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Original Research Article

Early onset and late onset preeclampsia-maternal and perinatal outcomes in a rural tertiary health center

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ABSTRACT

Background: Preeclampsia is main cause of morbidity and mortality both mother and fetus. Preeclampsia occurs in 10-17% of pregnancies. Preeclampsia was divided into early onset preeclampsia is occur at less <34 weeks of gestation age and late onset preeclampsia is occur at >34 weeks of gestation age. Early and late onset preeclampsia have different etiology and should be considered as different disease as there are difference in clinical manifestation, maternal and perinatal outcome, prognosis and complication.

Methods: An analytic observational study involving retrospective data done at RL Jalappa Hospital, Sri Devaraj Urs Medical College, Kolar. 217 women with singleton pregnancies with Pre eclampsia who were admitted and delivered in our hospital between June 2016 and May 2017 were recruited for this retrospective study.

Results: The results showed that the incidence of EOPE (27.6%) was lower than LOPE (72.4%). Diastolic blood pressure is significantly higher in EOPE compared to LOPE. Complications in perinatal outcomes such as low birth weight (<2500 gram) are more in EOPE (98.3%) compared to LOPE (45.2%) and asphyxia is more on EOPE (11.7%) compared to LOPE (1.3%). Stillbirth in EOPE (15%) is more than LOPE group (3.2%).

Conclusions: It is observed that EOPE incidence rate is lower than LOPE. Maternal and perinatal complications are greater in the EOPE group.

Keywords: Early and late onset preeclampsia, Fetal outcome, Maternal morbidity

INTRODUCTION

Pre eclampsia is a multisystem disorder occurring in pregnancy and puerperium which is characterised by development of hypertension of 140/90 mmHg and above after 20th week in a previously normotensive patient.¹ It is a global problem and complicates approximately 10-17% of pregnancies. The incidence of preeclampsia is 2 to 10% of all pregnancies in the world. According to WHO the incidence is 7 times greater in developing countries compared to developed countries.² Currently there has been a change in the definition and understanding of Preeclampsia, known as Early Onset Preeclampsia (EOPE) and Late Onset Preeclampsia (LOPE).early onset where preeclampsia occurs at <34 weeks gestational age

and late onset occurring at >34 weeks of gestation. Even though the presenting features overlap, there are differences in maternal and perinatal outcome, prognosis and complications. Early and late onset preeclampsia have different aetiologies and should be considered as different disease.³ Early onset preeclampsia is the most severe clinical variant of disease occurring 5-20% of all cases of preeclampsia and is associated with neonatal morbidity and mortality. Late onset preeclampsia occurring about 75-80% of all cases of preeclampsia; which are associated with maternal morbidity (metabolic syndrome, impaired glucose tolerance, obesity, dyslipidaemia, chronic hypertension), normal birth weight and normal placental volume.⁴

METHODS

217 women with singleton pregnancies with Pre eclampsia who were admitted and delivered in our hospital between June 2016 and May 2017 were recruited for this retrospective study.

Included patients (n=217) were divided into two groups: Early onset preeclampsia (EOPE) group consisting of 60 patients diagnosed before 34 weeks of gestation, Late onset preeclampsia (LOPE) group consisting of 157 patients who were diagnosed at or after 34 weeks gestation.

Exclusion criteria include pregnant women with essential hypertension or hypertension <20 weeks gestation, pre existing renal disease, multiple pregnancies, liver disorder, and epilepsy. The two groups were matched according to age, education, gestational age, parity, clinical findings (systolic blood pressure, diastolic blood pressure). Maternal outcomes include duration of treatment, mode of delivery and foetal outcomes include birth Weight, apgar score at 5th minute, intra uterine death (IUD), Still births were compared.

Statistical analysis

Data are expressed as n (%) and mean with standard deviation.

The data were tested for normal distribution and Student's t test, One-way ANOVA or the Mann-Whitney U test was used for comparison of two groups where appropriate. Statistical Package for the Social Sciences (SPSS) 17.0. A p value <0.05 was regarded as significant.

RESULTS

A total of 217 patients at >20 weeks of gestation with pre eclampsia were taken in this study divided into EOPE and LOPE groups (n=60 and 157) respectively.

Characteristics of preeclampsia were illustrated in Table 1; 148 (68.2%) patients were in the age group 20 to 35 yrs. Education of patients were upto high school 128 (59%). Majority of patients with pre eclampsia are multigravidas.

Based on the gestational age of PE, LOPE patients were 157 (72.4%) more than EOPE 60 (27.4%).

While the systolic blood pressure was on average 170.33±14.43 and mean diastolic 107.06± 9.557.

Table 2 illustrates EOPE were found more in multigravida while LOPE were found more in primigravida which is statistically significant.

Table 1: Characteristics of patients with preeclampsia.

Variable	N= 217
Age (years)	
<20	17 (7.8%)
20-35	148 (68.2%)
>35	52 (24%)
Education	
Elementary	57 (26.3%)
Middle school	60 (27.6%)
High school	74 (34.1%)
college	26 (12%)
Parity	
Primigravida	89 (41%)
Multigravida	128 (59%)
Gestational age (weeks)	
<34 (EOPE)	60 (27.6%)
≥34 (LOPE)	157 (72.4%)
Blood pressure (mmHg)	
Systolic	
Mean	170.36±14.93
Median	170
Range	120-230
Diastolic	
Mean	107.08±9.804
Median	110
Range	90-160

Table 2: Characteristics of patients EOPE and LOPE.

Variable	EOPE (n=60)	LOPE (n=157)	P value
Age (years)			
<20	5(8.3%)	24(15.3%)	0.0473
20-35	42(70%)	95(60.5%)	
>35	13(21.7%)	38(24.2%)	
Education			
Elementary	14 (23.4%)	38 (24.2%)	1.000
Middle school	16 (26.7%)	47 (30%)	
High school	25 (41.6%)	60 (38.2%)	
College	5 (8.3%)	12 (7.6%)	
Parity			
Primigravida	32 (53.3%)	92 (58.6%)	0.017*
multigravida	28 (46.7%)	65 (41.4%)	
Blood pressure :Systolic			
Mean	173.16±17.9	169.26±12.75	0.073
Median	170	170	
Range	140-250	120-230	
Diastolic			
Mean	109.71±10.76	106.05±8.71	0.002*
Median	110	110	
Range	100-150	60-160	

Table 3: Maternal and perinatal outcomes of patients EOPE and LOPE.

Variable	EOPE (n=60)	LOPE (n=157)	P value
Mode of delivery			
Caesarean section	32 (53.3%)	92 (58.6%)	0.366
Vaginal delivery	28 (46.7%)	65 (41.4%)	
Vaginal delivery			
Spontaneous	25 (89.3%)	15 (23.1%)	0.0001*
Forceps	2 (7.1%)	47 (72.3%)	
Vacuum	0 (0)	2 (3.1%)	
Episiotomy	1 (3.6%)	1 (1.3%)	
Length of hospital stay(days)			
Mean	5.59	5.32	0.0001*
Median	4	4	
Range (min-max)	1-92	1-35	
Birth weight			
< 2000gm	55 (91.6%)	24 (15.2%)	0.0001*
2000-2499gm	4 (6.7%)	47 (30.0%)	
>2500gm	1 (1.7%)	86 (54.8%)	
Apgar score (5min)			
Asphyxia	7 (11.7%)	2 (1.3%)	0.0001*
Not asphyxia	44 (73.3%)	150 (95.5%)	
Stillborn	9 (15%)	5 (3.2%)	

Blood pressure increases in both EOPE and LOPE but increase of diastolic pressure is more on EOPE than LOPE which is statistically significant.

Table 3 illustrates the comparison of outcomes in patients with EOPE and LOPE. Most deliveries are performed by caesarean section; 53% in EOPE and 58.6% in LOPE but not statistically significant. Spontaneous deliveries are more in EOPE group than in LOPE which is statistically significant. Duration of hospital stay is more in EOPE group than in LOPE which is significant. EOPE has more babies born weighing <2500 g (91.6%) where as in LOPE it weighs \geq 2500 grams (54.8%). Babies who are born in EOPE groups had asphyxia (11.7%) more than babies in LOPE group (1.3%).

DISCUSSION

Approaching as an early onset and late onset pre-eclampsia gives us better idea about understanding of the complex etiopathogenesis of this medical enigma.

The distinction between early onset and late onset PE is a relatively modern concept and is becoming widely accepted as a better indicator of disease significance⁵.

In our study the incidence of pre-eclampsia more in late onset (LOPE) than early onset preeclampsia (EOPE) group (72.4% vs 27.6%). Clinical characteristics like systolic blood pressure (SBP), diastolic blood pressure (DBP) were significantly higher in early onset (EOPE) group suggesting increased maternal total vascular

resistance which strengthens abnormal placentation as the probable cause for early onset subtype.

In our study gestational age at birth had a major impact on perinatal morbidity and mortality. Early onset group had early termination of pregnancy due to uncontrolled blood pressure, derangement of laboratory parameters and clinical manifestations. Perinatal mortality and still birth rates were higher in early onset preeclampsia (EOPE) than in late onset preeclampsia (LOPE).

Early onset preeclampsia (EOPE) is associated with increased risk of multi organ involvement including hepatic, hematologic, arterial, renal and adverse maternal and perinatal outcomes as compared with late onset preeclampsia (LOPE).

There are limited number of studies and reviews that have compared characteristics of early onset and late onset pre-eclampsia.

In a study conducted by Aziz, Mose, Indonesia shows maternal and perinatal morbidity and mortality is higher in EOPE group.⁶

Valensis et al. reported that patients who were diagnosed with EO-PE had higher total vascular resistance, while late-onset patients had low vascular resistance.⁷

In a study conducted by sreedevi atluri and nandish in mysore shows EOPE is more severe clinical entity than LOPE with high risk for life threatening maternal complications and fetal mortality.⁸

The distinction between EO- and LO-PE is a relatively modern concept and is becoming widely accepted as a better indicator of disease significance than the classic “mild” vs “severe” terminology.⁹ It was widely demonstrated that gestational age at birth had a major impact on perinatal outcome.¹⁰ Perinatal mortality and morbidity were demonstrated to be higher in women with EO-PE.¹¹⁻¹³ The results of the present study supported this observation.

CONCLUSION

Preeclampsia is a major complication in pregnancy with significant short term and long term consequences for both mother and foetus. currently, no therapies are available to slow or reverse the disease. A recent review by Sibai supported expectant management in selected cases of severe PE presenting before 34 weeks, good perinatal outcomes were reported. The provision of MgSO₄ 40% as a brain protector and corticosteroids to infant lung maturation can reduce complications of perinatal outcomes. CLASP (collaborative low-dose aspirin study in pregnancy) trial showed usage of low dose aspirin prophylactically in patients liable to early onset preeclampsia has better perinatal outcome.

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Ethical approval: The study was approved by the Institutional Ethics Committee

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