The Role of Hemic Hypoxia in the Development of Sensorineural Hearing Loss in Children Associated with Hepatitis B

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Authors' contributions

This work was carried out in collaboration among all authors. Author DSI designed the study, performed the statistical analysis, wrote the protocol and wrote the first draft of the manuscript. Authors AAK and ESA managed the analyses of the study. Author ESA managed the literature searches. All authors read and approved the final manuscript.

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ABSTRACT

Background: There are not enough studies and evidenced researches conducted related to this topic. Therefore, we studied fetal haemoglobin in various somatic diseases in children with sensorineural hearing loss associated with hepatitis B.

Materials and Methods: 26 children with sensorineural hearing loss associated with hepatitis B, aged from 5 to 18 years, were examined. The comparison group consisted of 8 children with sensorineural hearing loss without concomitant somatic pathology. The control group consisted of 12 healthy children. The compulsory examination plan for patients included generally accepted laboratory and instrumental diagnostic methods: complete blood count, urine, feces, Wasserman reaction, ECG.

Results: Hb concentration in blood inpatient children with sensorineural hearing loss of the associated chronic hepatitis B (CHB) was reduced significantly by 58% compared with the healthy children. In children with Sensorineural Hearing Loss (CHT) without CHB, the studied parameter

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dysfunction resulting from chronic hepatitis B. Endothelial for oxygen and its delivery can be disrupted by the delicate balance between the need of liver tissue angiogenesis and fibrosis. Consequently, the release of HIF-1 from stellate cells is stimulated, which affects the progression of angiogenesis and fibrosis. Therefore, HbF is a marker of tissue hypoxia not only in newborns but also in various adult pathologies. Despite the considerable interest in the study of fetal haemoglobin in various somatic diseases, in the available modern literature, we have not found data on its study in children with sensorineural hearing loss associated with hepatitis B.

**2. MATERIALS AND METHODS**

To solve the tasks for the period from 2016 to 2018. On the basis of the Republican Scientific
and Practical Medical Center of Pediatrics, 26 children with sensorineural hearing loss associated with hepatitis B, aged from 5 to 18 years, were examined. The comparison group consisted of 8 children with sensorineural hearing loss without concomitant somatic pathology. The control group consisted of 12 healthy children.

All patients were admitted to the hospital in the acute phase of sensorineural hearing loss associated with hepatitis. A detailed clinical diagnosis was made on the basis of the nature of complaints, anamnesis, clinical examination results, laboratory and instrumental methods of diagnosis, taking into account information from the patient's outpatient card (extracts of case histories of previous hospitalizations, data of dynamic observation of the patient in the clinic). In making the diagnosis, modern classifications of sensorineural hearing loss in children were used.

Criteria for the inclusion of patients in the study were proven viral etiology of sensorineural pathology; proven signs of HBV.

Exclusion criteria for the study were: primary pathology of the biliary system (primary sclerosing cholangitis, primary biliary cirrhosis), signs of secondary liver damage in patients with chronic diseases of the biliary tract and intestines (cholelithiasis, chronic cholecystitis, stenotic papillitis, Crohn's disease, ulcerative colitis); extrahepatic obstruction of the portal vein, associated with the consequences of surgical interventions, portal vein thrombosis, tum cholic pathology, congenital developmental abnormalities, injuries; Budd-Chiari syndrome; patients with fever associated with concomitant diseases (acute respiratory infections, pneumonia, acute intestinal infections, pyelonephritis, etc.); acute and chronic diseases of the broncho-pulmonary system.

All clinical, anamnestic and laboratory and instrumental data were entered into a detailed map developed by us. The map noted the patient's complaints, of which more often there was increased fatigue, weakness, memory and sleep disorders, dyspeptic disorders (nausea, belching, vomiting), pain and heaviness in the right hypochondrium, epigastric pain, nausea, loss of appetite. Complaints about shortness of breath and temper, cough, pain in the heart area, the presence of heartbeat and rhythm disturbances, and changes in blood pressure were investigated in detail.

The presence of jaundice, pruritus, fever, manifestations of hemorrhagic syndrome (gingival, nasal, gastroesophageal, hemorrhoidal bleeding), arthralgia, stool disorders, flatulence, and dysphagia was taken into account. An objective examination focused on manifestations of portal hypertension and signs of disease activity for the presence of liver signs (spider veins, palmar erythema), xanathomas, Dupuytren's contracture, lymph nodes, abdominal veins, ascites, peripheral edema, gynecomastia, and the size of the liver and spleen.

The compulsory examination plan for patients included generally accepted laboratory and instrumental diagnostic methods: complete blood count, urine, feces, Wasserman reaction, ECG. A study was conducted in the blood of total protein and protein fractions, immunoglobulins, Circulating Immune Complexes (CIC), lipoproteins, cholesterol, bilirubin, urea, creatinine, amylase of blood and urine, determined the activity of Alanine Aminotransferase (ALT) and Aspartate Aminotransferase (AST), GTP, alkaline phosphatase. The coagulogram was determined: fibrinogen content, XIII coagulation factor, fibrinolytic activity, fibrin monomers using ethanol test, fibrinogen-fibrin degradation products by the method.

For the isolation and purification of HbF, alkaline denaturation with 1.2 M NaOH, salting out with ammonium sulfate, gel filtration on a column with Sephadex G-25 (working buffer — 0.05 M phosphate buffer pH 7.4), and ion-exchange chromatography on DEAE Sephadex was used G-50 on 0.01 M Tris-chloride buffer pH 8.1. The quantitative determination of HbF was carried out by electrophoresis on an agar gel with sodium dodecyl sulfate. The HbF level in the control group was 2.26±0.02 g/l, which corresponds to literary data. Gender differences in the control group were absent.

2.1 Statistical Analyses

Data analysis was performed using the STATISTICA v.6.0 Windows XP application package. Descriptive statistics of the trait included arithmetic average (M), minimum and maximum values, median (Me) and interquartile range [Q25-Q75]. When comparing the obtained results, the Mann-Whitney test was used due to the inconsistency of the analyzed data with the law of normal distribution. The relationship
between signs was studied by the Spearman (R) correlation analysis method. Differences were considered statistically significant at p<0.05.

3. RESULTS AND DISCUSSION

As can be seen from the presented research results (Table 1), the haemoglobin values of blood in the examined children with sensorineural hearing loss of the associated CHB was significantly reduced by 58% compared with the healthy children. In children with CHT without CHB, the studied parameter decreased when compared with healthy children by 25%.

Other dynamics were noted with respect to fetal haemoglobin in the blood of the examined children with combined pathology. Analysis of the results showed a significant increase in the level of fetal haemoglobin in the blood of children with CHT associated with hepatitis B on average by 1.5 times, indicating hypoxia.

Hypoxia in chronic liver disease can be both local and systemic. Deficiency of oxygen entering the liver parenchyma occurs during chronic hepatitis as a result of several mechanisms that include vascular resistance, intrahepatic shunts, intravascular thrombus formation, reduction in the area of sinusoids and sinusoidal capillaries. Endothelial dysfunction (ED) plays a significant role in the development of these processes. ED is involved in the formation of portal hypertension that develops in chronic hepatitis, which leads to the formation of an extensive network of portal anastomoses, including and in the lung tissue with increased hypoxia and hypoxemia.

One of the objectives of this study was to establish the association of HbF changes with the concentration of such important markers of endothelial dysfunction as the vasoconstrictor ET-1 and the adhesive protein von Willebrand factor. As can be seen from the data of Table 1, the studied markers of endothelial dysfunction were significantly increased when comparing the obtained results with the values of healthy children. Consequently, a comprehensive assessment of HbF levels, markers of endothelial dysfunction, in combination with clinical data, provides much more information about the development of tissue hypoxia and hypoxemia in sick children with combined pathology, and also allows an additional assessment of the severity of the pathological process in the liver.

It was found that in children with sensorineural hearing loss associated with hepatitis B, in 30.8% of cases there was a simultaneous increase in the concentration of HbF and a decrease in blood oxygen saturation of 64.82 ± 5.43 mm Hg against 76.05 ± 6.11 mm Hg, indicating moderate hypoxemia. The increase in tissue hypoxia was also indicated by the results of arterial-venous blood difference in the examined children. For a more reliable confirmation of this version, we studied the activity of lactate dehydrogenase and aspartate

Table 1. Blood biochemical parameters in children with hepatitis B associated hearing loss

<table>
<thead>
<tr>
<th>No</th>
<th>Indicators</th>
<th>Healthy children n = 12</th>
<th>Children sensory hearing loss combined CHB n = 26</th>
<th>Children sensory hearing loss without CHB n = 8</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Haemoglobin content g/l</td>
<td>158,62±8,02</td>
<td>92,05±6,54*</td>
<td>119,32±8,13</td>
</tr>
<tr>
<td>2</td>
<td>Fetal haemoglobin (HbF) content (g/l)</td>
<td>2,26±0,12</td>
<td>3,41±0,18*</td>
<td>2,36±0,19</td>
</tr>
<tr>
<td>3</td>
<td>pO₂ of arterial blood (mm Hg)</td>
<td>76,05±6,11</td>
<td>64,82±5,43*</td>
<td>72,43±6,32</td>
</tr>
<tr>
<td>4</td>
<td>pO₂ arterio-venous blood difference (mm Hg)</td>
<td>42,35±3,12</td>
<td>24,89±1,57*</td>
<td>38,75±2,87</td>
</tr>
<tr>
<td>5</td>
<td>The content of desquamated endothelial blood cells (x10⁴/l)</td>
<td>2,34±0,22</td>
<td>4,78±0,34*</td>
<td>3,01±0,26*</td>
</tr>
<tr>
<td>6</td>
<td>Endothelin 1-retention (fmol/l)</td>
<td>0,93±0,12</td>
<td>1,740±13*</td>
<td>0,96±0,12</td>
</tr>
<tr>
<td>7</td>
<td>Von Willebrand factor (%)</td>
<td>76,51±4,27</td>
<td>112,36±7,11*</td>
<td>91,76±6,34*</td>
</tr>
<tr>
<td>8</td>
<td>Activity lactate dehydrogenase (U/l)</td>
<td>576,12 ±11,3</td>
<td>1028,74 ±13.85*</td>
<td>788,63 ±12,34</td>
</tr>
<tr>
<td>9</td>
<td>Aspartate aminotransferase activity (U/l)</td>
<td>20,61 ±1,44</td>
<td>69,71 ±5,32*</td>
<td>32,28 ±2,52*</td>
</tr>
</tbody>
</table>

Note: * - significance of differences P <0.05
aminotransferase in the blood of the examined children. As can be seen from the obtained results of the research, the activity of the studied enzymes exceeded the initial indicators, respectively, by 59% and 3.3 times.

An indicator of the functional state of endotheliocytes is von Willebrand factor and endothelin-1 [15]. As can be seen from the presented research results, the level of endothelin-1 in children with the combined form of the disease was significantly higher in comparison with healthy children [16]. The level of activity of von Willebrand factor in the blood plasma was also significantly higher than in healthy children.

Under physiological conditions, the endothelium produces a number of vasodilating and vasoconstrictive substances that support the necessary level of vascular tone [17,18]. Numerous studies have shown that endothelin-1 is the most potent vasoconstrictor factor currently known. Proved that vascular endothelial is the main source of endothelin-1 in vivo.

The von Willebrand factor is a complex multidimensional adhesive glycoprotein synthesized by endothelial cells. Functionally, it is a carrier-stabilizer for a procoagulant protein that circulates in the blood serum as a non-covalently bound complex and is an adhesion protein in hemostasis processes. The von Willebrand factor can bind collagen and possibly other endothelial structures and mediate platelet adhesion to the subendothelium through the binding of the glycoprotein Ib surface platelet receptor. Therefore, an increase in the level of von Willebrand factor activity is an indicator of endothelial damage [19].

In this study, in children with sensorineural hearing loss associated with hepatitis B, the reaction of endothelial dysfunction indicators was detected - a significant increase in endothelin-1 level and von Willebrand factor activity in the blood plasma and their interrelation with the level of oxygen partial pressure in the blood, which indicates a violation of vasoconstrictor therapy and adhesive endothelial function in this pathology. This fact is explained by the fact that with this pathology in children there are favourable conditions for the development of endothelial dysfunction, on the background of hypoxia, as well as disruption of the metabolic function of the endothelium, which can lead to an increase in the content of various biologically active substances [20].

In addition, an important sign of endothelial dysfunction is a change in the phenotypic activity of endotheliocytes, which results in the cells losing anticoagulant properties and enhancing the production of coagulation factors [21]. When exposed to a damaging factor (hypoxia), leukocytes, monocytes, mononuclear phagocytes are activated, and damage and proliferation factors are produced: free radicals, interleukin-1, tumor necrosis factor a, tissue factor, trombocyte growth factor, and other biologists reported that active substances acting on endotheliocytes. In this situation, endotheliocytes begin to intensively secrete vasoactive and prosclerogenic substances (endothelin, etc.), the accumulation of which stimulates fibrotic changes and vascular remodeling [22].

Thus, an important pathogenetic role of endothelial dysfunction in children with sensorineural hearing loss associated with hepatitis B has been shown.

4. CONCLUSION

Firstly, dependence of the indices of partial oxygen in the blood and, to a greater extent, HbF, on the blood content of the vasoconstrictor endothelin-1, von Willebrand factor, indicates the pathogenetic significance of the leading markers of endothelial dysfunction in the development of tissue hypoxia in children with sensorineural hearing loss combined liver disease.

Secondly, a combination of sensorineural hearing loss with hepatitis B in children. Pulmonary hypertension is associated with endothelial dysfunction (increased endothelin-1 concentration and von Willebrand factor activity).

Thirdly, determining the level of HbF in children with CHT of combined HBV can be used to diagnose chronic tissue hypoxia and helps to clarify the severity of the pathological process, which allows predicting the progression of the disease and their complications.

CONSENT AND ETHICAL APPROVAL

As per university standard guideline parents consent and ethical approval has been collected and preserved by the authors.

COMPETING INTERESTS

Authors have declared that no competing interests exist.
REFERENCES


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