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Evaluation of neonatal jaundice in the Makkah region

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The aims of this study were to detect the frequency at which the different types of neonatal jaundice occur in Makkah and to estimate the malondialdehyde (MDA) levels. This study included 239 neonates with neonatal jaundice, 20 anemic neonates and 21 healthy neonates. ABO incompatibility was observed in 31.6% of neonates with indirect hyperbilirubinemia, in 14.3% of those with early onset jaundice, in 9.5% of those with persistent jaundice, in 8.5% of those with physiological jaundice, in 5% of anemic neonates and in 12% of all neonates. glucose-6-phosphate dehydrogenase (G6PD) deficiency was observed in 10.5% of neonates with indirect hyperbilirubinemia, in 3.9% of those with physiological jaundice, in 11.1% of those with direct hyperbilirubinemia, in 12% of those with persistent jaundice, in 10% of anemic neonates and in 6.6% of all neonates. Rh incompatibility and polycythemia were found in 2.6% of neonates with indirect hyperbilirubinemia and in 0.4% of all neonates. In comparison to control group, MDA was significantly higher in all groups except for the anemic group. In conclusion, ABO incompatibility and G6PD deficiency frequently result in neonatal jaundice in Makkah, whereas Rh incompatibility and polycythemia are rare. The MDA level may serve as an indicator of oxidative stress.

Neonatal jaundice is a benign condition that often does not require intervention. Most cases of neonatal hyperbilirubinemia constitute physiological jaundice and do not have serious consequences. An important clinical challenge is to identify those newborns who are at risk of developing severe neonatal hyperbilirubinemia¹. There are many factors implicated in the development of pathological jaundice, including perinatal factors (e.g., birth trauma or infections), maternal factors (e.g., Rh or ABO incompatibility), neonatal factors (e.g., prematurity or polycythemia), and genetic factors (e.g., Crigler-Najjar's or Gilbert's syndrome)². In addition, the administration of drugs such as cephalosporins and glucose-6-phosphate dehydrogenase (G6PD) enzyme deficiency have been implicated in pathological jaundice². The mechanism by which hyperbilirubinemia occurs is through either increased bilirubin production (resulting from hemolysis, sepsis, blood extravasation or polycythemia) or increased enterohepatic circulation (resulting from prematurity, pyloric stenosis, delayed bacterial gut colonization, gastrointestinal tract immobility or obstruction), or decreased bilirubin elimination which occurs in Crigler-Najjar's and Gilbert's syndromes².

ABO incompatibility usually occurs in the offspring of women with type O blood and occasionally in mothers with both type A blood and high anti-B IgG levels³. ABO incompatibility exists in approximately 15–25% of all maternal/fetal pairs. However, ABO hemolytic disease of the newborn occurs in approximately 1% of group O mothers with high antenatal IgG antibody titers⁴. Currently, the use of prophylactic anti-D immunoglobulin greatly reduces the incidence of hemolytic disease of the newborn due to Rh incompatibility⁵.

G6PD deficiency is an X-linked recessive disease that results in clinical manifestations such as neonatal jaundice, chronic nonspherocytic anemia, infection and drug-induced hemolysis⁶. A role for G6PD deficiency in the genesis of neonatal hyperbilirubinemia has been reported⁷. The most serious complication of G6PD deficiency in newborns is kernicterus resulting from severe neonatal hyperbilirubinemia.

Infants who experience chronic or acute fetal hypoxia have a higher incidence of polycythemia, with hematocrit levels greater than 65%. Neonatal hyperbilirubinemia is a common clinical feature⁸.

Hyperbilirubinemia can also result from increased oxidative stress. It has been reported that bilirubin can function as a scavenger of reactive oxygen species⁹. Malondialdehyde (MDA) is a reactive metabolic product that results from the effects of reactive oxygen species on tissues and from a series of reactions that occur during lipid peroxidation¹⁰. Plasma MDA levels are a sensitive indicator of lipid peroxidation and thus of oxidative stress^{11,12}. Therefore, measuring the MDA concentration in hyperbilirubinemic newborns will help determine the relationship between oxidative stress and elevated bilirubin levels. The aim of this study was to determine the frequency at

which the different types of neonatal jaundice occur in the Makkah region. We also sought to determine the prevalence of ABO incompatibility, Rh incompatibility, G6PD deficiency, and polycythemia in neonates with different types of jaundice. Finally, we sought to estimate the levels of MDA and correlate them with the various parameters studied.

Results

This study included 239 jaundiced neonates, 20 anemic babies, and 21 healthy neonates (control group). The neonates consisted of 169 males and 111 females (a ratio of 3 male : 2 female). The numbers of neonates in each group was G1, $n = 21$, G2, $n = 38$, G3, $n = 129$, G4, $n = 9$, G5, $n = 42$, G6, $n = 21$, and G7, $n = 20$ neonates (Table 1). The total number of neonates with hyperbilirubinemia and anemia was 259. Of these neonates, 74 were preterm. The members of groups 2, 3, and 6 were significantly younger ($p < 0.001$) than those in the control group (G1). In contrast, those in group 7 were significantly older than those in G1 (Table 2). Six neonates (15.8%) were preterm in G2, 37 (28.7%) were preterm in G3, 1 (11.1%) was preterm in G4, 15 (35.7%) were preterm in G5, 9 (42.9%) were preterm in G6 and 6 (30%) were preterm in G7. No significant differences were found with regard to age among preterm and term infants in any of the groups ($p > 0.05$).

Several parameters were evaluated for each group. The results are summarized in Table 2. Comparisons between the different groups were performed using the Kruskal-Wallis test. We found highly significant differences between the groups in age, total bilirubin levels, direct bilirubin levels, G6PD activity, red blood cell counts and hemoglobin concentrations ($p < 0.001$). No significant differences in MDA levels were found ($p > 0.05$).

As shown in Figure 1, the frequencies of neonatal jaundice are as follows: physiological jaundice-53.9% (129/239), persistent jaundice-17.6% (42/239), indirect hyperbilirubinemia-15.9% (38/239), early onset jaundice-8.8% (21/239), and direct hyperbilirubinemia-3.8% (9/239).

The highest prevalence of ABO incompatibility was found in the indirect hyperbilirubinemia group (12/38, 31.6%) followed by the early onset jaundice group (3/21, 14.3%), the persistent jaundice group (4/42, 9.5%), the physiological jaundice group (11/129, 8.5%) and the anemic group (1/20, 5%). The frequency of ABO incompatibility in all studied groups was 31/259 (12%) (Fig. 2). ABO incompatibility was determined by clinical classification of the mother as type O and the neonate as type A or B.

Regarding hemoglobin concentrations in neonates with ABO incompatibility, no significant differences were observed between the control group and the indirect hyperbilirubinemia group, the physiological jaundice group, or the early onset jaundice group. However, the persistent jaundice group had significantly decreased hemoglobin levels than the control group (11.0 ± 1.4 , $p = 0.01$).

The frequency of Rh incompatibility was 1/38 (2.6%) in the indirect hyperbilirubinemia group. Rh incompatibility occurred in 0.4% of all neonates. The frequency of polycythemia was 2.6% (1/38) in the indirect hyperbilirubinemia group and 0.4% of all neonates. The direct Coombs' test was used in all cases. A positive result was found in only 5 cases of ABO incompatibility (16.1%) and in only one case of Rh incompatibility.

The frequency of G6PD enzyme deficiency among the different groups is shown in Table 3. The classification of G6PD deficiency was performed according to the WHO¹⁷. No cases of class I G6PD deficiency were found. However, 10 cases of class V were observed. Of these, 2 were in G2 (one male and one female), 6 were in G3 (five males and 1 female), 2 were in G5 (both female), and 2 were in G6 (both female). No significant differences were found between males and females regarding G6PD concentrations, with the exception of G6, the early onset of jaundice, which showed significantly higher G6PD concentrations in the female group than in the male group ($p > 0.05$ and $p < 0.05$, respectively) (Table 4). There was no significant difference in the concentrations of hemoglobin between the neonates with G6PD deficiency and the control group. The hemoglobin concentrations in each group were 13.7 ± 3.1 and 15.3 ± 2.0 , respectively.

Using the Spearman correlation, a significant negative correlation was found between MDA and hemoglobin levels in the persistent jaundice group ($r = -0.98$, $p = 0.01$). No other significant correlations were detected.

Discussion

Neonatal hyperbilirubinemia is associated with a variety of conditions. Severe NH poses a direct risk of permanent neurological sequelae. The early identification of neonates who are at a greater risk of developing severe neonatal hyperbilirubinemia is of paramount importance to preventing brain damage¹⁷.

In this study, the type of jaundice found at the highest frequency was physiological jaundice (53.9%) followed by persistent jaundice (17.6%), indirect hyperbilirubinemia, early onset jaundice and direct hyperbilirubinemia (Fig. 1). These findings are in line with various national and international studies^{18,19}. Physiological aspects that can contribute to neonatal hyperbilirubinemia include increased bilirubin production, less efficient hepatic conjugation, and enhanced bilirubin absorption by the enterohepatic circulation²⁰. Bilirubin has been proposed as an effective antioxidant, and modest elevations of bilirubin may be beneficial in neonates²¹.

ABO incompatibility and G6PD deficiency were found to be frequent causes of neonatal hyperbilirubinemia in our study, as well as in other studies. G6PD deficiency is the most common human genetic enzymopathy, affecting over 200 million individuals worldwide. It is closely associated with neonatal jaundice, chronic nonspherocytic hemolytic anemia, favism and food- or drug-induced acute hemolytic anemia²². In this study, class II G6PD enzyme deficiency

Table 1 | Classification of the neonates

Groups	Name	n	Male	Female	Criteria
G1	Normal neonates	21	13	8	Normal newborn without jaundice or anemia.
G2	Indirect hyperbilirubinemia	38	15	23	$\geq$ indirect bilirubin 5 mg/dl in 0 h, $\geq$ 10 mg/dl in 24–48 h of age, and $\geq$ 13 mg/dl in $\geq$ 48 h of age.
G3	Physiological jaundice	129	75	54	Bilirubin high but does not reach the cutoff for indirect hyperbilirubinemia in relation to the age of the subjects.
G4	Direct hyperbilirubinemia	9	7	2	Direct bilirubin $>$ 2 mg/dl ($>$ 20% of total bilirubin).
G5	Persistent jaundice	42	27	15	Persistent high bilirubin $>$ 2 w in full term baby; $>$ 3 w in preterm baby.
G6	Early onset jaundice	21	16	5	Appearance of jaundice at 0 to 24 h of age with indirect bilirubin $<$ 5 mg/dl.
G7	Anemic group	20	16	4	Anemia without jaundice.
	Total number	280	169	111

Table 2 | A comparison of the studied clinical and laboratory parameters in all groups

Groups studied	Mean $\pm$ SD	Median (range)	Age days	Total bilirubin mg/dl	Direct bilirubin mg/dl	Indirect bilirubin mg/dl	Malondialdehyde IU/l	Red blood cells $\times 10^{12}/l$	Hemoglobin g/dl
G1	Mean $\pm$ SD	Median (range)	17.4 $\pm$ 11.1	0.7 $\pm$ 0.26	0.1 $\pm$ 0.08	0.6 $\pm$ 0.2	4.0 $\pm$ 1.0	4.8 $\pm$ 0.7	15.5 $\pm$ 2.2
G2	Mean $\pm$ SD	Median (range)	21 (0–30)	0.85 (0.2–1.0)	0.1 (0.0–0.3)	0.7 (0.2–0.8)	3.7 (3.0–7.5)	4.8 (4.0–6.2)	15 (13–20)
G3	Mean $\pm$ SD	Median (range)	5.1 $\pm$ 3.9	12.6 $\pm$ 2.9	0.7 $\pm$ 1.6	12.0 $\pm$ 2.7	8.0 $\pm$ 4.5	4.8 $\pm$ 0.8	15.8 $\pm$ 3.1
G4	Mean $\pm$ SD	Median (range)	4.5 (0–14)***	13.0 (5–16)***	0.4 (0.1–10.1)	12.6 (4.8–15.7)***	6.0 (3.3–18.3)***	4.8 (3.3–6.2)	16.3 (10.4–21.6)
G5	Mean $\pm$ SD	Median (range)	5.9 $\pm$ 3.5	7.2 $\pm$ 2.9	0.4 $\pm$ 0.3	6.2 $\pm$ 2.9	7.8 $\pm$ 5.1	4.1 $\pm$ 0.8	14.7 $\pm$ 2.8
G6	Mean $\pm$ SD	Median (range)	5 (2–14)***	7.3 (1.1–12.5)***	0.3 (0.1–2.2)***	6.4 (0.4–11.8)***	6.7 (2.3–28)**	4.2 (2.5–6.2)**	14.7 (8.7–20.9)
G7	Mean $\pm$ SD	Median (range)	24.3 $\pm$ 6.7	6.9 $\pm$ 6.3	4.9 $\pm$ 3.4	2.0 $\pm$ 2.96	10.9 $\pm$ 2.9	3.8 $\pm$ 0.9	10.0 $\pm$ 2.4
G8	Mean $\pm$ SD	Median (range)	28 (14–30)	5.0 (3.0–3.0)**	3.9 (2.0–13.6)***	0.9 (0.26–9.4)**	11.0 (8–17)***	4.2 (2.5–4.7)**	10.0 (7.0–13.0)***
G9	Mean $\pm$ SD	Median (range)	20.8 $\pm$ 4.6	3.5 $\pm$ 2.7	0.7 $\pm$ 0.5	2.9 $\pm$ 2.8	9.0 $\pm$ 5.5	3.9 $\pm$ 0.7	11.9 $\pm$ 2.0
G10	Mean $\pm$ SD	Median (range)	21 (14–30)	2.5 (0.0–11.0)***	0.6 (0.1–2.0)***	1.8 (0–10.3)***	7.6 (3.5–19)**	3.8 (2.7–5.4)***	12.0 (8.0–15.0)***
G11	Mean $\pm$ SD	Median (range)	0.8 $\pm$ 0.4	3.8 $\pm$ 2.0	0.2 $\pm$ 0.1	3.5 $\pm$ 2.0	6.1 $\pm$ 0.5	4.7 $\pm$ 1.1	15.2 $\pm$ 2.9
G12	Mean $\pm$ SD	Median (range)	1 (0–1)***	3.0 (1.0–8.0)***	0.2 (0.1–5.0)***	2.8 (0.5–7.5)***	6.2 (5.8–6.5)*	5.2 (2.6–6.2)	16.0 (10.0–21.0)
G13	Mean $\pm$ SD	Median (range)	24.4 $\pm$ 11.4	0.53 $\pm$ 0.33	0.1 $\pm$ 0.08	0.73 $\pm$ 0.09	7.1 $\pm$ 4.6	3.8 $\pm$ 0.6	11.0 $\pm$ 1.7
G14	Mean $\pm$ SD	Median (range)	24.5 (0–60)*	0.9 (0.5–1.0)	0.1 (0.0–0.3)	0.8 (0.5–0.9)*	5.0 (4.3–14)	3.9 (2.7–4.7)***	12.0 (8.0–13.0)***

The values are expressed as the mean $\pm$ SD, median, and range. All comparisons were made with control subjects. Significant differences are indicated as * for $P < 0.05$, ** for $P < 0.01$, and *** for $P < 0.001$. Student's t test was used when the data were found to be normally distributed, and the Mann-Whitney U test was used for non-normally distributed data.

Figure 1 | The frequency of the different types of neonatal jaundice. The data are presented as a percentage (%) of the total number of neonates in each group.

was found only in males. However, class III deficiency was found in 4 males and 9 females. The hyperbilirubinemia that occurs in G6PD-deficient neonates is thought to be secondary to reduced hepatic conjugation and excretion of bilirubin, rather than a result of increased bilirubin production from hemolysis²³. This idea was confirmed by our study, as no significant differences in hemoglobin levels were observed between G6PD-deficient and normal groups.

The tendency of G6PD-deficient females to develop neonatal hyperbilirubinemia is known to be related to the degree of G6PD mosaicism²⁴. Red cell mosaicism results in G6PD-deficient and G6PD-normal cell types, and the proportion of these two cell types can vary enormously, ranging from normal activity to complete deficiency. For many years, G6PD-deficient heterozygotes were not regarded as being at risk for illness. However, some studies have shown that female heterozygotes are at risk of developing severe neonatal hyperbilirubinemia^{24,25}. This is in agreement with our study, as we found 7 females with class III G6PD deficiency out of 111 neonates with neonatal hyperbilirubinemia and only 2 cases of such deficiency in anemic infants lacking hyperbilirubinemia. Therefore, it is important to identify G6PD-deficient neonates in an effort to manage severe neonatal hyperbilirubinemia. In addition, our findings strengthen the notion that G6PD-deficient infants should be followed up after discharge.

In this study, no significant differences were found between males and females with regard to G6PD concentrations with the exception that those in the early onset jaundice group, for whom a significantly higher G6PD concentration was observed in females than in males ($p > 0.05$ and $p < 0.05$, respectively). As the gene encoding the G6PD enzyme is located on the X chromosome, females should have higher G6PD enzyme levels than males (because females have two copies of the X chromosome per cell). However, because of lyonization, enzyme levels in the female are not appreciably higher than in

Figure 2 | The frequency of ABO incompatibility in the different studied groups. The data are presented as a percentage (%) of the total number of neonates in each group.

Table 3 | The frequency of G6PD enzyme deficiency in each group

Groups	N	Class II (<10% = <1.2 U/gHb)		Class III 10–60% (1.2–7.3 U/gHb)		Total
		Males	females	Males	females	
G2	38	1/38 (2.6%)	—	—	3/38 (7.9%)	4/38 (10.5%)
G3	129	1/129 (0.8%)	—	—	4/129 (3.1%)	5/129 (3.9)
G4	9	—	—	1/9 (11.1%)	—	1/9 (11.1%)
G5	42	2/42 (4.7%)	—	3/42 (7.1%)	—	5/42 (12.0%)
G6	21	—	—	—	—	—
G7	20	—	—	—	2/20 (10%)	2/20 (10%)
Total	259	4	—	4	9	17/259 (6.6%)

males²⁶. In those with early onset jaundice, 2 females had higher G6PD enzymatic activity (more than 150% of the level of the control group). The remaining females had normal values. This could explain why, overall, females had significantly higher G6PD levels than did males.

This study is the first of its type performed in the Makkah region. We found that the frequency of G6PD enzyme deficiency in all studied neonates was 6.6%. In previous national studies, the frequency of this deficiency was found to be 2%, 30.3%, 2%, 30.6%, and 14.7% in Yanbu²⁷, Al-Hofuf²⁸, Riyadh²⁹, Al-Qatif³⁰, and Al Hasa³⁰, respectively. Studies from Iran³¹, Brazil³², and India³³ found the frequency of G6PD deficiency to be 2.1%, 4.6%, and 7.5%, respectively. By contrast, studies from China³⁴, Nigeria³⁵, and Thailand³⁶ found higher frequencies of G6PD deficiency, at 18.42%, 25.5%, and 38%, respectively. In our study, 88.2% (15/17) of G6PD-deficient patients developed neonatal jaundice, of which 33.3% (5/15) were in the physiological jaundice group and the remaining cases had other types of jaundice. Our results indicate a higher frequency of jaundice in neonates with G6PD deficiency. Previous studies reported that one-third of children with G6PD deficiency develop neonatal jaundice^{22,37}. Moreover, Tanphaichitr et al. (1995) reported that 49% of G6PD patients developed neonatal jaundice, of which 28.82% were classified as physiological and 20% as pathological³⁸. The prevalence of the different subtypes of jaundice could explain the different presentations in different geographical locations worldwide.

Neonates with blood group A or B born to group O mothers are defined as having ABO incompatibility³⁹. In our study, out of 259 neonates, 12% were ABO incompatible, which is in agreement with the national data²⁸ but is lower than results of international studies^{40,41}. The direct Coombs' test was positive in only 16.1% cases, and the remaining cases were negative, which is similar to previous reports^{42,43}. These reports showed that a positive direct Coombs' test in ABO incompatible neonates increases the risk of significant hyperbilirubinemia; however, when evaluated as a screening test for all newborns, the positive predictive value was found to be as low as 23% with readmission rates that were similar irrespective of the results from the Coombs' test in ABO-incompatible children.

Table 4 | A comparison between males and females with regard to G6PD concentration (U/gHb) among the different groups

Groups	Males Mean $\pm$ SD	Females Mean $\pm$ SD	P-value
G1	12.0 $\pm$ 2.5	12.4 $\pm$ 2.4	0.828
G2	14.5 $\pm$ 6.6	11.8 $\pm$ 4.2	0.203
G3	13.7 $\pm$ 4.8	12.3 $\pm$ 4.1	0.192
G4	9.5 $\pm$ 4.0	12.7 $\pm$ 3.1	0.498 [†]
G5	8.8 $\pm$ 4.2	13.8 $\pm$ 6.8	0.057
G6	13.9 $\pm$ 2.9	20.7 $\pm$ 0.9	0.013[†]
G7	11.2 $\pm$ 1.6	9.6 $\pm$ 4.4	0.578

Student's t test was used.

[†]The Mann-Whitney U test was used in these groups.

In most cases, ABO hemolytic disease of newborns causes hyperbilirubinemia without severe neonatal anemia, which is explained by the relatively few group A or B antigens on neonatal red blood cells and the presence of A and B antigens on other tissues and in body fluids⁴⁴. This result was confirmed by our study, as most ABO incompatible cases had normal hemoglobin levels. Additionally, in our study, ABO incompatibility and G6PD were common causes of neonatal jaundice. If these are present, either phototherapy or intravenous immunoglobulin or both are administered to prevent severe jaundice and to decrease the need for exchange transfusion. Phototherapy was initiated based on the American Academy of Pediatrics phototherapy chart, whereas intravenous immunoglobulin was administered in ABO cases if the total bilirubin was two lines below the exchange transfusion line, in accordance with the exchange transfusion chart.

In the indirect hyperbilirubinemia group, one neonate presented with Rh hemolytic disease of the newborn (a frequency of 2.6% in the indirect hyperbilirubinemia group and 0.4% of all neonates). This finding is in agreement with previous studies, which state that Rh hemolytic disease of the newborn is less common than previously owing to the administration of Rh Ig, which results in a greater than 90% reduction in the alloimmunization rate among treated women^{41,45}.

In this study, one neonate in the indirect hyperbilirubinemia group was polycythemic (2.6%). Although polycythemia can reflect normal fetal adaptation, it has been thought to be responsible for abnormalities in neonates⁴⁷.

MDA levels are a sensitive indicator of lipid peroxidation and thus of oxidative stress. Increased concentrations of free oxygen radicals in newborns damage the cell membrane through lipid peroxidation, and this damage may be associated with various pathologies such as hypoxic ischemic encephalopathy, intraventricular hemorrhage, necrotizing enterocolitis, and bronchopulmonary dysplasia⁴⁶. Bilirubin is an effective scavenger of oxidant radicals, and its concentration is increased during oxidative stress. In this study, the MDA concentrations were significantly higher in all of the neonates with the various types of jaundice than in the control group (Table 2). In agreement with our results, Yigit et al (1999) found MDA concentrations in infants with hyperbilirubinemia to be remarkably high⁴⁷. These MDA levels were thought to result from the response of the neonates to oxidative stress. It has been suggested that hyperbilirubinemia might play a defensive role against oxidative stress because the body's antioxidant mechanisms are not yet fully developed. It is known that biliary pigment can protect the cell membrane from lipid peroxidation⁴⁸. In addition, some neonates in our study had infections, perinatal asphyxia, and respiratory distress syndrome, and approximately 30% of our cases were preterm. All of these conditions are considered to contribute to the elevation of MDA concentrations^{48–51}.

In contrast to our results, Kumar et al (2007) found lower MDA levels in neonates with jaundice. This finding may arise from their study being conducted in neonates without high bilirubin levels⁵². At higher bilirubin levels, the antioxidant effects of bilirubin might be

negated by its direct cellular toxicity. In addition, Kumar et al excluded preterm neonates as well as those with infections, perinatal asphyxia, respiratory distress or major malformation, which were all included in our study. Furthermore, we found no significant correlation between bilirubin and MDA levels among the studied groups. This may indicate that factors other than MDA levels may contribute to the increased oxidative stress in our cases, such as stress and prematurity. Our results confirm those of Yigit et al (1998), who did not find a correlation between MDA levels and bilirubin concentrations in infants with non-hemolytic hyperbilirubinemia⁴⁸. Moreover, we found a significant negative correlation between MDA and hemoglobin levels in the group with persistent jaundice. This may indicate that anemia accompanies many pathological processes in neonates and can negatively affect prognosis. Thus, increased MDA levels may indicate oxidative stress⁵³.

One limitation of this study is that some factors that may have led to neonatal hyperbilirubinemia were not examined such as spherocytosis, thalassemia and sickle cell anemia.

Conclusions and recommendation. ABO incompatibility and G6PD deficiency are frequent causes of neonatal jaundice in Makkah, whereas Rh incompatibility and polycythemia are rarely observed. The MDA level could serve as an indicator of oxidative stress. Determinations of ABO and Rh incompatibility, G6PD deficiency and complete blood counts at birth are recommended to avoid serious complications.

Methods

Subjects. The protocol for this study was approved by the Umm Al-Qura University ethics committee. All participants gave informed consent in accordance with the declaration of Helsinki. The analytical methods were carried out in the research laboratory of the Biochemistry Department, Faculty of Medicine, Umm Al-Qura University in accordance with the approved guidelines.

This study was carried out from January 2011 to January 2013. All neonates were attended to at Alnour Hospital, Makkah, Saudi Arabia. The description and classification of all subjects is summarized in Table 1, with healthy neonates serving as a control group (G1). The jaundiced neonates were divided into 5 groups according to their indirect and direct bilirubin levels, the time of jaundice onset, and the persistence of jaundice¹³. Members of the indirect hyperbilirubinemia group (G2) were characterized as having increased levels of indirect bilirubin, where the concentration of indirect bilirubin was found to be ≥ 5 mg/dl at 0 h of age, ≥ 10 mg/dl at 24–48 h of age and ≥ 13 mg/dl at more than 48 h of age. In the physiological jaundice group (G3), the indirect bilirubin levels were not high enough to allow stratification based on age. In the direct hyperbilirubinemia group (G4), the levels of direct bilirubin were determined to be > 2 mg/dl ($> 20\%$ of total bilirubin). Group 5 (G5) was composed of neonates with high bilirubin levels that had persistent jaundice for > 2 weeks in full term neonates and > 3 weeks in preterm neonates. Group 6 was composed of neonates with early onset jaundice (G6), in which jaundice was observed in the first 24 h of life but indirect bilirubin remained < 5 mg. Neonates in the anemic group (G7) were free of jaundice.

Sample collection. Four ml of blood was collected from neonates at the time of admission under aseptic conditions. 2 ml of this blood was centrifuged for 15 minutes, and the serum was separated to allow MDA and total/direct bilirubin measurements. The remaining 2 ml of blood was collected in EDTA tubes and used to perform blood counts, blood group analyses, G6PD enzyme assays, and direct antiglobulin tests. In addition, 1 ml of blood was taken from each mother for blood group determination.

All participants were subjected to a:

1. Full clinical history.
2. Complete hemogram analysis using a Sysmex XT 2000i hematology analyzer (Sysmex Corporation of America, Long Grove, Illinois, USA), including red blood cells count, hemoglobin, packed cell volume, mean corpuscular hemoglobin, mean cell volume, mean corpuscular hemoglobin concentration, red cell distribution width, white blood cells count and platelet count.
3. ABO grouping of baby and mother (Dialab, Austria).
4. Direct antiglobulin test (Coombs' test) tube method. The reagent was supplied by Core Diagnostics LTD, Birmingham - United Kingdom.
5. Quantitative measurement of G6PD enzyme levels by the ultraviolet kinetic method (Trinity Biotech, USA)¹⁴.
6. MDA enzyme assay with the OxiSelect TBARS Assay Kit from Cell Biolabs, Inc. (USA)¹⁵.
7. Determination of direct and total bilirubin levels using the Dimension RxL max integrated chemistry system (Dade Behring, USA)¹⁶.

Statistical analyses. All statistical analyses in this study were performed using the SPSS program (version 16). All quantitative data are described as the mean $\pm$ SD, the median, or the range. Comparisons between the control group and all other groups were performed using the Student's t test or the Mann-Whitney U test, depending on the distribution of the data. The Kruskal-Wallis H test was used when comparing all groups, as some of the data were not normally distributed. In addition, the Spearman correlation was used when correlating MDA with other parameters studied. Statistical significance was defined as $p < 0.05$.

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Author contributions

A.E.: Design of the study, clinical selection, diagnosis and classification of the cases, writing and revising of the main manuscript text. E.E.N.E.: Design of the study, performance of biochemical laboratory investigations, preparing tables and figures, editing, styling, writing and revising of the main manuscript text. A.Z.: Design of the study, performance of hematological laboratory investigations, statistical analysis, writing and revising of the main manuscript text. S.M.: Performance of hematological laboratory investigations, writing and revising of the main manuscript text.

Additional information

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