

MATERNAL AND PERINATAL COMPLICATIONS IN PRE-ECLAMPSIA

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ABSTRACT

INTRODUCTION: Hypertensive disorders of pregnancy is one of the major causes of maternal morbidity/mortality leading to 10-15% of maternal deaths in low - and middle-income countries. Pre-eclampsia produces potentially lethal complications including abruptio placenta, disseminated intravascular coagulation, intracranial hemorrhage, hepatic failure, acute renal failure, and cardiovascular collapse. Intrauterine fetal growth restriction, intrauterine fetal demise and prematurity are the other related obstetric problems. The current study aimed to find the maternal and perinatal complications of pre-eclampsia in our context.

MATERIAL AND METHODS: A prospective study of 145 patients with the diagnosis of pre-eclampsia treated in a tertiary level teaching hospital between June 2011 to December 2013 was done. Hypertension in pregnancy, pre-eclampsia, severe pre-eclampsia and eclampsia were defined in the standard way. Maternal and perinatal complications were identified and analyzed.

RESULTS: Pre-eclampsia was detected in 6.02% of all deliveries. Maternal complications were abruptio placenta (11.03%), eclampsia (8.27 %), HELLP syndrome (4.12%), hypertensive encephalopathy (0.7%), acute renal failure (0.7%) and maternal death (0.7%). Fetal complications included preterm delivery (17.93%), intra uterine growth restriction (10.34 %), intra uterine death (8.30%) and neonatal death(2.80%).

CONCLUSION: Pre-eclampsia is associated with increased maternal-perinatal complications. The adverse outcome can be minimized by more widespread use of prenatal care, education of primary medical care personnel, prompt diagnosis of high-risk patients and timely referral to tertiary medical centers.

KEYWORDS: Pre-eclampsia; Eclampsia; Complications

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INTRODUCTION

Hypertensive disorder of pregnancy is one of the major causes of maternal morbidity and mortality. Overall, 10% - 15% of direct maternal deaths are associated with pre-eclampsia and eclampsia in low- and middle-income countries.^{1,2} It occurs in 12 - 22% of pregnancies depending on the population and the definitions used.³ Approximately 70 % of hypertensive disorders of pregnancy are gestational hypertension/ pre-eclampsia and 30 % is caused by chronic hypertension.⁴ Pre-eclampsia produces potentially lethal complications including abruptio placenta, eclampsia, disseminated intravascular coagulation, intracranial hemorrhage, HELLP syndrome, acute renal failure, and cardiovascular collapse. Most deaths in pre-eclampsia occur due to its complications and not due to hypertension per se. Intrauterine fetal growth restriction (IUGR), intrauterine fetal demise and prematurity are the most common fetal complications. All these clinical situations require prompt diagnosis and aggressive management in order to reverse adverse maternal-perinatal outcome. This study was carried out to find the extent of maternal and fetal complications of preeclampsia so that it will serve as a base line data for further studies on this subject matter.

MATERIAL AND METHODS

A prospective study of 145 patients with the diagnosis of pre-eclampsia treated between June 2011 to December 2013 in the department of obstetrics and gynecology of College of Medical Sciences Teaching Hospital was done. Patients were enrolled from both the antenatal clinics and antenatal ward. All the patients with pre-eclampsia who delivered in our centre were included. The patients with gestational hypertension and those with chronic hypertension or pre-eclampsia superimposed on chronic hypertension were excluded from the study. The diagnosis of pre-eclampsia and eclampsia was based on the criteria defined by national high blood pressure education programme working group on high blood pressure in pregnancy.⁵ Severe pre-eclampsia was diagnosed with one or more of the following criteria: blood pressure $\geq 160/110$, proteinuria of at least 5 g/24 h, oliguria (<600 ml/24 h or <30 -50 ml/h), intrauterine fetal growth restriction, oligohydramnios (amniotic fluid index < 50 mm), symptoms suggesting end organ failure such as headache, visual disturbances, epigastric pain, medical complications involving pulmonary edema, cerebral edema, acute renal insufficiency, HELLP syndrome (hemolysis, elevated liver enzymes, low platelet count). Blood pressure was controlled with alpha-Methyldopa or Nifedipine or both. Nifedipine was administered orally to control high blood pressure values $\geq 160/110$ mmHg. Magnesium Sulphate was used for the

control of seizure in eclampsia or for prevention of seizure in severe pre-eclampsia. The Pritchard regimen was followed (4 g intravenous along with 10 mg i.m loading dose followed by 5 mg intramuscular every 4 hourly until 24 h after delivery). Dexamethasone, four doses of 6 mg, was administered intramuscularly in 48 hrs to accelerate the fetal lung maturity in cases with gestational age of 28 - 34 weeks. Prostaglandin analogue or oxytocin infusion was preferred to induce or augment labour respectively according to Bishop Score in cases which were decided to be delivered.

Statistical analysis was done by using Statistical Package for the Social Sciences Software (SPSS) Programme for windows[®] version 18. Statistical significance was accepted at the 5 percent level (P value < 0.05). Chi-square test was used wherever applicable.

RESULTS

In this study out of 2305 deliveries conducted between June 2011 to December 2013, 145 (6.02%) patients were found to be complicated with pre-eclampsia. The mean age of the patient was 24.42 years. (Table 1)

Table 1: Demographic and clinical parameters

Variable(s)	Values(percentage)
Total number of patients	145
Age in years*	24.42 \pm 4.56 (16-37)
Gravida	
Primigravida	88 (60.7)
Multigravida	57 (39.3)
Parity	
Nullipara	95 (65.5)
Multipara	50 (34.5)
Period of gestation (weeks)*	37.47 \pm 3.0 (22.6-42.4)
Presenting symptoms and signs	
Ankle edema	82 (56.6)
Headache	55 (37.9)
Blurring of vision	22 (15.2)
Epigastric pain	14 (9.7)
Vaginal bleeding	9 (6.2)
Convulsion	5 (3.4)
Preeclampsia severity	
Mild	63 (43.4)
Severe	82 (56.6)

* Mean \pm SD (range)

Eighty-three percentages of the patients were in the age range

of 20- 34 years. There was similarity between patients of mild and severe pre-eclampsia in terms of age, gravida and parity. Majority of the mothers (70%) who were affected by pre-eclampsia were from rural area. Fifty eight percent of the patient were with term pregnancy, 38.6% were preterm and 2.6% were postterm at the time of presentation.

Twenty nine of the patients with severe pre-eclampsia received Inj. MgSO₄, while 12 of them had seizure (with proteinuria) and none of the patients in severe pre-eclamptic group who received Inj.MgSO₄ prophylactically developed seizure. Cesarean section was the mode of delivery in 50.3% of the patients, while 45.5% of them had normal vaginal delivery and 4.1% had undergone instrumental delivery. Among 145 patients with pre-eclampsia, 119 (82%) of the patients required anti-hypertensive medicines either during antepartum, intrapartum or postpartum period. Among them, 76 were in severe pre-eclampsia and 43 in mild pre-eclampsia group.

The most frequently encountered maternal complications were abruptio placenta, eclampsia and HELLP syndrome. (Table 2)

Table 2: Maternal complications of pre-eclampsia

Maternal complications	Mild pre-eclampsia	Severe pre-eclampsia	Total
Abruptio placenta	2	14	16
Eclampsia	3	9	12
Oligohydramnios	1	6	7
HELLP syndrome	0	6	6
PPH	1	0	1
Acute renal failure	0	1	1
Hypertensive encephalopathy	0	1	1
Hypertensive retinopathy	0	1	1
Maternal mortality	0	1	1
Total	7	39	46

PPH: post partum hemorrhage, HELLP: hemolysis, elevated liver enzymes, low platelet count

Other complications were oligohydramnios, acute renal failure, hypertensive encephalopathy, hypertensive retinopathy. Abruptio placenta was diagnosed in severely preeclamptic and mildly preeclamptic groups with incidence of 17.07 % and 3.17% cases respectively, with a statistically

significant difference (P= 0.002). Maternal mortality occurred in 1(0.7%) case due to pulmonary edema.

Among the fetal complications prematurity was the most common ((17.93%) followed by intra uterine growth restriction (10.34 %), intra uterine death (8.30%) and neonatal death (2.80%). (Table3)

Table 3: Fetal complications of pre-eclampsia

Fetal outcome	Mild pre-eclampsia	Severe pre-eclampsia	Total
Normal	49	39	88
Preterm	8	18	26
IUGR	2	13	15
IUD	2	10	12
NND	2	2	4
Total	63	82	145

IUGR: intra uterine growth restriction, IUD: intra uterine death, NND: neonatal death

Adverse fetal outcome and complications were significantly associated with the severity of pre-eclampsia (p= 0.03).

DISCUSSION

Pre-eclampsia is best described as a pregnancy-specific syndrome that can affect virtually every organ system. The exact incidence of pre-eclampsia is unknown but it has been reported to be approximately 5-8 %.⁶ It was identified in 6.02% of the total deliveries in our study. The mean age of the patients was 24.42 years, 60.7% were primigravida and 69.7% of the patients were unbooked. Singhal et al. reported that mean age of the patients was 24.04 years and 73% were primigravida, 82% of the patients were unbooked.⁷ Headache, blurring of vision and epigastric pain were the antecedent symptoms in 37.5%, 15.2% and 9.2% of the patients, respectively. Singhal et al. reported headache in 44%, epigastric pain in 20% and blurring of vision in 8%.⁷

Among 145 patients of pre-eclampsia, 56.6% had severe pre-eclampsia which was similar to the result of Yucesoy et al. (54.11%); however, there was differences in the incidence of mild pre-eclampsia (43.4 % vs. 34.50 %).⁸ Cesarean section was the mode of delivery in 50.3% of the

patients while 45.5% had normal vaginal delivery and 4.1% had undergone instrumental delivery. In the study done by Vidhyadhar et al. rate of caesarean delivery, vaginal delivery and instrumental delivery were 35%, 59% and 6% respectively.⁹ Vaginal delivery is recommended for the severe preeclamptic women in the absence of obstetric indication for caesarean section. Elective abdominal delivery may be preferred in cases before 32 weeks with IUGR and oligohydramnios.¹⁰ Among 145 patients with hypertensive disorder of pregnancy, 119 (82%) of the patients required anti-hypertensive medicines either during antepartum, intrapartum or postpartum period. Among them, 76 were in severe pre-eclampsia and 43 in mild pre-eclampsia group.

Overall maternal complications occurred in 31.7 % which comprised of abruptio placenta (11 %), eclampsia (8.3 %), oligohydramnios (4.8 %), HELLP syndrome (4.1 %) acute renal failure (0.7 %), post-partum hemorrhage (0.7 %), hypertensive encephalopathy (0.7 %) and hypertensive retinopathy (0.7 %). The incidence of abruptio placenta in severely preeclamptic and mildly preeclamptic group was 17.07 % and 3.17% respectively, the difference was statistically significant ($P= 0.002$). Our findings were similar to that reported by Vidhyadhar et al. in which the incidence of abruptio placenta in severe and mild preeclampsia were 19.04% and 5.12% respectively.⁹ Contrary to these only 7.7% of the severely pre-eclamptic patients had abruptio placenta.¹¹

In our study 8.3% of the pre-eclampsia was complicated by eclampsia which is less as compared to that reported by Vidhyadhar et al. (19% eclampsia).⁹ The incidence of eclampsia in severe pre-eclamptic patients was higher (11% vs. 6.3 %) compared to that of Nankali et al.¹¹ Oligohydramnios was present in 21.68% of the pre eclampsia.¹¹ This is in contrast to our result of 4.8% of oligohydramnios. None of the patients with mild pre-eclampsia had HELLP syndrome, while 7.3 % in the severe pre-eclampsia group developed the syndrome. There is wide variations (0.3-11 %) in the incidence of this syndrome in the literature.^{8,11} The incidence of PPH (2%) and renal failure (2%) were higher than our study.⁹

Maternal mortality occurred in 1 case (0.7%) and she was diagnosed as pulmonary edema. She was an unbooked patient who had few antenatal visits at health post and was referred for high blood pressure with dyspnea. Maternal mortality was nil in Tuffnell et al.¹² series, while Singhal et al.⁷ reported 8 % maternal mortality associated with pre-eclampsia. Most of the deaths (89.5%) were in unbooked women and the most common causes of death were acute renal failure, cardiopulmonary failure, disseminated intravascular

coagulopathy and cerebrovascular accident.¹³

Perinatal complications occurred in 39.3 % of the preeclampsia patients. Prematurity was the most common complication seen in 17.93 % of the deliveries. It was found in 11.1% of mild preeclampsia and 21.9% in severe preeclampsia, which was statistically significant ($p=0.03$). The rate of prematurity in mild preeclampsia and severe preeclampsia were 17.99% and 47.62% respectively.⁹ Singhal et al. reported prematurity in 67.33 % of preeclampsia patients.⁷ IUGR was the next common complication seen in 10.34 % of pre eclampsia. In contrary to our findings Singhal et al. reported the incidence of IUGR to be 23.47% in preeclampsia.⁷ In mild pre-eclampsia the IUGR was 10.3% and in severe pre-eclampsia it was 15.8%. The rate of IUGR in mild and severe pre-eclampsia were 7.69% and 26.19% respectively.⁹ One of the fetal complications was IUD which occurred in 8.3% of the pre-eclampsia cases. This figure is markedly low (8.3 % vs. 28.57%) as reported by Singhal et al.⁷ Perinatal mortality was 11% in our study. Perinatal mortality in literature ranged between 4.7% to 41.6%.^{7,9,12,14} This study has some limitations. First, it is a single centre based study so the results may not be applicable to other populations. Second, the sample size is relatively small. So we recommend for further multicenter study with larger sample size in order to know the extent of the problems of pre-eclampsia and measures to tackle them in our context.

CONCLUSION

Pre-eclampsia is associated with increased risk of adverse maternal and perinatal outcome. The complications of severe pre-eclampsia and eclampsia could be prevented by more widespread use of prenatal care, education of primary medical care personnel, prompt diagnosis of high-risk patients and timely referral to tertiary medical centers.

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