

# Role of Some Factors in Distribution of Neonatal Jaundice in Al-Najaf Province, Iraq

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#### Abstract

The present study occurred to detected on the factors in distribution of neonatal jaundice. During the first week of postnatal life affecting almost two thirds of term newborns, jaundice occurs in 60% of term newborns and 80% of preterm newborns in the first week of birth. Methods: The present study was conducted from September 2017 to January 2018. It was included 106 neonatal infants age (at born -17 day) treated with neonatal jaundice, who were Neonatology department at Al-Zahra Teaching Hospital located in Al-Najaf, Iraq. Data were collected using the newborns' medical records and interview sessions with the mothers. The results of this study were showed of 106 neonates jaundice infants indicated to significant difference p-value is <0.05 of age groups (<24 Hr. n= 46, 1-8 day n= 42, 9-16 n= 18), of neonatal jaundice infants in parameters included TSB, Hb, weight, and gestation age. The percent of neonatal jaundice rate was highest in male 62(59%) from female 44(41%). Increased of percent of neonatal jaundice rate in Gestation Age group >36 week n = 64(60.40%) more than group =< 35 week n = 42(39.60%), a statistically significant differences p < 0.05 decreased in TSB, Hb, weight, and gestation age of weight group = < 1500g n=22(20.8%), more than groups1600-2500 g n=32 (30.2%), and >2600 g n=52(49.1%) when compared between them. ABO results: the high percent rate in blood group O maternal (41.5%) when compared with A (32.1%), B (22.65%), and AB(3.8%) maternal blood groups. **Conclusion:** ABO and age gestation was an important factors may be effects of development physiological neonatal jaundice into harmful pathological neonatal jaundice and hyperbilirubinemia.

### Introduction

Neonatal jaundice is a common phenomenon, during the first week of postnatal life affecting almost two thirds of term newborns, jaundice occurs in 60% of term and 80% of preterm newborns in the first week of birth. (Mateo *et al*, 2013; Christensen *et al*, 2013). The mechanism of neonatal hyperbilirubinemia is multifactorial, including processes that contribute to extensive bilirubin load, or diminished bilirubin clearance (Lin *et al.*, (2015). Many studies have been performed to identify factors affecting neonatal hyperbilirubinemia. Based on the findings, several risk factors have been introduced as the determinants of jaundice including maternal and neonatal risk factors such as the newborn's age, ethnicity, Rh blood group, maternal diseases, neonate's gender, birth weight, frequency of nutrition and defecation, birth trauma, and history of jaundice among siblings (Bhutani 2

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2016; Oyedeji *et al*, 2015; Esmailepour *et al*, 2007). Although both genetic and environmental factors may contribute to the development of neonatal hyperbilirubinemia, the importance of genetically determined conditions has been increasingly recognized (Lin *et al*, 2008). Bilirubin overproduction that occurs in ABO, Rhesus or minor blood group incompatibilities with a positive direct antiglobulin test has been recognized as one of the major risk factors for development of severe hyperbilirubinemia in infants of 35 or more weeks' gestation. Maternal-fetal ABO blood group incompatibility, in which the mother has blood group O and the newborn has blood group A or B, occurs in 15-20% of all pregnancies. The hemolytic process results from maternal anti-A or anti-B immunoglobulin G (IgG) antibodies crossing the placenta and attaching to the appropriate antigens on the neonatal red cells. Resultant heme catabolism causes an increased indirect bilirubin (IB) production, leading to neonatal jaundice (Kaplan *et al*, 2009).

#### **Materials and Methods:**

The present study was conducted from September 2017 to January 2018. It was included 106 neonatal infants age (at born -17) day, treated with neonatal jaundice, who were Neonatology department at Al-Zahra Teaching Hospital located in Al-Njaf, Iraq. Data were collected using the newborns' medical records and interview sessions with the mothers. A checklist consisting of demographic, neonatal, and maternal information was used for data collection, in this study, the relationship between the severity of jaundice and predisposing factors including Age, gender, Gestation Age, birth weight, and ABO. Infants were divided into groups according Age (<24 Hr. n = 46, 1-8 day n = 42, 9-16 n = 18), gender (Male n = 62and Female n = 44), Gestation Age (=< 35 week n = 42, >36 week n = 64), birth weight (=< 1500 n = 22, 1600-2500 n = 32, >2600 n = 52), maternal and neonatal ABO were classified into four groups: A (n=34,32); B n=(24,34); AB (n=4,6); and O (n=44,34) respectively. 5.2milliliters of blood was collected in a plain bottle from the femoral vein. The samples were tested in the Department of Hematology Biochemistry clinical and Blood Bank of laboratory Al-Zahra Teaching Hospital. The mother 's and neonate 's blood groups were estimated along with hemoglobin levels, WBC, and total serum bilirubin of the neonate. Criteria for inclusion were all neonates with the diagnosis of jaundice as well as G6PD tests and Rh testing of mother and newborn.

#### Statistical analyses :

were performed using (SPSS 24.0 Inc., Chicago, IL, USA). Categorical variables were presented with absolute numbers and percentages for each groups. Testing of significance between groups regarding the analyzed parameters was performed with: Student independed t-test, Chi-square, and ANOVA test. The result was considered significant (p < 0.05). 3

#### **Results :**

The results of this study were showed of 106 neonates jaundice infants. **Table (1)** indicated to a significant difference p-value is <0.05 of age groups (<24 Hr. (n=46),1-8 day (n=42), 9-16 (n=18), of neonatal jaundice infants in parameters included TSB, Hb, weight, and gestation age. Therefore, no a significant difference in same of these parameters when compare between male and female neonatal jaundice infants **table (2)**, but the percent of neonatal jaundice rate was highest in male 62(59%) from female 44(41%) figure (1). table (6)

The result statistically significant in the **table (3)** and **figure (2)** cleared increased of percent of neonatal jaundice rate in Gestation Age group >36 week n = 64(60.40%) more than group=< 35 week n = 42(39.60%), but the Gestation Age wasn't effected (no significant differences) of TSB and study parameters levels between two groups.



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table (4) and figure (3) showed a statistically significant differences p < 0.05 decreased in TSB, Hb, weight, and gestation age of weight group = < 1500 g, more than groups 1600-2500 g, and >2600 g when compared between them, thus highest percent of neonatal jaundice rate in group >2600 g n=52(49.1%), but in n=22(20.8%) and n=32 (30.2%) of groups =< 1500 g and 1600-2500 g respectively. table (6).

Not statistically significant of study parameters in ABO groups of neonatal infants and those mothers table (4), also the result in the figure (4) indicated to percent neonatal jaundice rate about A (30.2%), B(5.7%), AB(32.1%) and O(32.1%) blood groups of neonatal infants respectively, but the high percent rate in blood group O maternal 41.5% when compared with A (32.1%), B (22.65%), and AB(3.8%) maternal blood groups respectively.

table(6). Tabl	le (1) Descriptiv	e Statistic of T	<b>SB</b> level	l and	study para	meters according a	
	gr	oups in neona	tal jauno	dice iı	nfants		
Age	Parameters	Mean	Std. Error		Minimum	Maximum	
-	Statistics						
<24 Hr.	TSB (mg/dl)	11.435 *	.8189		1.8	15.0	
N= 46							
WBC (cell*103)	16.830	1.4203		8.0		29.3	
HB (mg/dl)	18.383 *	.6100		13.2		23.0	
Weight (g)	2313.04 *	214.500		600		4250	
Gestation Age	33.43 *	.873		27		40	
(week)							
1-8 day	TSB (mg/dl)	15.905 *	.9025		4.3	18.9	
N= 42							
WBC (cell*103)	16.105	.7888		7.4		26.0	
HB (mg/dl)	16.390 *	.8661		6.8		21.0	
Weight (g)	2604.76 *	202.541		1100		4000	
Gestation Age	35.19 *	.642		28		38	
(week)							

1.0095

7.6

9.3

12.3

2000

35

17.2

22.0

20.0

3750

37

17.611 \*

1.3585

175.682

.7662

.200

9-16 day

WBC (cell\*103)

HB (mg/dl)

Weight (g)

(week)

Gestation Age

N=18

TSB (mg/dl)

16.556

17.089

36.11 \*

\*significant difference at p <0.05 between groups

2944.44 \*

ge

Table (2) Descriptive Statistic of TSB level and study parameters according Gender groups in neonatal jaundice infants

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Gender	Parameters	Mean	Std. Er	ror	Minimum	Maximum	13510
	Statistics						
Male	TSB (mg/dl)	15.97	.7409		2.1	17.5	
N= 62							
WBC (cell*103)	16.377	.9549		8.2		29.3	
HB (mg/dl)	17.068	0.6522		6.8		22.9	
Weight (g)	2629.03	146.313	3	1000	1	4250	
Gestation Age	34.65	0.582		28		39	
(week)							
Female	TSB	14.01	0.9573		1.8	18.9	
N=44	(mg/dl)						
WBC (cell*103)	16.664	1.1124		7.4		26.3	
HB (mg/dl)	17.805	0.6370		9.2		23.0	
Weight (g)	2404.55	232.246	5	600		4000	
Gestation Age	34.50	0.813		27		40	
(week)							

 Table (3) Descriptive Statistic of TSB level and study parameters according Gestation Age groups in neonatal jaundice

Parameters	Mean	Std. Error	Minimum	Maximum
Statistics				
TSB	16.88	0.6739	3.2	15.0
(mg/dl)				
13.548	1.0129	7.4	4	26.3
17.757	0.7587	6.8	8	23.0
1702.38	149.024	60	0	3250
TSB	13.08	0.8080	1.8	18.9
(mg/dl)				
18.431*	0.8325	9.3	3	29.3
17.122*	0.5882	8.3	3	22.9
3082.81*	109.931	20	000	4250
	Parameters Statistics TSB (mg/dl) 13.548 17.757 1702.38 TSB (mg/dl) 18.431* 17.122* 3082.81*	ParametersMeanStatisticsTSBTSB13.5481.012917.7570.75871702.38149.024TSB13.08(mg/dl)18.431*0.832517.122*0.58823082.81*109.931	Parameters         Mean         Std. Error           Statistics         TSB         16.88         0.6739           (mg/dl)         13.548         1.0129         7.4           17.757         0.7587         6.3           1702.38         149.024         60           TSB         13.08         0.8080           (mg/dl)         18.431*         0.8325         9.3           17.122*         0.5882         8.3           3082.81*         109.931         20	Parameters StatisticsMeanStd. ErrorMinimumStatistics16.880.67393.2TSB16.880.67393.2(mg/dl)13.5481.01297.417.7570.75876.81702.38149.024600TSB13.080.80801.8(mg/dl)18.431*0.83259.317.122*0.58828.33082.81*109.9312000

\*significant difference at p <0.05 between groups



 Table (4) Descriptive Statistic of TSB level and study parameters according Weight groups in neonatal jaundice infants

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Weight (g)	Parameters	Mean	Std. Err	or	Minimum	Maximum 🔇 💕		
	Statistics							
=< 1500	TSB	13.75	0.8057		4.3	11.5		
N= 22	(mg/dl)							
WBC	13.709	1.7667		7.4		26.3		
(cell*103)								
HB (mg/dl)	18.545	0.7836		13.5		23.0		
Gestation Age	e 29.82	0.519		27		32		
(week)								
1600-2500	TSB	17.36 *	1.1487		2.4	17.2		
N= 32	(mg/dl)							
WBC	15.463 *	1.0044		8.4		26.0		
(cell*103)								
HB (mg/dl)	17.050	1.0147		6.8		21.0		
Gestation Age	e 34.50 *	0.806		27		38		
(week)								
> 2600	TSB	13.85 *	0.8121		1.8	18.9		
N= 52	(mg/dl)							
WBC	18.312 *	0.9978		9.0		29.3		
(cell*103)								
HB (mg/dl)	17.077	0.6279		9.2		22.9		
Gestation Age	e 36.65 *	0.337		30		40		
(week)								
*significant d	lifforance at n	<0.05 hotwoor	aroung					

\*significant difference at p <0.05 between groups

 Table (5) Descriptive Statistic of TSB level and study parameters according ABO groups in neonatal jaundice infants

J						
ABO	Parameters	Mean	Std. Eri	ror	Minimum	Maximum
	Statistics					
Α	TSB (mg/dl)	14.22	1.1461		1.8	18.9
N= 32						
)3WBC (cell*10	17.488	1.2386		8.0		26.0
HB (mg/dl)	18.088	.7862		9.2		22.7
Gestation Age	34.94	.854		27		40
(week)						
Weight (g)	2571.88	253.762	2	1000		4250
B	TSB (mg/dl)	14.85	1.0832		2.1	17.5
N=34						
)3WBC (cell*10	14.818	1.0642		8.2		26.0
HB (mg/dl)	16.871	.4835		12.3		20.0
Gestation Age	34.47	.879		28		38
(week)						

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	Print IS	SN: 2073-8854	Colline	ISSN: 2311-6544	8
Weight (g)	2764.71	198.189	1000	) 4000	35114
AB	TSB (mg/dl)	15.43	3.6762	4.6 17	.2
N= 6					
)3WBC (cell*10	13.333 *	3.1248	7.4	18.0	
HB (mg/dl)	18.633	.8762	17.0	20.0	
Gestation Age	34.33	1.202	32	36	
(week)					
Weight (g)	1916.67	363.242	1250	) 2500	
0	TSB (mg/dl)	15.4	.8843	4.3 17	.5
N= 34					
)30WBC (cell*1	17.800	1.4393	9.3	29.3	
HB (mg/dl)	16.982	1.1403	6.8	23.0	
Gestation Age	34.41	.908	27	39	
(week)					
Weight (g)	2382.35	243.177	600	3900	
*significant diff	erence at p <0	.05 between gr	oups		



Table (6) Chi- Square test and p-values for groups of study

statistics	Gender	Gestatio	Weight	ABO	ÅBO	Age
	groups	n groups	groups	neonatal	maternal	groups
Chi-	3.057	4.566	13.208	21.245	33.019	12.981
Square						
df	1	1	2	3	3	2
p-value	.080	.033	.001	.000	.000	.002

#### Dissection

Various studies show large differences in the prevalence of neonatal jaundice in the world. This difference is more pronounced, especially for many factors that causes physiological and pathological neonatal jaundice leading to hyperbilirubinemia in newborn (Christensen *et al*, 2013; Nepal *et al*, 2010). Study included 106 cases of Neonatal jaundice cases. Range age of the (> 24 hr. to 17 day), among them male 62(59%) while 44(41%) were female. Neonatal jaundice was more common in male infants as compared to female infants in two different studies done by Mantani *et al* (2007)(62% vs. 38%) and Sharma *et al* (2006)(1.3:1), because the number of erythrocytes in male more than in female; But The results of (Ivana *et al*, 2014)



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showed that the incidence of jaundice does not be influenced by on the gender and that it is so common in male and female neonates. In present study, percentage of Gestation Age group =<35 week (39.60%,), were mean of (29.82±0.519) week babies having low birth weight (<1500 g), and TSB (16.88±0.76) mg/dl, whereas in Gestation Age group >36 week (60.40%), were mean about (36.65±0.33) week babies having normal weight (>2600) week were mean about (3082.81±109.93)g, and TSB (13.08 ±0.80) mg/dl, these results are agreed with the study of (Lee *et al.*, 2016), and a research of (Ivana *et al.*, 2014) showed that the occurrence of jaundice in newborns with a low birth weight is linked to low gestational age. he wasThe shared of jaundiced newborns of gestational age of fewer than 37 weeks in the total number of infants of the same age is 56%. Lin *et al.*, (2015) the literature revealed to incident neonatal jaundice are caused by unconjugated bilirubin in the blood resulted of breakdown of RBC and elevated bilirubin formation and because the neonatal liver is unable to removal bilirubin rapidly enough from the blood lead to hyberbilirubinemia.

In our study the result indicated to not statistically significant of study parameters in ABO groups of neonatal infants and those mothers, but percent neonatal jaundice rate about A (30.2%), B(5.7%), AB(32.1%) and O(32.1%) blood groups of neonatal infants respectively, but the high percent rate in blood group O maternal 41.5% when compared with A (32.1%), B (22.65%), and AB(3.8%) maternal blood groups respectively. These agree with other findings that jaundice resulting from incompatibility usually present higher levels of serum bilirubin (Ella et al., 2013; Vijaya et al., 2018) The bilirubin levels was high in the samples of babies with blood group A or B born to mothers with blood group O with 27.1 mg/100ml for the baby with blood group B. The difference was however not statistically significant. (Ella *et al.*, 2013). Another research from India evaluating 151 neonates also showed no difference in severity between O-A and O-B for hemolytic disease of the newborn (Bhat et al, 2012). The reason for this difference may have been that "A" and "B" antigens are weaker antigens and the distance between a/b antigen sites on the fetal red cells as compared to adult red cells is more. It is a very reliable indicator. To conclude, Neonatal Jaundice in the prematurity and perinatal period is related with the gestational age, weight birth, most of the cases were having idiopathic jaundice although and ABO- were not exceptional. Also prematurity and low birth weight were having higher levels of bilirubin. In any infant, 24 hours old any jaundice is considered pathologic and requires estimation.

#### References

1. Bhat Y, Pavan Kumar C. Morbidity of ABO haemolytic disease in the newborn. Paediatr Int Child Health, 2012; 32: 93-96.

2. Bhutani VK, Srinivas S, Castillo Cuadrado ME, Aby JL, Wong RJ, Stevenson DK. Identification of neonatal haemolysis: an approach to predischarge management of neonatal hyperbilirubinemia. Acta Paediatr. 2016;105(5):e189-194

3. Christensen RD, Lambert DK, Henry E, Eggert LD, Yaish HM, Reading NS, Prchal JT: Unexplained extreme hyperbilirubinemia among neonates in a multihospital healthcare system.Blood Cells Mol Dis 2013; 50: 105–109.

4. Dennery PA, Seidman DS, Stevenson DK. Neonatal hyperbilirubinemia. N Engl J Med. 2001;344(8):581-90.

5. Dufour DR, Monoghan WP. ABO hemolytic disease of the newborn: a retrospective analysis of 254 cases.Am J Clin Pathol 2010; 73: 369-373.

6. Esmailepour-Zanjani S, Safavi M, Jalali S, Abyane EE. Incidence and associated factoes of neonatal hyperbilirubinemia at Hedayat hospital. J Shahid Beheshti Sch Nurs Midwifery. 2007; 17(59):19-25 (Persian).





## Al-Kufa University Journal for Biology / VOL.8 / NO.1 / Year: 2016 Print ISSN: 2073-8854 Online ISSN: 2311-6544

Manjareeka. (2013) J Basic Clin Physiol Pharmacol 2013; aop Mishra et al.: Hematological profile in neonatal jaundice

8. Kaplan M, Na'amad M, Kenan A, et al. Failure to predict hemolysis and hyperbilirubinemia by IgG subclass in blood group A or B infants born to group O mothers.Pediatrics 2009; 123:132-137.

9. Lin Z, Fontaine J and Watchko JF. Coexpression of gene polymorphisms involved in bilirubin production and metabolism. Pediatrics. 2008;122(1):e156-62.

10. Mantani M, Patel A, Renge R, Kulkarni H. Prognostic value of direct bilirubin in Neonatal Hyperbilirubinemia. Indian J Pediatr 2007; 79: 819-22.

11. Mateo PC, Lee KS, Barozzino M, Sgro M. Management of neonatal jaundice varies by practitioner type. Can Fam Physician. 2013; 59(8):e379-86.

12. Nepal D, Banstola D, Dhakal AK ,Mishra U, Mahaseth C Clinico-Laboratory Profile and Immediate Outcomes of Hyperbilirubinemic Babies Admitted in Kanti Children Hospital Journal of Nepal Paediatric Society; January-June, 2010; 1 (30).

13. Oyedeji OA, Adeyemo TA, Ogbenna AA, Akanmu AS. Prevalence of anti-A and anti-B hemolysis among blood group O donors in Lagos. Niger J Clin Pract 2015;18(3):328-332.

14. Sharma P, Chhangani NP, Meena KR, Jora R, Sharma N, Gupta BD. Brainstem Evoked Response Audiometry (BAER) in neonates with hyperbilirubinemia. Indian J Pediatr2006; 73: 413-16.

15. Mesi I, Milas V, Meimurec M, and Rimar E. Unconjugated Pathological Jaundice in Newborns. Jaundice in Newborns, Coll. Antropol. 38 (2014) 1: 173–178.

16. Lin CH, Yang HC, Cheng CH, and Y CE. Effects of infant massage on jaundiced neonates undergoing phototherapy Italian Journal of Pediatrics (2015) 41:94

17. Lee BK, Le Ray I, Sun JY, Wikman A, Reilly M, Johansson S. Haemolytic and nonhaemolytic neonatal jaundice have different risk factor profiles. Acta Paediatr. 2016; 105: 1444–50.

18. Ella EE, Garba SA, Ogal WN. ABO Incompatibility and its role in neonatal jaundice in Zaria, Kaduna state of Nigeria. IJES. 2013;2(11):17-23.

19. Vijaya S. Kattimani, Ushakiran C. B.. Hemolytic disease of the new-born due to ABO incompatibility.IJCP. 2018, 5(2).

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