Umbilical Cord Blood Lipid Parameters and its Correlation with Birth Weight of Neonates

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Abstract: Introduction: Cardiovascular diseases are the leading cause of mortality in India. The important factor associated is dyslipidemia. A strong relationship has been seen in epidemiological studies between cholesterol and cardiovascular disease. The present study was planned to analyse cord blood lipid profile, apolipoproteins and atherogenic index and to see their relationship with birth weight. Method: Our study group consisted of 300 healthy fullterm newborn. The cord blood was collected immediately after a normal delivery. The blood was tested to determine lipid parameters and was correlated with birth weight. Results: The results showed that the levels of (TC), triglyceride, (HDL) and atherogenic index was higher whereas LDL was lower in high birth weight babies as compared to low birth weight and normal birth weight newborns. The correlation was positive with TC, LDL-Cholesterol, ApoB and atherogenic index but negative with TG, HDL-C and ApoA-I.

Keywords: Lipid profile, Apolipoproteins, Atherogenic index, Cord blood.

1. Introduction

Atherosclerosis is the major cause of global mortality and will continue to dominate trend in future. (1) Atherosclerosis is a process that begins in early in life and progresses silently for decades, some of scientist believe that atherosclerotic lesions may have its genesis during childhood. (2, 3) increased risk of CVD is associated with lifestyle and various medical conditions, such as hypercholesterolemia, hypertension, smoking, obesity and inadequate physical activity. (4)

A foetus needs a considerable amount of cholesterol for development of tissues and organs. After birth lipid transport system is transformed from one containing low VLDL and LDL levels to adult system with a relatively high LDL levels which continues to increase with age. Cord blood contains all adult lipoproteins and apolipoproteins. (5) Abnormal lipid profiles in childhood persist into adult life and elevated ApoB in adult have been linked to atherosclerosis in later life. (6) On the basis of research conducted have shown that the high prevalence of atherosclerosis risk factor and increased morbidity and mortality due to cardiacological diseases within the urban areas is more than rural areas. (7, 8) Moreover, several maternal and fetal factors, such as hypertension, diabetes, obesity and low or high birth weight can influence fetal plasma lipids. (9)

Disorders of lipid metabolism and elevated levels of low density lipoprotein (LDL-C), total cholesterol (TC) and apolipoproteinB (ApoB) levels in young adults have been connected with cardiovascular diseases in the life. (7) Dirisamer et al (2006) have showed evidence that levels of ApoB, ApoB/apoA-1 ratio as well as LDL-C concentrations are sensitive indicators for later coronary heart disease in children. (8) ApoB/ ApoA-1 ratio known as atherogenic index is sensitive marker to track coronary artery disease is found to be measured closely during first year of life. (10)

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Screening patients for ApoA-I and ApoB levels may significantly improve ability to properly evaluate CAD risk. (11) with Low birth weight is associated the increased incidence of CVD, hypertension and type II diabetes (12) Changes in blood lipids in low birth weight newborns with relative insulin intolerance can increase the risk of CVD in adulthood. LBW is a risk of later atherosclerotic diseases that is equal to smoking or hypertension at puberty. (13) Therefore it seems that there is some relation between birth weight and mortality from CVD in adulthood. Hence present study was planned to check the correlation of lipid parameters and birth weight of newborns.

2. Material and Methods

The present prospective study was conducted n the department of Biochemistry, Geetanjali Medical College and hospital, Udaipur. A total of 300 healthy neonates (following healthy normotensive pregnancy) were included in the study and were divided on the basis of birth weight.

Inclusion criteria for mothers:
Healthy mother only on iron folic acid and calcium supplementation.

Exclusion criteria for mothers:
History with alcoholism, smoking hypertension, thyroid disorders, diabetes mellitus, renal diseases, hypercholesterolemia, twins, liver diseases, tuberculosis and asthma, pregnancy induced hypertension.

Inclusion criteria for neonates:
Gestational age between 35-42 weeks and Absence of congenital anomalies.
Exclusion criteria for neonates
Congenital malformations, Neonates born to mother with maternal illness, Neonates with perinatal problems like hypoglycemia, pathological jaundice, Instrumental delivery including extraction and also Neonates with hypoxic ischemic encephalopathy and sepsis.

Sample collection
After delivery and cord clamping umbilical venous blood was taken from maternal umbilical end. Serum was separated and analyzed for lipid profile (total cholesterol, Triglyceride, HDL-C, LDL-C, and VLDL-C) and Apolipoproteins (ApoB, ApoA-1).

Statistical Analysis
The mean and standard deviation has been used to define data in each group. The data were correlated with birth weight using Pearson correlation.

3. Observation and Results

4. Results
A total of 300 newborns, 33 (11%) were low birth weight, 171 (57%) were normal birth weight and 96(32%) were high birth weight neonates were studied. The mean total serum lipid levels in these newborns were compared and mean and standard deviation were calculated. Table 1 shows mean and SD values in different groups. The results showed that the levels of total cholesterol (TC), triglyceride (TG), high density lipoprotein cholesterol (HDL-C), low density lipoprotein cholesterol (LDL-C), very low density lipoprotein cholesterol (VLDL-C) and Apolipoproteins (ApoB, ApoA) were significantly higher in low birth weight neonates compared to normal birth weight neonates. The results are shown in Table 2.

Figure 1: Correlation scatter plots with different parameters

Table 1: Mean ± SD level of Lipid profile in cord blood in different newborn having different weight.

<table>
<thead>
<tr>
<th>S.N.</th>
<th>Parameters</th>
<th>Low birth weight &lt;2.5 kg (N=33)</th>
<th>Normal birth weight 2.5-4.0 kg (N=171)</th>
<th>High Birth weight &gt;4.0 kg (N=96)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Total Cholesterol</td>
<td>67.90±17.13</td>
<td>68.78±16.92</td>
<td>70.71±16.83</td>
</tr>
<tr>
<td>2</td>
<td>Triglyceride</td>
<td>47.72±21.10</td>
<td>52.90±23.50</td>
<td>54.27±24.91</td>
</tr>
<tr>
<td>3</td>
<td>HDL-C</td>
<td>27.15±9.79</td>
<td>29.52±10.62</td>
<td>30.10±10.98</td>
</tr>
<tr>
<td>4</td>
<td>LDL-C</td>
<td>31.21±14.15</td>
<td>28.67±14.97</td>
<td>29.75±16.27</td>
</tr>
<tr>
<td>5</td>
<td>VLDL-C</td>
<td>9.54±4.22</td>
<td>10.58±4.70</td>
<td>10.85±4.98</td>
</tr>
<tr>
<td>6</td>
<td>ApolipoproteinB</td>
<td>31.70±7.30</td>
<td>31.16±7.50</td>
<td>31.40±5.26</td>
</tr>
<tr>
<td>7</td>
<td>ApolipoproteinA-1</td>
<td>52.34±9.01</td>
<td>52.92±9.43</td>
<td>52.30±8.79</td>
</tr>
<tr>
<td>8</td>
<td>ApoB/ApoA-1</td>
<td>0.63±0.22</td>
<td>0.62±0.23</td>
<td>0.60±0.11</td>
</tr>
</tbody>
</table>

Table 2: Correlation of birth weight with lipid parameters

<table>
<thead>
<tr>
<th>S.No.</th>
<th>PARAMETERS</th>
<th>Mean ± SD</th>
<th>r-value</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Total Cholesterol mg/dL</td>
<td>70.47±16.79</td>
<td>0.430</td>
<td>0.0004</td>
</tr>
<tr>
<td>2</td>
<td>Triglyceride mg/dL</td>
<td>54.80±23.90</td>
<td>-0.083</td>
<td>0.205</td>
</tr>
<tr>
<td>3</td>
<td>HDL-C mg/dL</td>
<td>31.16±10.59</td>
<td>-0.111</td>
<td>0.134</td>
</tr>
<tr>
<td>4</td>
<td>LDL-C mg/dL</td>
<td>27.27±15.20</td>
<td>0.259</td>
<td>0.004</td>
</tr>
<tr>
<td>5</td>
<td>VLDL-C mg/dL</td>
<td>10.15±0.78</td>
<td>0.598</td>
<td>0.000</td>
</tr>
<tr>
<td>6</td>
<td>ApolipoproteinB mg/dL</td>
<td>30.68±6.97</td>
<td>0.273</td>
<td>0.0005</td>
</tr>
<tr>
<td>7</td>
<td>ApolipoproteinA-1 mg/dL</td>
<td>54.36±8.89</td>
<td>-0.187</td>
<td>0.060</td>
</tr>
<tr>
<td>8</td>
<td>ApoB/ApoA-1</td>
<td>0.61±0.19</td>
<td>0.304</td>
<td>0.000</td>
</tr>
</tbody>
</table>
density lipoprotein (HDL) and atherogenic index were higher whereas low density lipoproteins was lower in high birth weight babies as compared to low birth weight and normal birth weight newborns. The differences were statistically non significant.

Table2 and Figure1 showed correlation among the measured parameters and weight of the newborn. The correlation of lipid profile, apoliproteins and atherogenic index with birth weight were showed. The positive correlation was found with total cholesterol, LDL-Cholesterol, Apo B and atherogenic index but TG, HDL-C and ApoA-1 were negatively correlated. Total cholesterol was positively correlated with high significance. Triglyceride and HDL-cholesterol were negatively correlated without significance. LDL- cholesterol showed positive correlation with high significance.

The Apolipoproteins were also estimated and correlated with neonatal weight. Apolipoprotein A -1 showed negative correlation like HDL- cholesterol and was not significant. Apolipoprotein B was positive correlated with significance. The ratio or atherogenic index was positive correlated with significance and same was the correlation with LDL-C/ HDL-C.

5. Discussion

The cord blood lipid parameters screening of neonates in this part of southern Rajasthan were almost comparable with other reports. Atherosclerosis originates during childhood and the serum lipid levels are a key factor in the process. The observation in cord blood reveals to study the risk factor variable in the early stage. The observations in infancy provide a background for the studies in older children and adults. According to newborn weight the subjects were divided into three groups, group I were low birth weight neonates with weight less than 2.5kg, Group II normal birth weight neonates having weight 2.5kg to 4kg and group III as high birth weight neonates with more than 4kg.

The mean cholesterol levels were 70.47± 16.79 mg/dl which is less than the reports of Puspedra et al, 2013 and other where the values were 83.30± 28.10, 76± 19, 94± 32, 79± 17 and even 105.6± 17.1. The values were higher ranging from 64± 19 has been noted in other studies. (16, 17, 18) The mean triglyceride levels noted in present study were much lower than the adult values. Perhaps the quiescent state of fat utilization in the fetus and the absence of need for fat mobilization is responsible for low serum triglyceride at birth. The values of HDL-C and LDL-C were similar to those reported in other studies. (19)

Fetal growth retardation establishes a lifelong irreversible atherogenic profile and men with low birth weight have been reported to have an atherogenic profile. In the present study apolipoproteinA-1 and apolipoproteinB were estimated and correlated with birth weight. The results showed that the ApoB is most strongly associated with ischemic heart diseases risk. A decrease in ApoB of 10% was associated with 22% reduction in risk of ischemic heart disease. (20) In our study the Apo B levels in the cord blood were found to be 30.68± 6.97 which was similar to the values of 28.10± 16.90 reported by a past study. (21)

Birth weight is a measure of fetal growth which summates body size, body length and subcutaneous fat. Studies have shown that babies who have a reduced birth weight in relation to gestation tend as adults to develop syndrome “X” a combination of hyptension, non- insulin dependent diabetes mellitus, disordered lipids, hyperinsulinemia, obesity and abdominal fatness.(22) The present study demonstrated that cord blood cholesterol level do not have significant association with the birth weight as has been reported by kumar et al.(23) Birth weight below and above 2.5kg did not find any significant association with any of the other lipid parameters except LDL and Apolipoprotein B. In studies by Desai also showed no association (24) but in study by Mathur a significantly lower cholesterol was found among babies weighing lower than 2.5 kg. Though atherogenesis is a pediatric process has been known for decades, the reversibility of the injuries in this early phase of life is open to speculate. (25)

6. Conclusion

The findings of the study again reaffirm the link between prenatal factors and cord blood lipid parameters.

7. Acknowledgement

We sincerely thank R.N.T. Medical College and associated hospital, Udaipur and Geetanjali Medical College and Hospital, Udaipur for extending all the facilities for conducting the work. The authors are also grateful to authors / editors / publishers of all those articles, journals and books from where the literature for this article has been reviewed and discussed.

References


