Original Article

# **Early Neonatal Morbidities in Late Preterm Neonates**

Neonatal Morbidities

Nasir Khan<sup>1</sup>, Khyal Muhammad<sup>2</sup>, Fiaz Ahmed<sup>3</sup>, Zaheer Abbas<sup>2</sup>, Rifayat Ullah Afridi<sup>4</sup> and Ejaz Hussain<sup>2</sup>

# **ABSTRACT**

**Objective:** To determine the distribution of early neonatal morbidities in late preterm infants.

Study Design: Descriptive study

**Place and Duration of Study:** This study was conducted at the department of Pediatrics and Neonatology Ayub teaching hospital, Abbottabad from May 2018 to December 2019.

**Materials and Methods:** After taking approval from ethical committee, data was collected from all neonates admitted to department of neonatology, who were born late preterm with gestation of less than 37 weeks but with 34 completed weeks. Total 147 neonates were included in this study. In all neonates who were included in this study, morbidities were evaluated from birth till 7<sup>th</sup> day of life through clinical examination and relevant investigations and were recorded on proforma.

**Results:** Mean neonatal age was 4 days with SD  $\pm$  3.74. Fifty-six percent neonates were male and 44% neonates were female. More over 25% neonates had hyperbilirubinemia, 28% neonates had sepsis, 20% neonates had intrauterine growth restriction, 4% neonates had transient tachypnea of newborn, 15% neonates had hypoglycemia, 16% neonates had respiratory distress syndromes and 13% neonates had apnea.

**Conclusion:** Our study concludes neonatal morbidities like hyperbilirubinemia, sepsis, intrauterine growth restriction, transient tachypnea of newborn; hypoglycemia, respiratory distress syndromes, and apnea are associated with late preterm births.

**Key Words:** Early neonatal morbidities, late preterm, infants

Citation of article: Khan N, Muhammad K, Ahmed F, Abbas Z, Afridi RU, Hussain E. Early Neonatal Morbidities in Late Preterm Neonates. Med Forum 2020;31(10):75-78.

### INTRODUCTION

Preterm delivery is one of the most significant cause of neonatal morbidity and mortality. Globally preterm deliveries are occurring due to various medical and obstetrical conditions mostly occurring in the late preterm period that results in morbidities of the new born. Late preterm neonates, born with period of gestation less than 37 weeks but 34 completed weeks are considered normal newborns and are kept in well-infant nursery units under the similar protocols as that of the term infants and sent home before sufficient observational period. Late preterm babies are not physiologically as full-grown as term babies so should not be considered functionally term in any aspect.

- <sup>1.</sup> Women Medical College Abbottabad.
- <sup>2</sup> Ayub Teaching Hospital Abbottabad.
- <sup>3.</sup> Women & Children Hospital
- <sup>4.</sup> Department of Pediatrics, Nasseer Teaching Hospital, Peshawar.

Correspondence: Dr. Nasir Khan, Assistant Professor Women Medical College, Abbottabad.

Contact No: 0311-5529571

Email: drnasirkhan1234@gmail.com

Received: April, 2020 Accepted: July, 2020 Printed: October, 2020 Late preterm infants are at high threat of morbidity and endangered outcome. These babies are at notably high short and long term unfavorable outcomes compare to term babies with a list of neonatal problems documented in literature. Some of these problems include feeding difficulties, hypoglycemia, respiratory distress syndromes (RDS), intrauterine growth retardation, sepsis, apnea, jaundice (hyperbilirubinemia) and transient tachypnea of the newborn. In a study in Pakistan by Haroon et al. He Respiratory distress syndrome was documented as 16.5%, High level of bilirubin of 37.9%, Hypoglycemia was reported about 5.2%, Growth retardation was 24.8%, Sepsis was documented about 4.9%, Transient tachypnea of newborn was reported in 7.0%, Apnea was documented about 15.3 in a Pakistani.

As the late preterm group is associated with greater morbidity compare to term neonates so this study is undertaken to identify the early neonatal morbidities in later preterm babies. Prior awareness of the morbidities associated with late preterm bodies is helpful for the health care provides to anticipate and manage potential complications in late preterm infants. Accurate estimate of the risks of morbidities is required to enable healthcare provider to take timely measures to improve the outcome.

#### MATERIALS AND METHODS

After taking hospital ethics committee approval this descriptive study was conducted at Pediatric

Department, Ayub Teaching Hospital Abbottabad from May 2018 to December 2019. Sample size of 147 was calculated using previous study. 11 Sampling technique applied was Consecutive non-probability sampling. All the late preterm (34 to 36 weeks of gestation) infants of both genders and of age up to 7 days admitted to neonatology unit of Ayub Teaching Hospital, Abbottabad were included while the term infants, infants with congenital anomies, syndromes, early preterm, multiple births and surgical conditions were excluded. Pretest counseling was given to parents. After written consent from the parents, those neonates fulfilling the above mentioned criteria were assessed for gestational age by menstrual period. In every baby who had require admission to neonates unit from birth to first 7 days of life, morbidities such as respiratory distress syndrome, hypoglycemia, sepsis, transient high respiratory rate of newborn, apnea and jaundice were evaluated. Infants were evaluated daily till 7 days of life through clinical examination or investigation for development of any of the morbidities mentioned above. Any of the predefined medical conditions resulting in post-delivery inpatient hospital observation and admission were assessed by physical examination as well as through relevant investigations. All the observations were done under supervision of an experience pediatrician.

All the above mentioned information including name, age, gender and address were recorded on a predesigned proforma.

Data was analyzed using SPSS version 21. Quantitative variables like age, gestation (weeks), weight were described

in terms of means+ standard deviation. Categorical data like gender and early neonatal morbidities (hyperbilirubinemia, sepsis, intrauterine transient tachypnea restriction, of newborn, hypoglycemia, respiratory distress syndrome and apnea) were described in the terms of frequency and percentages. All results were presented as tables and diagrams. Data was stratified by gender, age, gestation (weeks) & weight in term of neonatal morbidities. Post stratification chi -square test was used at 5% level of significance.

#### **RESULTS**

Table No 1. Neonatal morbidity (n= 147)

Morbidity	Frequency	Percentage
Hyperbilirubinemia	37	25%
Sepsis	41	28%
Intrauterine growth	29	20%
Restriction		
Transient tachypnea of	6	4%
Newborn		
Hypoglycemia	22	15%
Respiratory distress	24	16%
Syndrome		
Apnea	19	13%

Table No 2. Stratification of neonatal morbidity w.r.t age distribution

w.i.t age uist	w.r.t age distribution							
Morbidity	Status	1-4	4-7 days	Total	P value			
		days						
Hyperbili-	Yes	26	11	37				
rubinemia	No	75	35	110	0.8127			
Total		101	46	147				
Sepsis	Yes	28	13	41				
	No	73	33	106	0.9462			
Total		101	46	147				
Intrauterine	Yes	20	9	29				
growth	No	81	37	118	0.9733			
restriction								
Total		101	46	147				
Transient	Yes	4	2	6				
tachypnea of	No	97	44	141	0.9123			
newborn								
Total		101	46	147				
Hypoglycem	Yes	15	7	22				
ia	No	86	39	125	0.9540			
Total		101	46	147				
Respiratory	Yes	17	7	24				
distress	No	84	39	123	0.8060			
syndrome								
Total		101	46	147				
Apnea	Yes	13	6	19				
	No	88	40	128	0.9770			
Total		101	46	147				

Table No. 3: Stratification of neonatal morbidity w.r.t. gender distribution

gender distribution					
Morbidity	Status	Male	Female	Total	P value
Hyperbilirubinemia	Yes	21	16	37	
	No	61	49	110	0.8903
Total		82	65	147	
Sepsis	Yes	23	18	41	
	No	59	47	106	0.9618
Total		82	65	147	
Intrauterine growth	Yes	16	13	29	
restriction	No	66	52	118	0.9412
Total		82	65	147	
Transient tachypnea	Yes	3	3	6	
of newborn	No	79	62	141	0.7709
Total		82	65	147	
Hypoglycemia	Yes	12	10	22	
	No	70	55	125	0.8992
Total		82	65	147	
Respiratory distress	Yes	13	11	24	
syndrome	No	69	54	123	0.8617
Total		82	65	147	
Apnea	Yes	11	8	19	
_	No	71	57	128	0.8425
Total		82	65	147	

In this study age distribution among 147 neonates was analyzed as 101(69%) neonates were in age range 1-4 days, 46(31%) neonates were in age range 4-7 days. Mean age was 4 days with SD  $\pm$  3.74 Gender distribution among 147 neonates was analyzed as 82(56%) neonates were male and 65(44%) neonates were female. Gestational weeks among 147 neonates were analyzed as 56(38%) neonates had 35 weeks of gestation while 91(62%) neonates had 36 weeks of

gestation. Mean Gestational weeks was 36 weeks with SD  $\pm$  2.341 Weight distribution among 147 neonates was analyzed as 26(18%) neonates had weight <1.5 kg while 106(72%) neonates had weight range 1.5-2.5 Kg. Mean weight was 1.7 kg with SD  $\pm$  1.116.

Table No. 4: Stratification of neonatal morbidity w.r.t gestational week

gestational week					
Morbidity	Status	35	36	Total	P value
		weeks	weeks		
Hyperbilirubinemia	Yes	14	23	37	0.9703
	No	42	68	110	
Total		56	91	147	
Sepsis	Yes	16	25	41	
	No	40	66	106	0.8853
Total		56	91	147	
Intrauterine growth	Yes	11	18	29	
restriction	No	45	73	118	0.9838
Total		56	91	147	
Transient tachypnea of	Yes	2	4	6	
newborn	No	54	87	141	0.8063
Total		56	91	147	
Hypoglycemia	Yes	8	14	22	
	No	48	77	125	0.8561
Total		56	91	147	
Respiratory distress	Yes	9	15	24	
syndrome	No	47	76	123	0.9477
Total		56	91	147	
Apnea	Yes	7	12	19	
_	No	49	79	128	0.9041
Total		56	91	147	

Table No. 5. Stratification of neonatal morbidity w.r.t weight

Morbidity	Status	35	36	Total	P
-		weeks	weeks		value
Hyperbili-	Yes	14	23	37	0.9703
rubinemia					
	No	42	68	110	
Total		56	91	147	
Sepsis	Yes	16	25	41	
	No	40	66	106	0.8853
Total		56	91	147	
Intrauterine	Yes	11	18	29	
growth	No	45	73	118	0.9838
restriction					
Total		56	91	147	
Transient	Yes	2	4	6	
tachypnea of	No	54	87	141	0.8063
newborn					
Total		56	91	147	
Hypoglycemia	Yes	8	14	22	
	No	48	77	125	0.8561
Total		56	91	147	
Respiratory	Yes	9	15	24	
distress	No	47	76	123	0.9477
syndrome					
Total		56	91	147	
Apnea	Yes	7	12	19	
	No	49	79	128	0.9041
Total		56	91	147	

Neonatal morbidity among 147 neonates was analyzed as 37(25%) neonates had hyperbilirubinemia, 41(28%) neonates had sepsis, 29(20%) neonates had intrauterine growth restriction, 6(4%) neonates had transient tachypnea of newborn, 22(15%) neonates had hypoglycemia, and 24 (16%) neonates had respiratory distress syndromes while 19 (13%) neonates had Apnea. (Table 1).

# **DISCUSSION**

There has been a concomitant rise in the rate of morbidities among newborn delivered as preterm gestation.

Our study showed that among 147 neonates 69% neonates were in age range 1-4 days, 31% neonates were in age range 4-7 days. Mean neonatal age was 4 days with SD±3.74. Fifty six percent neonates were male and 44% neonates were female. Thirty eight percent neonates had 35 weeks of gestation while 62% neonates had 36 weeks of gestation. Mean Gestational weeks was 36 weeks with SD  $\pm$  2.341. Eighteen percent neonates had weight <1.5 kg while 106(72%) neonates had weight range 1.5-2.5 Kg. Mean weight was 1.7 kg with SD  $\pm$  1.116. More over 25% neonates had hyperbilirubinemia, 28% neonates had sepsis, 20% neonates had intrauterine growth restriction, 4% neonates had transient tachypnea of newborn, 15% neonates had hypoglycemia, and 16% neonates had respiratory distress syndromes while 13% neonates had

Similar results were observed in a study conducted by Haroon A et al<sup>11</sup> in which Respiratory distress syndrome in

16.5%, Growth retardation in 24.8%, high level of bilirubin in 37.9%, sepsis was 4.9%, hypoglycemia in 5.2%, transient high respiratory rate in 7.0% and apnea in 15.3%.

In another study conducted by Femitha P et al<sup>12</sup> in which respiratory distress syndrome was 12.4%, hyperbilirubinemia was 28.7%, sepsis was 20.8%, and Hypoglycemia was 5.2%. while in a study at Brazil<sup>13</sup> the growth retardation was 26.1%, transient tachypnoea was 25.9% and apnoea was 6.3% while 30% sepsis, hypoglycemia in 10.3% and feeding difficulty in 15.8% late preterm neonates were recorded in Jordan.<sup>14</sup>

Tiwari et al<sup>15</sup> reported that among late preterm 13.06% developed respiratory distress 52.56% late preterm had jaundice, 10.99% episodes of hypoglycemia, Hypothermia occurred in 7.94% late preterm neonates, 4.24% late preterm experienced one or more episodes of apnea. 18.06% late preterm babies had feeding problems, 9.79% term babies had confirmed sepsis.

In a study conducted by Binarbasi P et al<sup>16</sup>, hypothermia was noted in 14.5% of late preterm neonates and feeding difficulty in 19.1% in late preterm. Ligginc GC et al<sup>17</sup> observed that incidence of

apnea in 6% late preterm babies. In another study<sup>18</sup> the incidence of sepsis in late preterm was 10.3%.

This variation may be due to climatic condition of study places, difference in cut off temperature for consideration of hypothermia or differences in timing of study.

# **CONCLUSION**

Our study concludes that the frequency of early neonatal morbidities like hyperbilirubinemia, sepsis, intrauterine growth restriction, transient tachypnea of newborn, hypoglycemia, respiratory distress syndromes, apnea are associated in late preterm infants. Prior awareness of the morbidities associated with late preterm bodies is helpful for the health care provides to anticipate and manage potential complications in late preterm infants. Accurate estimate of the risks of morbidities is required to enable healthcare provider to take timely measures to improve the outcome.

#### **Author's Contribution:**

Concept & Design of Study: Nasir Khan

Drafting: Khyal Muhammad,

Fiaz Ahmed

Data Analysis: Zaheer Abbas, Rifayat

Ullah Afridi, Ejaz

Hussain

Revisiting Critically: Nasir Khan, Khyal

Muhammad

Final Approval of version: Nasir Khan

**Conflict of Interest:** The study has no conflict of interest to declare by any author.

#### REFERENCES

- 1. American College of Obstetricians and Gynecologists. Practice bulletin no. 159: management of preterm labor. Obstet Gynecol 2016;127(1):e29-38.
- 2. Natarajan G, Shankaran S. Short and long term outcomes of moderate and late preterm infants. Am J Perinatol 2016;33:305-17.
- Williams JE, Pugh Y. the late preterm: A population at risk. Crit Care Nurs Clin North Am 2018;30:431-43.
- 4. Ko HS, Jang YR, Yun H, Wie JH, Choi SK, Park IY, et al. Late preterm infants, early term infants and Timing of elective deliveries; current status in

- a Korean medical center. J Matern Fetal Neonatal Med 2019;32:1267-74.
- 5. Horgan MJ. Management of the late preterm infants: Not quite ready for prime time. Pediatr Clin North Am 2015;62:439-51.
- 6. Vogel JP, Chawanpaiboon S, Moller AB, Watananirun K, Bonet M, Lumbiganon P. The global epidemiology of preterm birth. Best Pract Res Clin Obstet Gynaecol 2018;52:3-12.
- 7. Torchin H, Ancel PY, Jarreau PH, Goffinet F. Epidemiology of preterm birth: Prevalence, Recent Trends, Short and Long-Term outcomes. J Gynecol Obstet Biol Repord (Paris) 2015;44:723-31.
- 8. Huff K, Rose RS, Engle WA. Late preterm infants: Morbidities, mortalities and management recommendations. Pediatr Clin North Am 2019; 66:387-402.
- 9. Pike KC, Lucas JSA. Respiratory consequences of late preterm birth. Paediatr Respir Rev 2015; 16:182-8.
- 10. Dani C, Poggi C, Pratesi S. Bilirubin and oxidative stress in term and preterm infants. Free Radic Res 2019;53:2-7.
- Haroon A, Ali SR, Ahmed S, Maheen H. Short-term neonatal outcome in late preterm vs. term infants. J Coll Physicians Surg Pak 2014;24(1): 34-8
- 12. Femitha P, Bhat BV. Early neonatal outcome in late preterms. Ind J Pediatr 2012;79(8):1019-24.
- 13. de Araújo BF, Zatti H, Madi JM, Coelho MB, Olmi FB, Canabarro CT. Analysis of neonatal morbidity and mortality in late-preterm newborn infants. J Pediatr 2012;88(3):259-66.
- 14. Salah OA. Unfavorable outcomes associated with late preterm birth: observations from Jordan. J Pak Med Assoc 2011;61(8):769-72.
- 15. Tiwari SK, Kumar N, Kumar S, Prabha R. A study in late preterm babies for early neonatal outcome. Ind J Neonatal Med Res 2017;5(1): PO01-PO05.
- 16. Binarbasi P, Akin Y, Narter F, Telatar B, Polatglu E, Agzikuru. Mortality and morbidity in late–preterm Newborn. Turk Arch Ped 2013; 48:17-22.
- 17. Liggins GC, Howie RN. A controlled trial of antepartum glucocorticoid treatment for prevention of respiratory distress syndrome in premature infants. Pediatr 1972; 50:515-25.
- 18. Sahana, Adarsh E, Sunil, Rajnish, Sreekrishna. Short term outcome of late preterm. Int J of Med and Appl Sci 2014;3(1): 206-09.