Correlation Between Maternal and Neonatal Vitamin B12 Serum Levels in Fullterm Neonates with Physiologic Jaundice

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Abstract

Introduction: Vitamin B12 (vitB12) deficiency causes failure of red blood cells (RBC) maturation leading to cell lysis causing excess heme production resulting in indirect hyperbilirubinemia.

Purpose: We aimed to assess vitB12 serum levels in neonates with jaundice and compare these levels with that of apparently healthy neonates and detect the impact of the maternal vitB12 on both vitB12 and bilirubin of their neonates.

Patients and methods: Complete blood count (CBC), C-reactive protein (CRP), and VitB12 were assessed in both groups; 60 full term neonates with jaundice (Group I) and 60 full term controls. VitB12 levels were assessed for the mothers of all included neonates.

Results: VitB12 in jaundice neonates were significantly lower than that of controls. Maternal vitB12 was positively correlated with neonatal vitB12 and negatively correlated with neonatal total serum bilirubin levels (TSB).

Conclusion: Low maternal vitB12 affects both vitB12 and bilirubin levels of their neonates.

Introduction

Physiologic jaundice is caused by a combination of increased bilirubin production (secondary to accelerated destruction of erythrocytes and decreased excretory capacity secondary to low levels of ligandin in hepatocytes) and low activity of the bilirubin-conjugation enzyme uridine diphospho glucuronyltransferase (UDPGT) [1].

VitB12 is essential for maturation and proliferation of RBCs [2]. Deficiency of VitB12 causes failure of RBC maturation leading to cell lysis which causes excess heme production resulting in indirect hyperbilirubinemia [3].

Hemolysis in patients with vitB12 deficiency is a well-recognized phenomenon and it is due to intramedullary destruction of erythrocytes (ineffective erythropoiesis). Homocysteine increases the risk of hemolysis in vitB12 deficiency and there is a high frequency of vitB12 deficiency in asymptomatic patients with homozygous methylenetetrahydrofolate reductase (MTHFR) C677T mutation, a known cause of hyperhomocysteinemia [4].

Patients demonstrated complete resolution of hemolysis with simultaneous normalization of serum homocysteine levels after vitB12 treatments [5].

Aim of the Work

The aim of this study is to assess vitB12 serum levels in neonates with physiologic jaundice, compare them with those of apparently healthy neonates and detect the impact of maternal serum vitB12 on both vitB12 and serum bilirubin of their neonates.

Patients and Methods

One hundred twenty full term neonates and their mothers were enrolled from Neonatal Care Unit and Obstetric Department of Minia University Hospital, Minia, Egypt in the period from February to December 2017.
Neonates with gestational age less than 37 weeks and more than 42 weeks, ages less than 3 days and more than 7 days, neonates with bad general condition (absent or weak moro or suckling), organomegaly; neonates with Rh or ABO incompatibility, Glucose 6-phosphate Dehydrogenase Deficiency (G6PD), hypothyroidism, polycythemia, +ve CRP, Direct bilirubin >20%, CBC: signs of infection (thrombocytopenia, leukopenia leukocytosis, differential count of WBCs shows shift to left) were excluded from our study.

**Laboratory investigations**

Five ml of venous blood were obtained from each neonate under sterile venipuncture and divided as follow:

One ml was added to EDTA containing tube for performing CBC.

Four ml were put on a plain plastic tube, incubated for 20 minutes at 37°C and centrifuged at 3000 r.p.m for 10 minutes. The serum aliquot was divided into two parts; one for serum bilirubin and CRP while the remaining part stored at -20°C in Eppendorf cups until processing serum vitB12.

**CBC assessment**

Done by automated cell counter symex KX21N (JAPAN).

**Serum bilirubin**

Done by automated chemical analyzer Konelab 60 i (FILAND).

**CRP**

Done by qualitative and semiquantitated determination latex agglutination spin react (SPAIN) [6,7].

**Serum VitB12 level**

By quantitative method using ELISA kit. Glory Science Co, Ltd (CHINA, HONG KONG).

VitB12 levels were also measured for the mothers of all included neonates.

**Statistical methods**

The collected data were coded, tabulated, and statistically analyzed using SPSS program (Statistical Package for Social Sciences) software version 24.

Descriptive statistics were done for parametric quantitative data by mean, standard deviation and minimum & maximum of the range, while they were done for categorical data by number and percentage.

Analyses were done for parametric quantitative data between the two groups using independent samples T test.

Analyses were done for qualitative data using Chi square test.

Correlation between two quantitative variables was done by using Pearson's correlation coefficient. Weak (r=0-0.24), fair (r=0.25-0.49), moderate (r=0.5-0.74), strong (r=0.75-1).

Simple linear regression analysis was used for prediction of vitB12 level.

ROC curve analysis of vitB12 was used for prediction of jaundice severity. The level of significance was taken at (P value < 0.05).

**Results**

There was no significant difference between the two studied groups regarding the demographic and the laboratory data as shown in tables 1 and 2. There was no significant difference between the two groups regarding the levels of Hb, platelets, WBC and neutrophils. Also, both groups had -ve CRP.

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**Table 1: Demographic data of the studied groups.**

|                      | Control (n=60) | Patient (n=60) | P value |
|----------------------|---------------|---------------|---------|
| Age                  |               |               |         |
| Range (3-7)          | (3-7)         | (3-7)         | 0.413   |
| Mean ± SD            | 4.3±1.3       | 4.4±1.1       |         |
| Sex                  |               |               |         |
| Male                 | 29(48.3%)     | 26(43.3%)     | 0.583   |
| Female               | 31(51.7%)     | 34(56.7%)     |         |
| Mode of delivery     |               |               |         |
| SVD                  | 22(36.7%)     | 24(40%)       | 0.707   |
| CS                   | 38(63.3%)     | 36(60%)       |         |
| Gestational age      |               |               |         |
| Range (37-41)        | (37-41)       | (37-41)       | 0.930   |
| Mean ± SD            | 38.4±1        | 38.4±1        |         |
| Weight               |               |               |         |
| Range (2.5-3.7)      | (2.5-3.6)     | (2.5-3.6)     | 0.413   |
| Mean ± SD            | 3.1±0.4       | 3.1±0.3       |         |

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**Table 2: CBC, CRP findings in both groups.**

|                      | Control (n=60) | Patient (n=60) | P value |
|----------------------|---------------|---------------|---------|
| Hb                   |               |               |         |
| Range (14-18.9)      | (14.3-19)     | (14.3-19)     | 0.696   |
| Mean ± SD            | 16.4±1.4      | 16.5±1.4      |         |
| Platelets            |               |               |         |
| Range (158-451)      | (150-448)     | (150-448)     | 0.767   |
| Mean ± SD            | 293±79.7      | 288.4±91.5    |         |
| WBCs                 |               |               |         |
| Range (4.5-10.9)     | (4-11)        | (4-11)        | 0.650   |
| Mean ± SD            | 7.5±1.8       | 7.7±2         |         |
| Neutrophils          |               |               |         |
| Range (48-67.7)      | (48.5-65.1)   | (48.5-65.1)   | 0.606   |
| Mean ± SD            | 55.7±5.2      | 55.2±4.7      |         |
| CRP                  | -ve           | -ve           |         |
TSB levels were significantly higher in patients than controls (15.7 ± 2.9 and 3 ± 0.8 respectively, p < 0.001*). On contrast, neonatal vitB12 levels were significantly lower in patients than in controls (226.4 ± 25.5 and 421.8 ± 81.3 respectively, p < 0.001*). VitB12 levels were also significantly lower in the patients’ mothers than that of controls (185.1 ± 16 versus 217.7 ± 24) (Table 3).

Neonatal vitB12 levels were negatively correlated with their serum bilirubin levels (r = -0.873, p < 0.001*) (Table 4A).

Maternal vitB12 levels were negatively correlated with neonatal serum bilirubin levels and positively correlated with neonatal vitB12 levels (r = -0.923, p < 0.001* and r = 0.828, p < 0.001* respectively) (Table 4B).

By using Linear Simple Regression analysis, we can predict serum vitB12 level from TSB level by the following equation:

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\text{VitB12} = 345.251 + (-7.572 \times \text{TSB})
\] (Table 5).

### Table 3: comparison between Neonatal TSB, VitB12 and maternal VitB12 between both groups.

| Patient group       | Neonatal TSB | P value |
|---------------------|--------------|---------|
| Controls (n=60)     | (12.4-4.9)   | <0.001* |
| Patients (n=60)     | (9-20.5)     |         |
| Mean ± SD           | 15.7±2.9     |         |

### Table 4A: correlation between neonatal vitB12 and other parameters.

| Parameter          | R   | P value |
|--------------------|-----|---------|
| Age                | 0.040 | 0761    |
| Gestational age    | 0.182 | 0.164   |
| Weight             | -0.071 | 0.588   |
| TSB                | -0.873 | <0.001* |

### Table 4B: correlation between maternal vitB12 and other parameters.

| Parameter          | r    | P value |
|--------------------|------|---------|
| Neonatal TSB       | -0.923 | <0.001* |
| Neonatal vitB12    | 0.828 | <0.001* |

### Table 5: Linear simple regression analysis for prediction of VitB12 levels from TSB level.

| B     | P value | R | R² | Regression equation |
|-------|---------|---|----|---------------------|
| Constant | 345.251 | <0.001* | 0.873 | 0.762 | VitB12 = 345.251 + (-7.572 x TSB) |
| TSB    | -7.572  | <0.001* |      |                |
Intramedullary destruction or hemolysis of fragile and abnormal red blood cell precursors is the result of ineffective erythropoiesis secondary to defective DNA and cell maturation. The hemolytic picture may resemble microangiopathic hemolytic anemia. If such condition persisted, it may deplete iron storage and lead to concomitant iron deficiency anemia [13].

In newborns, the functional capacity of liver is only about 1% of that of the adult. RBC lysis and low functional capacity of liver result in inefficient clearance of bilirubin and so indirect hyperbilirubinemia develops [14].

Our result is in agreement with that of Eroglu et al. [3] who found that the mean vitB12 level in patients was 119.9 ± 43.9 ng/L (42.35 - 178 ng/L) while in controls was 286.17 ± 97.43 ng/L (207.90 - 624.10 ng/L). They also stated that low vitB12 level has been observed as a risk factor in neonatal hyperbilirubinemia.

In our study, vitB12 levels in mothers of neonates with jaundice were lower than their levels in mothers of controls and the maternal vitB12 level was positively correlated with neonatal vitB12.

This result was in agreement with that of Obeid et al., 2017 [11] who stated that women who began pregnancy with depleted stores (low or very low plasma cobalamin) will give birth to depleted infants who are likely to develop deficiency symptoms during the first few weeks or months postpartum. Newly ingested cobalamin during pregnancy and lactation is transferred to the child and is not likely to correct cobalamin status in depleted women [11].

VitB12 deficiency is more common in pregnant, lactating women and their young children in developing countries due to malabsorption which is exacerbated by low dietary intake of the vitamin [15,16].

VitB12 levels fall progressively throughout pregnancy in most healthy women. This decline begins in the first trimester and levels may drop to 50 percent of pre-pregnancy values by the third trimester [17]. That is why in our study, maternal vitB12 levels in both groups were lower than the normal adult levels.

The normal physiologic changes during pregnancy contribute to the fall in vitB12 levels. Hemodilution (in which the plasma volume increases by 40 to 50 percent) lowers the vitamin concentration. Because the kidneys filter blood at a higher rate during pregnancy, more vitB12 will be lost in urine. VitB12 is actively transferred across the placenta to the developing fetus. VitB12 levels in umbilical cord and fetal blood are twice as high as in maternal blood, indicating preferential delivery to the fetus [17].

This is the first study showing that maternal vitB12 was negatively correlated with neonatal hyperbilirubinemia.

There is an evidence of increased vitB12 absorption during pregnancy, with the newly absorbed vitB12 being more important to placental transport than maternal liver stores. So that, lack of vitB12 rich diet of the mother may cause vitB12 deficiency in newborn [3]. So, it is recommended that vitB12 supplement is necessary to avoid both neonatal vitB12 deficiency and hyperbilirubinemia.

We could also recommend vitB12 intake for neonates with exaggerated physiologic jaundice.

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Authors' Contributions

Reem A. Abdel Aziz and Mostafa A. Abu Elela conceived the study, carried out its designing, coordinated the implementation, helped to perform the statistical analysis and drafted the manuscript. RA designed the study, participated in the analysis and interpretation of data and revised the statistics and final draft of the manuscript. MA was responsible for interpretation of laboratory data of patients and revision of the manuscript. All authors read and approved the final manuscript.

Conflicting Interests

The authors declare that there are no potential conflicts of interest with respect to the research, authorship, and/or publication of this article.

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