the present study. Mean UA PI in the present study is 1.01 with a standard

deviation of 0.21. Abnormal UA PI value has a sensitivity of 46.81% in predicting low birth wt  $<$ 2.5kg and specificity of  $>$ 83.96% with a diagnostic accuracy rate of 66.50% for a  $p$  value  $<$ 0.0001. Abnormal UA PI value has a sensitivity of 53.62% and specificity of 81.68% in determining APGAR  $<$ 7 with an accuracy rate of 72% for  $p$  value  $<$ 0.0001. In this study. the UA PI has a sensitivity of 50.66% and specificity of 81.6% in predicting adverse neonatal outcome. It has a PPV of 62.29% and NPV of 73.38%. Smitha et al., studied umbilical artery PI, middle cerebral artery PI and MCA/UA PI ratio. Out of 100 cases studied, 54 cases had abnormal UA PI and 46 cases had normal UA PI. 50 cases out of 54 had low birth weight which is 92.60% of the total abnormal UA PI Doppler. UA PI has a sensitivity of 90.26%, specificity of 80.57% and accuracy of 84% on predicting adverse pregnancy outcome.<sup>(19)</sup> Padmini C P et al studied on 80 cases of which abnormal UA PI is seen in 93% cases and showed adverse perinatal outcome which proved to be more sensitive in their study compared to other UA Doppler indices.<sup>(18)</sup> Jyoti Yadav et al. found that UA PI had highest predictive power with sensitivity (84.21%), specificity(93.54%), PPV (88.88%) and accuracy (90%) followed by UA S/D having sensitivity (80.64%), PPV (76.92%) and accuracy (86.5%). Specificity and negative predictive power was same for all the UA indices.<sup>(18)</sup> Yoon BH et al demonstrated that an abnormal umbilical artery Doppler waveform is a strong and independent predictor of adverse perinatal outcome in patients with preeclampsia.<sup>(26)</sup> Present study is comparable with that of Jyoti Yadav et al, B.N. Lakhar et al<sup>1</sup>, Hazra et al and Gutierrez et al where their studies also showed UA PI has more specificity than sensitivity.<sup>(18,21,27,28)</sup> But it is not comparable with the study of Smitha K et al as their study showed sensitivity of UA PI is more than specificity.<sup>(19)</sup>

**Table: 32. COMPARISON OF UA PI WITH VARIOUS STUDIES**

|   | Sensitivity | Specificity | Positive predictive value | Negative predictive value |
|---|-------------|-------------|---------------------------|---------------------------|
| <b>B.N. Lakhar et al<sup>(21)</sup></b> | 58.80%      | 81.80%      | 22.20%                    | 95.70%                    |
| <b>Hazra et al<sup>(27)</sup></b>       | 25%         | 92.60%      | 12.50%                    | 95.60%                    |
| <b>Jyoti yadav et al<sup>(18)</sup></b> | 84.21%      | 93.54%      | 88.88%                    | 90.62%                    |
| <b>Gutierrez et al<sup>(28)</sup></b>   | 44.00%      | 99.00%      | 80.00%                    | 97.00%                    |
| <b>Smitha K et al<sup>(19)</sup></b>    | 90.26%      | 80.57%      | 82.24%                    | 88.35%                    |
| <b>PRESENT STUDY</b>                    | 50.66%      | 81.60%      | 62.29%                    | 73.38%                    |

**Figure: 23. COMPARISON OF STATISTICAL DATA OF UA PI WITH OTHER STUDIES**



**COMPARISON OF CP RATIO AND PERINATAL OUTCOME WITH OTHER STUDIES:**

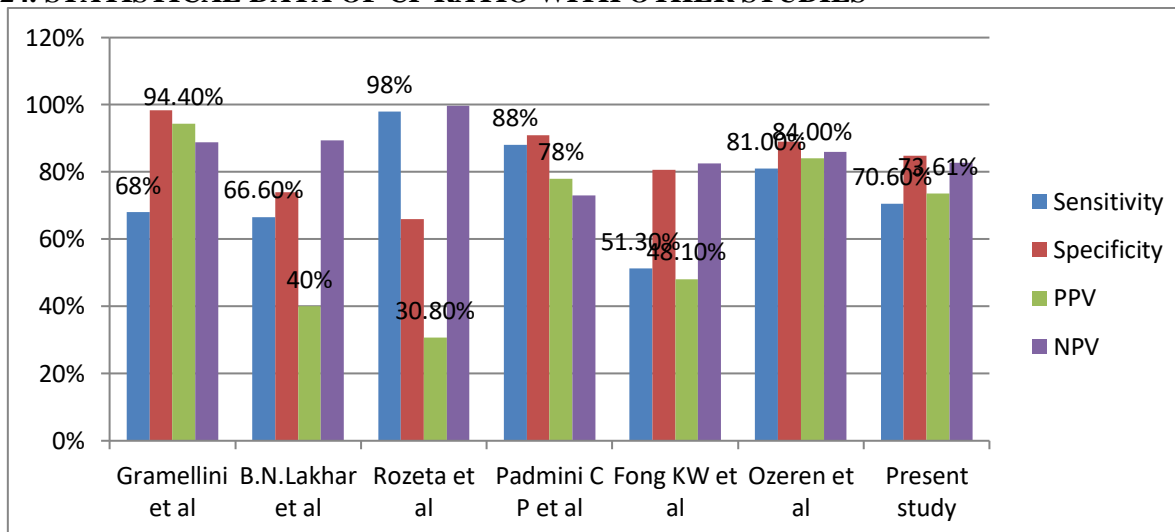
In the present study, cerebroplacental ratio has a better prediction for low birth weight, APGAR<7, NICU admissions and perinatal death compared to UA PI with a better sensitivity, specificity, positive and negative predictive values. But CP ratio and UA PI has no significant association with colour of liquor. Cerebroplacental ratio < 1.08 has a sensitivity of 61.7% and specificity of 86.79 % in predicting low birth weight <2.5kg which was statistically significant with a p value < 0.0001. Abnormal UA PI value has a sensitivity of 46.81% in predicting low birth wt <2.5kg and specificity of >83.96% with a diagnostic accuracy rate of 66.50% for a p value<0.0001. Cerebroplacental ratio < 1.08 has a sensitivity of 75.36% and specificity of 84.73% in predicting APGAR <7 which was statistically significant with a p value < 0.0001. Abnormal UA PI value has a sensitivity of 53.62% and specificity of 81.68% in determining APGAR<7 with an accuracy rate of 72% for p value <0.0001. Cerebroplacental ratio < 1.08 has a

very good specificity of 84.8%, sensitivity of 70.6%, PPV of 73.61%, NPV of 82.8% in predicting adverse perinatal outcome. It has a better diagnostic accuracy rate of 79.50% when compared with UA PI alone. Cerebroplacental ratio < 1.08 has a very low accuracy rate of predicting colour of liquor with an accuracy rate of 28%. Abnormal UA PI value has no significance in relation to colour of liquor as sensitivity and specificity is 21.52% and 35.71% respectively with very low positive and negative predictive values. Wladimiroff et al reported that CP ratio does not vary significantly between 30 and 40 weeks of gestation Hence it is possible to use a single cut off value for CP ratio after 30 weeks. (29) Arbeille et al also found CP ratio to be constant during last 10 weeks of gestation and suggested a cut off value of 1.08 above which values are considered normal and below 1.08 values are abnormal. (10) Mari G et al, have suggested that cerebral Doppler indices are associated with adverse perinatal outcome, while others like Gramellini et al and Ozeren M et al have proposed the cerebral-umbilical ratio as a better predictor of adverse perinatal outcome. (23,30,31)

**Table: 33. COMPARISON OF CP RATIO WITH VARIOUS STUDIES**

|   | Sensitivity | Specificity | Positive predictive value | Negative predictive value |
|---|-------------|-------------|---------------------------|---------------------------|
| <b>Gramellini et al<sup>(23)</sup></b>  | 68%         | 98.40%      | 94.40%                    | 88.80%                    |
| <b>B.N.Lakhar et al<sup>(21)</sup></b>  | 66.60%      | 73.90%      | 40%                       | 89.40%                    |
| <b>Rozeta et al<sup>(32)</sup></b>      | 98%         | 66%         | 30.80%                    | 99.70%                    |
| <b>Padmini C P et al<sup>(20)</sup></b> | 88%         | 91%         | 78%                       | 73%                       |
| <b>Fong KW et al<sup>(33)</sup></b>     | 51.30%      | 80.60%      | 48.10%                    | 82.50%                    |
| <b>Ozeren et al<sup>(30)</sup></b>      | 81.00%      | 89.00%      | 84.00%                    | 86.00%                    |
| <b>PRESENT STUDY</b>                    | 70.60%      | 84.80%      | 73.61%                    | 82.80%                    |

**Figure: 24. STATISTICAL DATA OF CP RATIO WITH OTHER STUDIES**



Present study is comparable with that of Ozeren et al and Gramellini et al who also concluded that CP ratio had higher sensitivity and positive predictive value compared to UA PI.<sup>(23,30)</sup> Present study is not correlating with Fong KW et al who showed a sensitivity and PPV of 51.3% and 48.1% respectively, the difference may be because they included small for

gestational age cases in their study.<sup>(33)</sup> The present study also doesn't correlate with that of Rozeta et al where their study showed higher sensitivity of CP ratio compared to specificity which is not as that of the present study. But the study correlates with respect to their high NPV(99.70%) which is similar to the present study where the NPV is 82.8%.

**COMPARISON OF DIAGNOSTIC ACCURACY:**

Our study shows highest diagnostic accuracy for CP ratio, followed by umbilical artery pulsatility index.

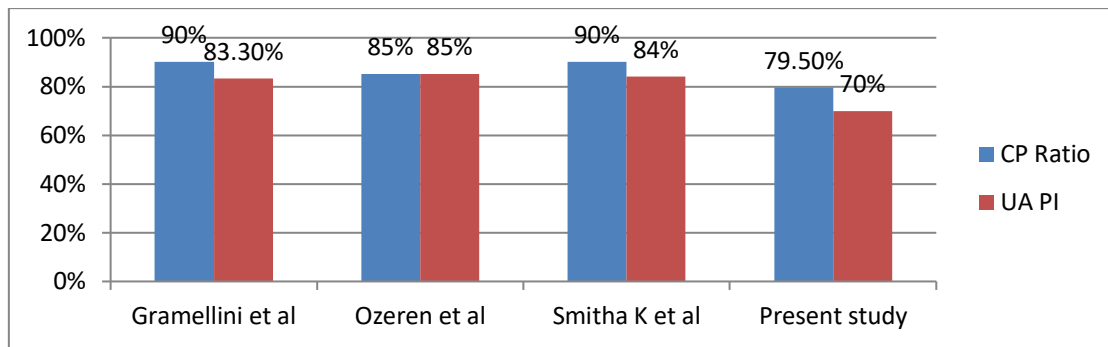
**Table: 34. COMPARISON OF DIAGNOSTIC ACCURACY WITH VARIOUS STUDIES**

|  | CP Ratio | UA PI  |
|--|----------|--------|
| <b>Gramellini et al<sup>(23)</sup></b> | 90.00%   | 83.30% |
| <b>Ozeren et al<sup>(30)</sup></b>     | 85.00%   | 85.00% |
| <b>Smitha K et al<sup>(19)</sup></b>   | 90.00%   | 84.00% |
| <b>PRESENT STUDY</b>                   | 79.50%   | 70%    |

The present study is comparable with Gramellini et al and Smitha et al in terms of diagnostic accuracy where cerebroplacental ratio has been proved to have been better predictor compared to UA PI alone.<sup>(19,23)</sup> Present

study is not correlating with Ozeren et al where the diagnostic accuracy is same for both cerebroplacental ratio and UA PI.<sup>(30)</sup>

**Figure: 25. ANALYSIS OF DIAGNOSTIC ACCURACY OF CP RATIO AND UA PI WITH VARIOUS STUDIES**



**CONCLUSION:**

The present study draws inference that hypertensive disorders of pregnancy are most common in Primigravida and the mean age group is 25.4yrs with a standard deviation of 3.48. The study states that cerebroplacental ratio is a better tool for assessing immediate perinatal outcome. It has good accuracy for determining low birth weight and APGAR <7. UA PI also has a better accuracy for determining immediate perinatal outcome but out of comparison, Cerebroplacental ratio seems to be a better predictor in the present study. Overall diagnostic accuracy for the present study is 79.5% for CP Ratio and 70% for UA PI. Doppler velocimetry which is a non invasive tool has proved to detect early foetal compromise and in hypertensive disorders of pregnancy. It also aids in planning a timely delivery and thereby reducing both perinatal morbidity as well as perinatal mortality. Compared to other methods of foetal monitoring, the use of Doppler has proved to be more sensitive in detecting foetal compromises early and helps in aiding appropriate and timely delivery.

**REFERENCES:**

1. Hypertension in pregnancy. Report of the American College of Obstetricians and Gynecologists’ Task Force on Hypertension in Pregnancy. *Obstet Gynecol.* 2013 Nov;122(5):1122–31.
2. 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;2015:614747.
3. Pijnenborg R, Bland JM, Robertson WB, Dixon G, Brosens I. The pattern of interstitial trophoblastic invasion of the myometrium in early human pregnancy. *Placenta.* 1981 Dec;2(4):303–16.

4. De Wolf F, De Wolf-Peeters C, Brosens I, Robertson WB. The human placental bed: electron microscopic study of trophoblastic invasion of spiral arteries. *Am J Obstet Gynecol.* 1980 May 1;137(1):58–70.
5. American College of Obstetricians and Gynecologists, American College of Obstetricians and Gynecologists, editors. *Hypertension in pregnancy.* Washington, DC: American College of Obstetricians and Gynecologists; 2013. 89 p.
6. Nicolaides K, Rizzo G, Hecher K, Ximenes R. *Doppler in Obstetrics.* :162.
7. Alfirevic Z, Stampalija T, Gyte GM. Fetal and umbilical Doppler ultrasound in high-risk pregnancies. *Cochrane Database Syst Rev.* 2010 Jan 20;(1):CD007529.
8. Simanaviciute D, Gudmundsson S. Fetal middle cerebral to uterine artery pulsatility index ratios in normal and pre-eclamptic pregnancies. *Ultrasound Obstet Gynecol.* 2006;28(6):794–801.
9. Alfirevic Z, Stampalija T, Gyte GM. Fetal and umbilical Doppler ultrasound in high-risk pregnancies. *Cochrane Database Syst Rev.* 2010 Jan 20;(1):CD007529.
10. Arbeille P, Body G, Saliba E, Tranquart F, Berson M, Roncin A, et al. Fetal cerebral circulation assessment by Doppler ultrasound in normal and pathological pregnancies. *Eur J Obstet Gynecol Reprod Biol.* 1988 Dec;29(4):261–73.
11. Bahado-Singh RO, Kovanci E, Jeffres A, Oz U, Deren O, Copel J, et al. The Doppler cerebroplacental ratio and perinatal outcome in intrauterine growth restriction. *Am J Obstet Gynecol.* 1999 Mar;180(3 Pt 1):750–6.
12. Sharma DK, Minhas DA, Sharma DrS. Cerebro-placental ratio as a predictor of neonatal outcome in hypertensive disorders of pregnancy.



Int J Clin Obstet Gynaecol. 2021 Jan 1;5(1):69–72.

13. Bonnevier A, Maršál K, Brodzski J, Thuring A, Källén K. Cerebroplacental ratio as predictor of adverse perinatal outcome in the third trimester. *Acta Obstet Gynecol Scand.* 2021 Mar;100(3):497–503.

14. Acharya G, Wilsgaard T, Berntsen GKR, Maltau JM, Kiserud T. Reference ranges for serial measurements of umbilical artery Doppler indices in the second half of pregnancy. *Am J Obstet Gynecol.* 2005 Mar;192(3):937–44.

15. Madazli R, Budak E, Calay Z, Aksu MF. Correlation between placental bed biopsy findings, vascular cell adhesion molecule and fibronectin levels in pre-eclampsia. *BJOG Int J Obstet Gynaecol.* 2000 Apr;107(4):514

16. Hershkovitz R, Kingdom JC, Geary M, Rodeck CH. Fetal cerebral blood flow redistribution in late gestation: identification of compromise in small fetuses with normal umbilical artery Doppler. *Ultrasound Obstet Gynecol Off J Int Soc Ultrasound Obstet Gynecol.* 2000 Mar;15(3):209–12.

17. Ebbing C, Rasmussen S, Kiserud T. Middle cerebral artery blood flow velocities and pulsatility index and the cerebroplacental pulsatility ratio: longitudinal reference ranges and terms for serial measurements. *Ultrasound Obstet Gynecol Off J Int Soc Ultrasound Obstet Gynecol.* 2007 Sep;30(3):287–96.

18. Devender Kumar JY. Role of umbilical artery Doppler velocimetry in predicting the adverse perinatal outcomes in hypertensive disorder of pregnancy. *Int J Reprod Contracept Obstet Gynecol.* 2017 Oct;6(10).

19. K S, K S, T M. Study of Doppler waveforms in pregnancy induced hypertension and its correlation with perinatal outcome. *Int J Reprod Contracept Obstet Gynecol.* 2017 Jan 2;3(2):428–33.

20. P PC, Das P, R C, Adithya MS. Role of Doppler indices of umbilical and middle cerebral artery in prediction of perinatal outcome in preeclampsia. *Int J Reprod Contracept Obstet Gynecol.* 2017 Feb 23;5(3):845–9.

21. Lakhkar BN, Rajagopal KV, Gourisankar PT. Doppler prediction of adverse perinatal outcome in PIH and IUGR. *Indian J Radiol Imaging.* 2006 Feb;16(01):109–16.

22. Kanmani, S. Evaluation of Doppler Cerebroplacental Ratio as a Predictor of Adverse

Perinatal Outcome in High Risk Pregnancy [Chennai.]: Madras Medical College; 2017.

23. Gramellini D, Folli MC, Raboni S, Vadora E, Merialdi A. Cerebral-umbilical Doppler ratio as a predictor of adverse perinatal outcome. *Obstet Gynecol.* 1992 Mar;79(3):416–20.

24. Berkowitz GS, Mehalek KE, Chitkara U, Rosenberg J, Cogswell C, Berkowitz RL. Doppler umbilical velocimetry in the prediction of adverse outcome in pregnancies at risk for intrauterine growth retardation. *Obstet Gynecol.* 1988 May;71(5):742–6.

25. Fairlie FM, Moretti M, Walker JJ, Sibai BM. Determinants of perinatal outcome in pregnancy-induced hypertension with absence of umbilical artery end-diastolic frequencies. *Am J Obstet Gynecol.* 1991 Apr;164(4):1084–9.

26. Yoon BH, Lee CM, Kim SW. An abnormal umbilical artery waveform: a strong and independent predictor of adverse perinatal outcome in patients with preeclampsia. *Am J Obstet Gynecol.* 1994 Sep;171(3):713–21.

27. Hazra SK, Dash KK, Chaudhuri A, Ghosh MK, Banerjee D, Guha S. A Prospective Study of Doppler Velocimetry in Pregnancy-induced Hypertension in a Rural Population of a Developing Country. *J Basic Clin Reprod Sci.* 2013;2(2):127–31.

28. Romero Gutiérrez G, Aguilar Barajas I, Chávez Curiel A, Ponce Ponce de León AL. [Prediction of fetal wellbeing with Doppler flowmetric profile in pregnant hypertensive women]. *Ginecol Obstet Mex.* 2001 Dec;69:480–6.

29. Wladimiroff JW, vd Wijngaard JA, Degani S, Noordam MJ, van Eyck J, Tonge HM. Cerebral and umbilical arterial blood flow velocity waveforms in normal and growth-retarded pregnancies. *Obstet Gynecol.* 1987 May;69(5):705–9.

30. Ozeren M, Dinç H, Ekmen U, Senekayli C, Aydemir V. Umbilical and middle cerebral artery Doppler indices in patients with preeclampsia. *Eur J Obstet Gynecol Reprod Biol.* 1999 Jan;82(1):11–6.

31. Mari G, Moise KJ, Deter RL, Kirshon B, Carpenter RJ, Huhta JC. Doppler assessment of the pulsatility index in the cerebral circulation of the human fetus. *Am J Obstet Gynecol.* 1989 Mar;160(3):698–703.

32. Shahinaj R, Manoku N, Kroi E, Tasha I. The value of the middle cerebral to umbilical artery Doppler ratio in the prediction of neonatal

outcome in patient with preeclampsia and gestational hypertension. *J Prenat Med.* 2010;4(2):17–21.

33. Fong KW, Ohlsson A, Hannah ME, Grisaru S, Kingdom J, Cohen H, et al. Prediction of perinatal

outcome in fetuses suspected to have intrauterine growth restriction: Doppler US study of fetal cerebral, renal, and umbilical arteries. *Radiology.* 1999 Dec;213(3):681–9.