



CODEN [USA]: IAJPBB

ISSN: 2349-7750

INDO AMERICAN JOURNAL OF
PHARMACEUTICAL SCIENCES<http://doi.org/10.5281/zenodo.3928059>Available online at: <http://www.iajps.com>

Research Article

MATERNAL AND PERINATAL OUTCOME IN PRETERM
PRELABOR RUPTURE OF MEMBRANESDr Mahnoor Shahid¹, Dr Maliha Younis¹, Dr Talia Saeed¹¹Fatima Jinnah Medical University, Lahore

Article Received: May 2020

Accepted: June 2020

Published: July 2020

Abstract:

Introduction: Preterm premature rupture of membranes (PPROM) occurs in 3% of pregnancies and is responsible for approximately one third of all preterm births. **Objectives of the study:** The main objective of the study is to analyze the maternal and perinatal outcome in preterm prelabor rupture of membranes. **Material and methods:** This cross sectional study was conducted in Ganga Ram Hospital, Lahore during June 2019 to January 2020. Patients admitted in Obstetrics and Gynaecology Unit through outpatient & casualty department with gestational age between 24 to 37 weeks with preterm premature rupture of membrane (PPROM) confirmed by ultra sound and clinical examination were included regardless of their age. **Results:** A total of 72 female patients were included in the study. Mean age and gestational age of the patients was 25.36 ± 3.8 years and 32.09 ± 1.78 weeks. 56.9% (n=41) patients had 15-25 years of age and 43.1% (n=31) patients fell in age group 26-40 years. Gestational age distribution showed that majority of the patients i.e. 58.3 (n=42) had gestational age 29-32 weeks and 41.7% (n=30) had 33-36 weeks of gestational age. **Conclusion:** It is concluded that most common maternal morbidity associated with PPRM was chorioamnionitis, that of neonatal morbidity was prematurity and its complications.

Corresponding author:**Dr. Mahnoor Shahid,**

Fatima Jinnah Medical University, Lahore

QR code



Please cite this article in press Mahnoor Shahid et al, *Maternal And Perinatal Outcome In Preterm Prelabor Rupture Of Membranes.*, Indo Am. J. P. Sci, 2020; 07(07).

INTRODUCTION:

Preterm premature rupture of membranes (PPROM) occurs in 3% of pregnancies and is responsible for approximately one third of all preterm births. The incidence of preterm premature rupture of membrane averages from 0.7 to 2.1% and accounts for about 20 to 40% cases of PROM before 37 weeks of gestation. For surviving preterm neonates, there may be significant health consequences with lasting disabilities, including respiratory problems, hearing and vision impairment, cerebral palsy, and mental retardation [1].

Delivery of fetus before thirty seven weeks (259 days) is labeled as preterm birth. Preterm birth is multifactorial and its incidence has increased one and half fold in last few years and is a leading cause of fetal mortality worldwide. About 75% mortality and morbidity of fetus and mother occurs during preterm birth [2]. Pattern and mechanism of preterm labor is unknown but in all cases force full uterine contractions are common. Uterine contractions may be physiological and pathological that ends with preterm delivery of fetus [3].

Preterm labor is an unresolved issue in obstetrical profession and has lot of challenges for fetal and maternal outcomes. Preterm labor can be stopped with use of tocolytic medication, antibiotics and corticosteroids; in cases of preterm labor progesterone can be used for maintenance of tocolytic effect [4]. Many techniques have been used and recommended to stop the preterm labor. Use of excessive water to prevent dehydration and complete bed rest are famous but do not have any effect in inhibition of preterm labor. It has been shown that parenteral to colytic agents have better outcome as compared to oral [5].

The fetal and neonatal morbidity and mortality risks are significantly affected by severity of oligohydramnios, duration of latency, and gestation at PROM. The primary complication for the mother is risk of infection. Complications of PROM for the fetus and newborn consist of prematurity, fetal distress, cord compression, deformation and altered pulmonary development leading to pulmonary hypoplasia and pulmonary hypertension, necrotizing enterocolitis (NEC) and neurologic disorder. Infectious morbidities in

mother, fetus and newborn have been related to both PROM and prolonged rupture of membranes [6].

Objectives of the study

The main objective of the study is to analyze the maternal and perinatal outcome in preterm prelabor rupture of membranes.

MATERIAL AND METHODS:

This cross sectional study was conducted in Ganga Ram Hospital, Lahore during June 2019 to January 2020. Patients admitted in Obstetrics and Gynaecology Unit through outpatient & casualty department with gestational age between 24 to 37 weeks with preterm premature rupture of membrane (PPROM) confirmed by ultra sound and clinical examination were included regardless of their age. Patients with multiple gestation, obstetric bleeding, fetal anomaly, allergic reactions, intrauterine death and patients who refused to participate in the study were excluded. Successful tocolysis was labeled as prolongation of pregnancy for minimum 48 hours (corticosteroids were given in this time for fetal lung maturity) after therapy of tocolysis. Failed or unsuccessful tocolysis was labeled when delivery occurred within 48 hours. After admission, detailed workup including history, general physical examination, abdomen and pelvic examination and relevant /specific investigations were noted.

Statistical analysis

All data were analyzed using SPSS version 21. Post stratification chi square test was applied to see effect modification. $P < 0.05$ was considered as significant.

RESULTS:

A total of 72 female patients were included in the study. Mean age and gestational age of the patients was 25.36 ± 3.8 years and 32.09 ± 1.78 weeks. 56.9% (n=41) patients had 15-25 years of age and 43.1% (n=31) patients fell in age group 26-40 years. Gestational age distribution showed that majority of the patients i.e. 58.3 (n=42) had gestational age 29-32 weeks and 41.7% (n=30) had 33-36 weeks of gestational age. The cause of mortality in the present study was sepsis. All of them were between 28 weeks-29 weeks and latency period of <24hrs. Sepsis was the cause of neonatal mortality in our study.

Table 01: Analysis of Maternal morbidity

Maternal complications	Early PROM	Late PROM
Chorioamnionitis	12%	2%
Abruption	16%	4%
No complications	72%	94%

Table 2. Analysis of perinatal mortality

Gestational age	Latency period (hrs)	Birth weight (kg)	Cause of mortality
29 weeks	12	1.3	Sepsis, pulmonary hemorrhage
28 weeks	06	1.2	Sepsis, DIC
28 weeks	05	1.0	Still born

DISCUSSION:

Chorioamnionitis was more common in indicated deliveries as they had prolonged latency period and they were prone to infection during this period. In a study by Bryden et al also showed 67% of chorioamnionitis among early PPRM were in indicated delivery.12 Another study by Jenny et al showed 1.6% of chorioamnionitis in late PPRM.11 In the present study (16%) had abruption in early PPRM, no abruption in late PPRM [7]. Due to their adverse effects and short outcomes these drugs were replaced with oxytocin, Ca+ channel openers and blockers. A new drug nifedine has been introduced for reduction of uterine contractions [8]. It is a Ca+ channel antagonist that blocks the Ca+ influx in cells of myometrium and oral intake gains peak plasma level within 45 minutes and has half life 2-3 hours their use [9]. In our study, we observed that in nifedipine 88.9% patients had prolonged pregnancy while in nitroglycerine 61.1% patients had prolong pregnancy. Conde-Agudelo et al conducted a study in 2011 on comparison between nifedipine and magnesium sulphate in management of preterm labor. They reported that nifedipine was more effective in α 2-adrenergic agonist as Mg sulphate is also [10-11].

CONCLUSION:

It is concluded that most common maternal morbidity associated with PPRM was chorioamnionitis, that of neonatal morbidity was prematurity and its complications.

REFERENCES:

- Jung EJ, Byun JM, Kim YN, Lee KB, Sung MS, Kim KT, et al. Antenatal magnesium sulfate for both tocolysis and fetal neuroprotection in premature rupture of the membranes before 32 weeks' gestation. *J Maternal-Fetal Neonatal Med* 2017;26:1-11.
- Lipi LB, Begum N, Alam UK, Jahan R, Rahman MM, Rumana R. Study on role of magnesium Sulphate as a Tocolytic agent in preventing preterm labour. *J Dhaka Med Coll* 2015;22:179-84.
- Habib H, Mustafa SG, Gul Y. Transdermal Nitroglycerine as a Tocolytic in Preterm Labor. *IJSRP* 2014;4:1-4.
- Rust OA, Bofill JA, Arriola RM, Andrew ME, Morrison JC. The clinical efficacy of oral tocolytic therapy. *AJOG* 1996;175:838-42.
- Magann EF, Bass D, Chauhan SP, Sullivan DL, Martin RW, Martin JN. Antepartum corticosteroids: disease stabilization in patients with the syndrome of hemolysis, elevated liver enzymes, and low platelets (HELLP). *AJOG* 1994;171:1148-53.
- Iams JD, Romero R, Culhane JF, Goldenberg RL. Primary, secondary, and tertiary interventions to reduce the morbidity and mortality of preterm birth. *The Lancet* 2008;371(9607):164-75.
- Asghar A, Xia L. To Compare Efficacy of Nifedipine and Nitroglycerine as Tocolytic Agent in Preterm Labor patients. *PJMHS* 2014;4:1-5.
- Chan L, Sahota D, Yeung S, Leung T, Fung T, Lau T, et al. Side-effect and vital sign profile of nifedipine as a tocolytic for preterm labour. *Hong Kong Med J* 2008;14:267-72.
- Perveen S, Araainuddin J, Naz S. Short term tocolytic efficacy of transdermal nitroglycerine. *Med Channel* 2010;16:152-4.
- Noor S, Nazar AF, Bashir R, Sultana R. Prevalance of PPRM and its outcome. *J Ayub Med Coll Abbottabad*. 2007;19(4):14-7.
- Bartfield MC, Carlan SJ. The home management of preterm premature ruptured membranes. *Clini Obstet Gynecol*. 1998;41(3):503-14.