

DOI: <http://dx.doi.org/10.18203/2320-1770.ijrcog20184147>

Original Research Article

Role of amniotic fluid index on maternal and neonatal outcomes among obstetric women with preterm premature rupture of membranes

Jennifer Britto J.*, Sailatha R., Amrita Priscilla Nalini

Department of Obstetrics and Gynecology, Chettinad Hospital and Research Institute, Chennai, Tamil Nadu, India

Received: 09 August 2018

Accepted: 04 September 2018

*Correspondence:

Dr. Jennifer Britto J.,

E-mail: drjenni12345@gmail.com

Copyright: © the author(s), publisher and licensee Medip Academy. This is an open-access article distributed under the terms of the Creative Commons Attribution Non-Commercial License, which permits unrestricted non-commercial use, distribution, and reproduction in any medium, provided the original work is properly cited.

ABSTRACT

Background: Preterm Premature rupture of Membrane (PPROM) is one of the major complications of a pregnant women with risk factors like low socioeconomic class, infections, STDs etc. Maternal complications such as Chorioamnionitis, Abruptio placenta, sepsis and neonatal complications like neonatal sepsis, RDS, intraventricular hemorrhage are common with those with PPRM. The objectives of the present study were to find the association between AFI values and maternal and fetal outcomes in patients with preterm premature rupture of membranes

Methods: A cross sectional study was conducted in a Multispecialty Teaching Hospital, North Chennai, Tamil Nadu with the sample size of 100. The study participants included pregnant women with gestational age of 28 to 34 weeks presenting with PPRM. Maternal outcomes included clinical diagnosis of chorioamnionitis, placental abruption, meconium in liquor, fetal distress, prolapsed cord and mode of delivery. Neonatal outcomes include neonatal hospitalization in NICU, APGAR score at 1st minute and 5 minutes, early neonatal sepsis neonatal death. Significance of difference in means was calculated using independent t test.

Results: Total study participants were divided into 2 groups with Amniotic Fluid Index (AFI) of >5 and <5. Latency period, Chorioamnionitis, placental abruption and mode of delivery were compared between the groups. Chorioamnionitis was present in 8% and 12% in group 1 and group 2 respectively. Neonatal outcomes like NICU admission, RDS and neonatal sepsis were also compared between the groups. NICU admission was required for 18% and 48% in group 1 and 2 respectively. APGAR scores at 1 and 5 minutes were found to be lower among those with Group 2. (p=0.000).

Conclusions: PROM is associated with increased maternal and neonatal complications. AFI has been proven a role in predicting maternal and neonatal outcomes in PROM. Identifying the risk factor and its treatment remains the main mode of prevention for PROM.

Keywords: Abruptio placenta, Chorioamnionitis, NICU, PPRM

INTRODUCTION

Preterm premature rupture of the membranes (PPROM) is one of the fatal complications during pregnancy with significant perinatal complications. PPRM is defined as the rupture of the amniotic membranes before 37 weeks of gestation and before the onset of labour.¹ PPRM constitutes one third of the preterm births and it complicates about 3% of pregnancies.² Predisposing

factors of PPRM are low social economic class, previous history of preterm labour, smoking, infections, sexually transmitted diseases, genetic and/or enzymatic abnormalities, nutritional deficiencies, intervention such as circlage and amniocentesis.³⁻⁵

Major maternal risks are chorioamnionitis (35%), abruption (19%) and sepsis (<1%). Abruptio placenta is frequently seen if the rupture of membrane occurred

before 28 weeks of gestation.⁶ In the presence of intrauterine infection or oligohydromnios, the risk of abruption is high.⁷

Fetal morbidity due to PPROM includes pulmonary hypoplasia, Respiratory Distress Syndrome, sepsis, intraventricular hemorrhage and neonatal sepsis.⁸ Presence of adequate amniotic fluid prevent infections and thereby reducing the occurrence of PPROM. Thus oligohydramnios is also considered as risk factor for PPROM.⁹ When the presentation is non cephalic in PPROM there are risks such as shorter latency and increased neonatal morbidity, to be increased when oligohydramnios is present.¹⁰ In addition, there are reports about unfavourable impact by noncephalic presentation of the fetus with PPROM on the antepartum, intrapartum and neonatal risks, which were primarily due to cord prolapse.¹¹ The objective of the present study was to find the association between AFI values and maternal and fetal outcomes in patients with preterm premature rupture of membranes.

METHODS

The study was conducted in a Multispecialty Teaching Hospital located in North Chennai of Tamil Nadu, India. The study was a cross sectional type of study. The study included patients presenting to Labour room with Preterm PROM from 28 to 34 weeks period of gestation (POG). Inclusion criteria was pregnant women with gestational age 28 to 34 weeks presenting with leaking per vaginam. Exclusion criteria includes - symptoms of chorioamnionitis at the time of admission, history of previous caesarean section or previous uterine surgery, non-cephalic presentation, Intrauterine growth restriction, maternal diseases complicating pregnancy such as preeclampsia, diabetes, eclampsia etc.

The study was carried out between June 2015- October 2016. Sample size was calculated with the formula for cross sectional studies, i.e. $N = Z^2pq/L^2$.

The incidence of preterm premature rupture of membranes from previous studies in India ranges from 3.0-10.0% of all deliveries.¹² With the incidence of PPROM at 7%, $Z=1.96$ and error of 5%, the sample size calculated was 100.

Patients fulfilling inclusion criteria and willing to participate in the study were included successively-consecutive sampling method was followed. AFI value of the previous visit was taken to divide the study population into two groups. In the present study, two groups were made-those with AFI >5 and those with AFI < 5. Thus, enrollment was continued till each group consisted 50 patients.

Participants who fulfilled the inclusion criteria were hospitalized, admitted in labour room and are monitored for 12hours in view of emerging contractions, bleeding or

possible start of delivery after NST and fetal heart rate monitoring. Those with the complaints of leaking per vaginam with confirmed leaking were determined by using sterile speculum examination using the pooled fluid and the secretions were examined by fern test under microscopic and litmus paper test. Ultrasonogram was performed on all the participants during first 12 to 24 hours to measure amniotic fluid in four abdominal quadrants in order to determine AFI. Participants were divided into 2 groups according to their AFI. Supportive management such as administration of single course of betamethasone were done to the patients at admission (2 doses of inj betamethasone 12mg, 24 hours apart) for fetal lung maturity.

Antibiotic prophylaxis for the participants includes inj ampicillin 1gram i.v. 6hourly for first 48hours was given and then followed by cap amoxicillin 500mg 8hourly orally till the delivery. During hospitalization fetal heart rate was monitored for every 2 hours, daily non-stress tests was performed for fetus with gestational age >28weeks.

The patients were observed for clinical symptoms of chorioamnionitis such as fever, uterine tenderness, maternal tachycardia, fetal tachycardia, laboratory symptoms (leucocytosis). If symptoms suggested start of clinical chorioamnionitis, antibiotics were given and if delivery is not started, labour was induced.

Maternal outcomes such as period of latency, clinical diagnosis of chorioamnionitis, placental abruption, meconium in liquor, fetal distress, prolapsed cord and mode of delivery. Neonatal outcomes such as neonatal hospitalization in NICU, duration in NICU stay, APGAR score at 1st minute and 5 minutes, RDS, early neonatal sepsis and early neonatal death.

Data analysis was done with Statistical Package for Social Sciences (SPSS IBM) version 21.0. The qualitative variables are described in the form of proportions and quantitative variables are described in the terms of mean, range and standard deviation. Data was checked for normality before applying appropriate tests of significance.

Significance of difference in means was calculated using independent t test. Significance of p value was taken as $p < 0.05$. Ethical permission was obtained from Institutional Ethics Committee of the hospital. The confidentiality of the study participants was maintained at all points of the study.

RESULTS

Baseline characteristics

Among the total 100 (100%) study participants- there were two groups with each group consisted of 50. Group1 were those with AFI >5 and Group 2 with AFI <5. The

age of the study participants varied between 20 years to 35 years. The mean (SD) age of study participants in group 1 and group 2 was 28(\pm 2.96) and 29(\pm 2.92) respectively. (Table 1)

Mean (SD) hemoglobin of participants in Group 1 was 10.8(\pm 2.5) gms whereas it was 11.9(\pm 1.6) in group 2. Random blood sugar values were 93 and 92mg/dl in group1 and group 2 respectively.

Table 1: Baseline parameters of study participants. (n=100).

Profile of study participants	Group 1 N=50 N (%)	Group 2 N=50 N (%)
Age group		
\leq 20 years	1(2)	0
21-30 years	38(76)	31(62)
>30 years	11(22)	19(38)
Gravida		
Primigravida	38(76)	40(80)
Multigravida	12(24)	10(20)
Residence		
Urban	9(18)	11(22)
Rural	41(82)	39(78)

Ultrasound features

With ultrasound, certain parameters such as gestational age, amniotic fluid index, estimated fetal weight and placental position were determined.

Mean (SD) values of gestational age of group 1 and 2 were 31(\pm 2) and 32(\pm 2) respectively. Mean (SD) value of group 1 and 2 amniotic fluid indices was 10.2(\pm 2.2) and 4.5(\pm 0.7) respectively. Estimated fetal weight (in kilogram) of group 1 and 2 was 2.6(\pm 0.23) and 2.0(\pm 0.29) respectively. Comparison of maternal outcomes such as latency period, chorioamnionitis, placental abruption and mode of delivery was done with two groups. Chorioamnionitis was present in 4(8%) of participants in group1 and 6(12%) in group 2. Latency period of < 48 hours in 82% of study participants in group 1 and 92% of study participants. Latency period of > 48 hours in 18% and 8% in group 1 and group 2 respectively.

Abruptio placenta was seen in 10% of study participants in group 1 and 14% of study population in group 2. Chorioamnionitis, abruption placenta and cord prolapsed were found to be higher in group 2 (AFI<5). Proportion of caesarean section was found to be much higher in group 2 when compared to group 1. This was found to be significant (P value =0.042) (Table 2).

Table 2: Distribution of study participants according to maternal outcomes. (n=100).

Maternal outcomes	Group 1, N=50, N (%)	Group 2, N=50, N (%)	OR (95%CI)	P value
Period of latency				
<48 hours	41(82)	46(92)	0.396 (0.11-1.38)	0.137
>48 hours	9(18)	4(8)		
Chorioamnionitis				
Yes	4(8)	6(12)	1.00 (0.32-3.08)	0.121
No	46(92)	44(88)		
Abruptio placenta				
Present	5(10)	7(14)	0.683 (0.20-2.31)	0.583
Absent	45(90)	43(86)		
Meconium stained liquor				
Yes	11(22)	10(20)	0.243 (0.01-1.78)	0.346
No	39(78)	40(80)		
Cord prolapse				
Yes	1(2)	2(4)	0.787 (0.21-1.56)	0.252
No	49(98)	48(96)		
Mode of delivery				
Vaginal	20(40)	12(24)	2.068 (1.70-6.11)	0.042
Caeserean section	30(60)	38(76)		

Chi square test applied, p value <0.05 is significant

Comparison of neonatal outcomes such as NICU admission, RDS and neonatal sepsis was done with two groups. Among the neonatal outcomes, higher NICU admission was required in group 2 and this was found to

be statistically significant. (p value =0.001). NICU admission was required in 18% of the neonates in Group 1 whereas 48% of neonates required NICU admission in Group 2. RDS developed in 12 % and 8% of neonates in

Group 1 and Group 2 respectively. Neonatal mortality among group 1 was 4% whereas in group 2 it was 2%. Likewise, more than 60 days NICU stay was required in

20% of group1 neonates. Upto 28 days admission in NICU was required in 24% of group 1 neonates and 48% of group 2 neonates (Table 3).

Table 3: Distribution of study participants according to neonatal outcomes. (n=100).

Neonatal outcomes	Group 1, N=50, N (%)	Group 2, N=50, N (%)	OR (95%CI)	p value
NICU admission				
Yes	9 (18)	24 (48)	2.238 (1.09-3.59)	0.001
No	41 (82)	26 (52)		
Duration of neonatal stay				
0-28 days	12 (24)	24 (48)	-	0.560
28-45 days	18 (36)	18 (36)		
46-60 days	10 (20)	8 (16)		
>60 days	10 (20)	0		
Respiratory distress syndrome				
Yes	4 (8)	6 (12)	0.306 (0.059-1.59)	0.043
No	46 (92)	44 (88)		
Neonatal sepsis				
Yes	2 (4)	7 (14)	1.397 (0.44-4.367)	0.128
No	48 (96)	43 (86)		
Early neonatal death				
Yes	2 (4)	1 (2)	2.12 (0.786-4.32)	0.140
No	48 (96)	49 (98)		

Chi square test applied, p value <0.05 is significant

Mean (\pm SD) of 1-minute APGAR among group 1 was 7.84(\pm 0.58) and 5.5(\pm 1.41) in group 2. Mean (\pm SD) of 5-minute APGAR among group 1 was 8.96(\pm 0.49) and 7.38(\pm 1.42) in group 2. APGAR scores at 1 and 5

minutes were compared between the two groups. It was found from the analysis that APGAR scores at 1 and 5 minutes were found to be lower among those with Group 2 when compared to those Group 1. This was found to be statistically significant. (p =0.000) (p=0.000) (Table 4)

Table 4: Comparison of mean APGAR scores among groups (N=100).

Variables	Mean (SD)	t value	Mean Difference	95% CI	p value
APGAR (1')					
Group1	7.84 (±0.58)	10.52	2.28	1.850-2.710	0.000
Group 2	5.5 (±1.41)				
APGAR (5')					
Group 1	8.96 (±0.49)	7.40	1.58	1.156 -2.004	0.000
Group 2	7.38 (±1.42)				

Independent t test applied, p value<0.05 is significant

DISCUSSION

In the present study,80% of study participants were from rural area and 20% were from urban area. PPROM was found frequently with low socioeconomic class and rural area. Higher chances of infections due to unhygienic living conditions in rural area is an independent risk factor for PPROM.¹³⁻¹⁴ PPROM was not age specific. Majority (69%) belong to age group 21-30 years of age. Similar finding was reported by Gandhi M et al (77.6%)

and Kumari BR et al (49%). 78% were primigravida in present study. This is similar to previous study finding as well- 60.7%, 58%.^{15,16}

Chorioamnionitis was present in 6(12%) of participants in group2 and 4(8%) in group 1. Overall incidence of chorioamnionitis in the study population was 20%. In present study it was found that higher chorioamnionitis was found in group 2 when compared to group 1. Ekin A et al found that AFI < 5 cm demonstrated a significantly a

higher rate of clinical chorioamnionitis.¹⁷ Yu et al reported that 17.8% of pregnancies and Jiaswal AA et al reported clinical chorioamnionitis (11.9%).^{18,19} Majority of the study participants required caesarean section. In Group 1 60% and in group 2, 76% required caserean section for delivery of the baby. Overall, 70% had caesarean section in present study. In a study by Mohakar et al reported 65% patients had vaginal delivery and 25% required LSCS which is lower than present study findings.¹²

NICU admission was required in 18% of the neonates in Group 1 whereas 48% of neonates required NICU admission in Group 2. Totally in both groups the NICU admission was 66%. Higher NICU admission in group 2 was found to be statistically significant. This is similar to previous study finding by Yu et al which reported 72.9% NICU admission.¹⁸ RDS developed in 12 % and 8% of neonates in Group 1 and Group 2 respectively. Over all 10% had RDS in present study. In a study by Patil S et al twenty six percent newborn suffered from respiratory distress syndrome.²⁰ RDS is not significantly associated with two groups. Likewise, in a study by Khanal S et al which reported that Respiratory system related problems like birth asphyxia, respiratory distress syndrome, apnea and pneumonia were common in both group but not statistically significant ($p>0.05$).²¹

CONCLUSION

PPROM is associated with increased maternal and perinatal complications especially when AFI<5. Caserean section rates, NICU admission, low mean APGAR scores and neonatal sepsis were found to be higher in PPRM with low amniotic fluid. Identification of risk factors, prompt treatment of infections and antibiotics following high vaginal swab culture might reduce the complications associated with PPRM. Prevention of risk factors and prediction of these morbidities are important in the management of PPRM.

Funding: No funding sources

Conflict of interest: None declared

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

REFERENCES

1. Beckmann, Charles. *Obstetrics and Gynecology*, 7e. Philadelphia: Wolters Kluwer Health/Lippincott Williams and Wilkins. pp. Chapter 17: Premature Rupture of Membranes. 2014;169-173.
2. Meis PJ, Ernest JM, Moore ML. Causes of low birth weight births in public and private patients. *Am J Obstet Gynecol.* 1987;156(5):1165-8.
3. Al-Riyami N, Al-Shezawi F, Al-Ruheili I, Al-Dughaiishi T, Al-Khabori M. Perinatal Outcome in Pregnancies with Extreme Preterm Premature Rupture of Membranes (Mid-Trimester PROM). *Sultan Qaboos Univ Med J.* 2013;13(1):51-6.
4. Bendon RW, Faye-Petersen O, Pavlova Z, Qureshi F, Mercer B, Miodovnik M, et al. Fetal membrane histology in preterm premature rupture of membranes: comparison to controls, and between antibiotic and placebo treatment. *Pediatr Dev Pathol.* 1999;2(6):552-8.
5. Mercer B, Milluzzi C, Collin M. Periviable birth at 20 to 26 wk of gestation: proximate causes, previous obstetric history and recurrence risk. *Am J Obstet Gynecol.* 2005;193(3 Pt 2):1175-80.
6. Gonan R, Hannah ME, Milligan JE. Does prolonged premature rupture of the membranes predispose to abruption placenta? *Obstet Gynecol.* 1989;74(3):347-50.
7. Ananth CC, Oyelese Y, Srinivas N, Yeo L, Vintzileos AM. Preterm premature rupture of membranes, intrauterine infection, and oligohydramnios: risk factors for placental abruption. *Obstet Gynecol.* 2004;104(1):71-7.
8. Mercer BM. Preterm premature rupture of the membranes. *Obstet Gynecol.* 2003;101(1):178-93.
9. Vintzileos AM, Campbell WA, Nochimson DJ. Degree oligohydramnios and pregnancy outcome in patients with PROM. *Obstet Gynecol.* 1985;66(2):162-7.
10. Mercer BM, Rabello YA, Thurnau GR. The NICHD-MFMU antibiotic treatment of preterm PROM study: Impact of initial amniotic fluid volume on pregnancy outcome: NICHD- MFMU Network. *Am J Obstet Gynecol.* 2006;194(2):438-45.
11. Lewis DF, Robichaux AG, Jaekle RK. Expectant management of preterm premature rupture of membranes and non-vertex presentation: what are the risks? *Am J Obstet Gynecol.* 2007;196(6): 566e1-6.
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):285-8.
13. Hargar 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 multicentric case control study. *Am J Obstet Gynecol.* 1990;163(1):130-7.
14. Gomez R, Romero R, Edwin SS, David C. Pathogenesis of preterm labor and preterm premature rupture of membranes associated with intrauterine infection. *Infect Dis Clinic North Am.* 1997;11(1):135-76.
15. Gandhi M, Shah F, Panchal C. Obstetric Outcomes In Premature Rupture Of The Membrane (Prom). *Inter J Gynecol Obstet.* 2012;16(2):1-5.
16. Kumari BR, Sailaja C, Usha P. Foetomaternal outcome in cases of premature rupture of membrane (prom) at term: an experience in our institute. *J. Evolution Med Dent Sci.* 2016;5(64):4508-11.
17. Souza AS, Patriota AF, Guerra GV, de Melo BC, Santos AC, Torres junior AC. Amniotic fluid volume and maternal outcomes in women with preterm premature rupture of membranes. *Rev Bras Gynecol Obstet.* 2014;36(4):146-51.

18. 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.
19. Jaiswal AA, Hariharan C, Dewani DKC. Study of maternal and fetal outcomes in premature rupture of membrane in central rural india. *Int J Reprod Contracept Obstet Gynecol*. 2017;6(4):1409-12.
20. Patil S, Patil V. Maternal and Foetal Outcome in Premature Rupture of Membranes. *IOSR J Dent Med Sci*. 2014;13(12):56-83.
21. Khanal S, Zhang W, Rajbhandari Shrestha N, Dahal GR. A comparative study of outcome of preterm neonate with and without history of preterm premature rupture of membrane. *Nepal Med Coll J*. 2009;11(2):99-103.

Cite this article as: Britto JJ, Sailatha R, Nalini AP. Role of amniotic fluid index on maternal and neonatal outcomes among obstetric women with preterm premature rupture of membranes. *Int J Reprod Contracept Obstet Gynecol* 2018;7:4171-6.