ISSN: 2319-7064 SJIF (2022): 7.942

Incidence and Risk Factors of PPROM and its Maternal and Neonatal Outcomes at Different Gestational Ages

Dr. Jasmine Gulati¹, Dr. Arvind², Dr. Kulwinder Kaur³, Dr. Ashwani Kumar⁴

Abstract: Introduction: In countries with high and low incomes, preterm premature rupture of membranes (PPROM) is a major contributor to perinatal, neonatal, and maternal morbidity and mortality. The impact of PPROM is exacerbated in developing countries as a result of various factors related to the standard of healthcare provided and socioeconomic factors. Aims & Objectives: To determine the incidence and risk factors of PPROM. We also assess the obstetric and neonatal outcomes at different gestational ages. Methodology: This was a prospective study conducted in the Department of Obstetrics and Gynecology, Bebe Nanki Mother and Child Care Centre, Guru Nanak Dev Hospital, Amritsar on the pregnant women attending the indoor or outdoor services between February 2021 and May 2022 with preterm premature rupture of membrane (PPROM). Results: Nearly 63.6%neonates had noneonatal morbidity while 36.4%had neonatalmorbidity in form of jaundice, RDS, septicemia, meningitis and hypoglycemia. Maternal outcome and Neonatal outcome at different gestational age was found to be statistically significant (p<0.05). Latency period, abnormal vaginal discharge, history of PROM etc. were found to be significant maternal risk factors on statistical analysis (p<0.05). Conclusion: Based on the findings of the present study, we conclude that abnormal vaginal discharge, vaginal bleeding, previous premature rupture of membrane etc. were associated with preterm premature rupture of membrane. Thus, early screening and treatment of urinary tract infections and abnormal vaginal discharges were recommended to reduce the risk of preterm premature rupture of membrane.

Keywords: Culture and sensitivity, neonatal outcome, chorioamniotis, maternal risk factors, CRP

1. Introduction

Prelabor rupture of membranes (PROM), or the rupture of membranes before the onset of labor and before 37 weeks of gestation, occurs in 2–3% of pregnancies. It is a severe pregnancy complication that carries risks like premature birth, infection of the mother or the baby, and respiratory distress syndrome. [1, 2] The major risk factors for PPROM include preterm prelabour rupture of membranes in a prior pregnancy, vaginal bleeding, anaemia/low BMI (Body Mass Index<19kg/m2), preterm labour, low socioeconomic status, other uteroplacental and foetal factors etc.

Most studies addressed the fact that in cases of expectant management of PPROM, combined use of antibiotics and corticosteroids appear to confer greater benefits in the reduction of adverse neonatal outcomes (e.g. respiratory distress syndrome (RDS) and intracranial hemorrhage) in the absence of maternal and fetal infectious compromise. Although concern persists over the development of resistant strains of organisms involved in neonatal sepsis, current data support the use of antibiotics in this setting.[3,4] Nor is there currently great evidence to support routine use of tocolytics in cases with PPROM; however, tocolysis for a limited period of time (48 h) after preterm amniorrhexis may justify consideration. There is paucity of data regarding incidence, risk factors and outcome of PPROM in Indian population. Hence, the present study aimed to determine the incidence and risk factors of PPROM and evaluate the obstetric and neonatal outcomes at different gestational ages. This study was being done with a view to improve the safety and outcomes in pregnant women with PPROM.

2. Material & Methods

Study design: Prospective observational study Place of Study: Guru Nanak Dev Hospital, Amritsar Sample size: 300pregnant women with preterm premature rupture of membrane

Sampling technique: This was a prospective, observational study

Period of study: February 2021 - May 2022

Inclusion Criteria

All the pregnant women with pregnancies <37 weeks of gestational age with premature rupture of membranes.

Exclusion Criteria:

- Multiple pregnancies.
- Intrauterine growth restriction.
- Fetal Anomalies.
- Intrauterine fetal death (IUFD).

Clinical examination included the following:

- Pulse
- · Blood pressure
- Abdominal palpation to confirm presentation
- Uterine contraction
- Fetal heart

Other Investigations included the following:

- Complete Blood Count
- Platelet count
- Total Leukocyte Count
- Differential Leukocyte Count
- C Reactive Protein
- Liver Function Tests
- Renal Function Tests
- Urine culture
- Vaginal culture and sensitivity tests
- Ultrasonography

Volume 12 Issue 5, May 2023

www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

Paper ID: MR23511124024 DOI: 10.21275/MR23511124024 954

ISSN: 2319-7064 SJIF (2022): 7.942

All data was tabulated and subjected to appropriate statistical analysis using SPSS version 21.0.

Ethical approval was obtained from institutional ethical committee and written consent was obtained from all patients after explaining in detail the entire research protocol.

3. Results

Descriptive

The majority of study participants (65%) were in the age group 21 to 30 years with 57.3% from the rural areas, whereas remaining 42.7% resided in the urban areas. The majority of participants in the study (60%) belonged to lower middle class, 26.7% had completed education up to middle school, followed by 19.3% participants who completed intermediate. (Table 1)

Table 1: Patient demographics

Characteristics	N (%)					
Age Group (Years)						
<20	43 (14.3)					
21-30	195 (65.0)					
>30	62 (20.7)					
Residence						
Rural	172					
Urban	128					
Socioeconomic Status						
Upper Class	18 (6.0)					
Upper Middle	6 (2.0)					
Lower Middle	180 (60.0)					
Upper Lower	42 (14.0)					
Lower	54 (18.0)					
Educational Qualification						
Illiterate	48 (16.0)					
Primary School	32 (10.7)					
Middle School	80 (26.7)					
High School	45 (15.0)					
Intermediate	58 (19.3)					
Graduate And Above	37 (12.3)					

Incidence

In the present study,150 participants (50%) were between 33-37 weeks of gestation,123 participants (41%) between 28-32 weeks and 27 participants (9%) with gestational age <28 weeks.

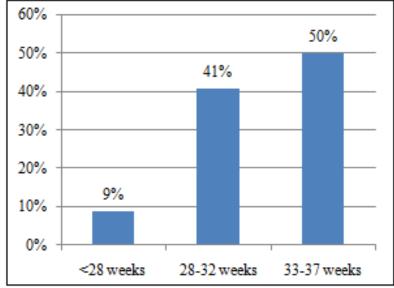


Figure 1: Distribution of study participants with PPROM at different gestational ages

Licensed Under Creative Commons Attribution CC BY

955

ISSN: 2319-7064 SJIF (2022): 7.942

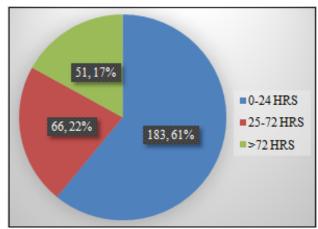


Figure 2: Represents the graphical distribution of latency period at different gestational ages.

Risk Factors

Table 2 depicts distribution of risk factors at different gestational ages. Vaginal bleeding, abnormal vaginal discharge, history of PROM, history of preterm birth, mode of delivery, hours of leakage and diabetes mellitus were

found to be significant risk factors on statistical analysis (p<0.05).

956

Table 2: Risk factors of PPROM at different gestational ages

		Tat	ole 2: Ris.	k factors o	f PPROM a		gestational ag			
Risk Factors		Total		Gestational Age <28 Weeks (N=27)		Gestational Age 28-32 Weeks (N=123)		Gestational Age 33-37 Weeks (N=150)		P Value
		NO.	%AGE	NO.	%AGE	NO.	%AGE	NO.	%AGE	
Vaginal Bleeding	Yes	54	18.0%	18	66.7%	22	17.9%	14	9.3%	<0.00001
	No	246	82.0%	9	33.3%	101	82.1%	136	90.7%	
Abnormal Vaginal Discharge	Yes	222	74.0%	13	48.1%	99	80.5%	110	73.3%	0.0023
	No	78	26.0%	14	51.9%	24	19.5%	40	26.7%	
II:	Yes	133	44.3%	20	74.1%	53	43.1%	60	40.0%	0.0043
History of Prom	No	167	55.7%	7	25.9%	70	56.9%	90	60.0%	
History of LITI	Yes	33	11.0%	3	11.1%	14	11.4%	16	10.7%	0.9823
History of UTI	No	267	89.0%	24	88.9%	109	88.6%	134	89.3%	
History of	Yes	64	21.3%	18	66.7%	20	16.3%	26	17.3%	<0.00001
Preterm Birth	No	236	78.7%	9	33.3%	103	83.7%	124	82.7%	
Foetal	Cephalic	252	84.0%	20	74.1%	102	82.9%	130	86.7%	0.2371
Presentation	Breech	48	16.0%	7	25.9%	21	17.1%	20	13.3%	
II£I1	<12 Hr	133	44.3%	20	74.1%	53	43.1%	60	40.0%	0.0043
Hours of Leakage	>12 Hr	167	55.7%	7	25.9%	70	56.9%	90	60.0%	
A : -	Yes	160	53.3%	15	55.6%	75	61.0%	70	46.7%	0.0602
Anemia	No	140	46.7%	12	44.4%	48	39.0%	80	53.3%	
Diabetes Mellitus	Yes	168	56.0%	8	29.6%	70	56.9%	90	60.0%	0.005
	No	132	44.0%	19	70.4%	53	43.1%	60	40.0%	
Hypertension	Yes	34	11.3%	1	3.7%	17	13.8%	16	10.7%	0.3030
	No	266	88.7%	26	96.3%	106	86.2%	134	89.3%	

P<0.005 was considered significant.

Table 3 depicts distribution of neonatal risk factors at different gestational ages. Birth weight, resuscitation and outcome were found to be significant neonatal risk factors on statistical analysis (p<0.05).

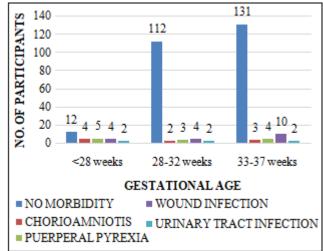
Volume 12 Issue 5, May 2023 www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

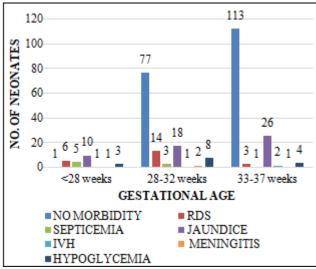
ISSN: 2319-7064 SJIF (2022): 7.942

Table 3.	Neonatal	risk factors	at different	gestational ages

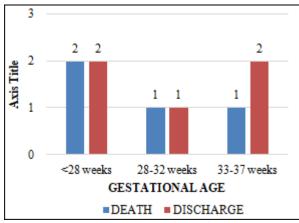
Tuble of Hooling it is interest geometrical ages										
Neonatal Characteristics		Total		Gestational Age <28 Weeks		Gestational Age 28-32 Weeks		Gestational Age 33-37 Weeks		P Value
				No.	%AGE	No.	%AGE	No.	%AGE	
		Birth Weight (KG)	<1.5	40	13.30%	6	22.20%	12	9.70%	22
1.5-2.0	80		26.70%	3	11.10%	30	24.40%	47	31.30%	
2.0-2.5	98		32.70%	14	51.90%	61	49.60%	23	15.30%	
>2.5	82		27.30%	4	14.80%	20	16.30%	58	38.70%	
APGAR	<5	2	0.70%	1	3.70%	1	0.80%	0	0%	0.081
Score at 5 Min	>5	298	99.30%	26	96.30%	122	99.20%	150	100%	
D '(()	Yes	64	21.30%	18	66.70%	20	16.30%	26	17.30%	< 0.00001
Resuscitation	No	236	78.70%	9	33.30%	103	83.70%	124	82.70%	<0.00001
Congenital Anomaly	Yes	6	2.00%	2	7.40%	2	1.60%	2	1.30%	0.1077
	No	294	98.00%	25	92.60%	121	98.40%	148	98.70%	
Outcome	Death	14	4.70%	9	33.30%	4	3.30%	1	0.70%	< 0.00001
	Discharge	286	95.30%	18	66.70%	119	96.70%	149	99.30%	



Maternal Outcomes



Neonatal Outcomes



Outcomes with chorioamnionitis

Outcomes

Figure 3 shows graphical presentation of Maternal, Neonatal outcome and outcome in patients with chorioamnionitis at different gestational age.

4. Discussion

In the present study,65% of the participants were in the age group 21 to 30 years. Surayapalem S et al. [3] observed that PROM was common in age group of 20-24 years with mean age of 22.6 years and SD of 2.8 years. In our study, 57.3% of participants were from rural areas and 60% belonged to lower middle class. Dars S et al. [4] also concluded that 72% of them belonged to lower middle class. The factors leading to higher incidence of PPROM in low socio-economic status include poor hygiene, malnutrition, anaemia, stress, over exertion, high parity, recurrent genitourinary infections etc.

Latency period, vaginal bleeding, abnormal vaginal discharge, history of PROM, history of preterm birth, mode of delivery, hours of leakage and diabetes mellitus were found to be significant maternal risk factors. Kannan MP et al. [5] identified that diabetes mellitus, poor weight gain and history of previous preterm birth were the factors associated with PPROM significantly. Bouvier D et al.[6] observed that the specific risk factors for PPROM were history of PPROM, nulliparity, gestational diabetes, and low level of education. Addisu et al. [7] identified significant risk factors for PPROM were abnormal vaginal discharge, urinary tract

957

Volume 12 Issue 5, May 2023

www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

ISSN: 2319-7064 SJIF (2022): 7.942

infection, history of premature rupture of membranes and vaginal bleeding.

In the present study, 61% patients had latency period below 24 hours and 73.3% with gestational age of 33-37 weeks had latency period <24 hours which is statistically significant. In a study by Yu Het al.[8], there was a strong inverse correlation between gestational age at PPROM and the latency period. The duration of the latency period was significantly longer among woman with PPROM before 34 weeks as observed by Test G et al. [9]

Similar to findings by Addisu D et al. [7] vaginal bleeding and previous history of PROM were found to be a significant risk factor in this study. Women who had a history of vaginal bleeding in current pregnancy were 2.58 times more likely to develop PPROM. Pregnant women who had history of PROM were 3.31 times more likely to develop PPROM. Similar to findings by Assefa NE et al. [10] a significant association was noted between abnormal vaginal discharge and PPROM. Abnormal vaginal discharge is a common symptom of genital infections. Inflammatory cells produced by genital infections are involved in weakening of the fetal membranes among pregnant women thus causing PPROM. [11]

Having gestational diabetes mellitus was found to be a significantly associated factor for preterm rupture of membrane in the present study. This is consistent with findings of Bouvier D et al.[6], and Harger JH et al.[11]. Maternal morbidity was noticed in 15% cases, which is comparable to findings of Mohokar SA et al. [12], Surayapalem S et al. [3] and Galletta MAK et al.[13]. However, in study by Pisoh DW et al.[14], only 10.53% of women who had PPROM had an unfavourable outcome.

In the present study, birth asphyxia, septicemia and respiratory distress syndrome were main causes for neonatal mortality. Kannan MP et al. [5] found birth asphyxia, low birth weight and prematurity to be significant neonatal risk factors. Neonatal morbidity was observed in 36.3% cases in this study with most common being jaundice, followed by respiratory distress syndrome, septicaemia and IVH. Neonatal outcome at different gestational age was found to be statistically significant in this study. In a similar study by Singh N et al. [15], birth asphyxia was seen in 12% of patients, jaundice in 12%, and septicemia in 4% of the subjects. The neonatal mortality rate in present study was 4.7%. Yu H et al. [8] and Mohokar SA et al. [12] reported neonatal mortality rate of 7.4% and 15% respectively. A low mortality rate of 3.3% was observed by SinghN et al. [15]

5. Conclusion

Majority of PPROM patients were with gestational age 33-37 weeks with maternal age 21-30 years in our study. Latency period, abnormal vaginal discharge, vaginal bleeding, history of prom, history of preterm birth and hours of leakage were found to be major maternal risk factors and birth weight, apgar score at 1 minute and resuscitation were the major neonatal risk factors. Hospitalisation of the study group population with PPROM contributes to increased economic burden. An appropriate and accurate diagnosis of

PPROM is critical to optimize pregnancy outcome. Patient is suggested that timely diagnosis and management of PPROM will allow obstetric care provider to optimize perinatal outcome and minimize neonatal mortality.

References

- [1] Rib DM, Sherer DM, Woods JR Jr. Maternal and neonatal outcome associated with prolonged premature rupture of membranes below 26 weeks' gestation. Am J Perinatol 1993;10:369–373.
- [2] Cox SM, Williams ML, Leveno KJ. The natural history of preterm ruptured membranes: What to expect of expectant management. Obstet Gynecol 1988;71:558–562.
- [3] Surayapalem S, Cooly V, Salicheemala B. A study on maternal and perinatal outcome in premature rupture of membranes at term. International Journal of Reproduction, Contraception, Obstetrics and Gynecology. 2017;6(12):5368.
- [4] Dars S, Malik S, Samreen I, Kazi RA. Maternal morbidity and perinatal outcome in preterm premature rupture of membranes before 37 weeks gestation. Pak J Med Sci. 2014;30(3):626-9.
- [5] Kannan MP, Bhat BV, Sharma R, Ferns S, Rani R. Perinatal mortality- Identification of risk factors. Journal of Obstetrics and Gynaecology of India. 2003;53(3):240-3.
- [6] Bouvier D, Forest JC, Blanchon L, Bujold E, Pereira B, Bernard N, et al. Risk Factors and Outcomes of Preterm Premature Rupture of Membranes in a Cohort of 6968 Pregnant Women Prospectively Recruited. J Clin Med. 2019;8(11):1987.
- [7] Addisu D, Melkie A, Biru S. Prevalence of Preterm Premature Rupture of Membrane and Its Associated Factors among Pregnant Women Admitted in Debre Tabor General Hospital, North West Ethiopia: Institutional-Based Cross-Sectional Study. Obstet Gynecol Int. 2020;2020:4034680.
- [8] Yu H, Wang X, Gao H, You Y, Xing A. Perinatal outcomes of pregnancies complicated by preterm premature rupture of the membranes before 34 weeks of gestation in a tertiary center in China: A retrospective review. Biosci Trends. 2015; 9(1):35-41.
- [9] Test G, Levy A, Wiznitzer A, Mazor M, Holcberg G, Zlotnik A, et al. Factors affecting the latency period in patients with preterm premature rupture of membranes. Arch Gynecol Obstet. 2011;283(4):707-10.
- [10] Assefa NE, Berhe H, Girma F, Berhe K, Berhe YZ, Gebreheat G, et al. Risk factors of premature rupture of membranes in public hospitals at Mekele city, Tigray, a case control study. BMC Pregnancy Childbirth. 2018;18(1):386.
- [11] Harger JH, Hsing AW, Tuomala RE, Gibbs RS, Mead PB, Eschenbach DA, Knox GE, Polk BF. Risk factors for preterm premature rupture of fetal membranes: a multicenter case-control study. Am J Obstet Gynecol. 1990;163(1 Pt 1):130-7.
- [12] Mohokar SA, Bava AK, Nandanwar YS. Analysis of maternal and perinatal outcome in cases of preterm premature rupture of membranes. Bombay Hospital J. 2015;57(3):285-90.

958

Volume 12 Issue 5, May 2023 www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

ISSN: 2319-7064 SJIF (2022): 7.942

- [13] Galletta MAK, Bittar RE, Agra I, Guerra ECL, Francisco RPV, Zugaib M. Epidemiological profile of patients with preterm premature rupture of membranes at a tertiary hospital in São Paulo, Brazil. Clinics (Sao Paulo). 2019;74:e1231.
- [14] Pisoh DW, Mbia CH, Takang WA, Djonsala OGB, Munje MC, Mforteh AA, et al. Prevalence, Risk Factors and Outcome of Preterm Premature Rupture of Membranes at the Bamenda Regional Hospital. Open Journal of Obstetrics and Gynecology.2021; 11:233-51.
- [15] Singh N, Pattnaik L, Panda SR, Jena P, Panda J. Fetomaternal Outcomes in Women Affected With Preterm Premature Rupture of Membranes: An Observational Study From a Tertiary Care Center in Eastern India. Cureus. 2022; 14(5):e25533.

Author Profile

Dr. Jasmine, junior resident, Department of Obstetrics and Gynaecology, Government Medical College, Amritsar

Dr. Arvind, Intern, Maulana Azad Medical College, Delhi

Dr. Kulwinder Kaur, Associate Professor, Department of Obstetrics and Gynaecology, Government Medical College, Amritsar

Dr. Ashwani Kumar, Professor, Department of Pediatrics, Government Medical College, Amritsar

Volume 12 Issue 5, May 2023 www.ijsr.net

Licensed Under Creative Commons Attribution CC BY

959