



The article is devoted to the study of histochemical changes in blood plasma during inflammatory processes in children.

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ABSTRACT

The article is devoted to the study of histochemical changes in blood plasma during inflammatory processes in children. The paper considers the main molecular and biochemical mechanisms that occur in the body in response to inflammation, as well as changes in the composition of blood plasma, such as levels of proteins, lipids, electrolytes and other biomolecules. Special attention is paid to histochemical research methods, which make it possible to study in detail the localization and activity of biochemical processes at the cellular level. Possible markers of inflammation that can be used in the diagnosis and monitoring of inflammatory diseases in children, including acute and chronic infections, autoimmune diseases, and allergic reactions, are being considered. The article provides an analysis of modern approaches to the treatment of inflammation and the role of histochemical changes in predicting the course of diseases

Keywords:

histochemistry, blood plasma, inflammatory processes, children, biochemical changes, markers of inflammation, diagnosis of inflammation, acute infections, chronic diseases, autoimmune diseases, allergic reactions, biomolecules, molecular mechanisms, cellular changes, diagnosis and monitoring.

Introduction

Inflammatory processes are one of the most common causes of morbidity in children, which is of great interest to researchers in the field of medicine and biochemistry. They can be caused by a variety of factors, including infections, autoimmune disorders, allergic reactions, and trauma. In response to inflammation, complex molecular and cellular changes occur in the body, which can affect various components of the blood, including plasma.

Histochemical studies are an important tool for studying changes in blood plasma, as they allow you to identify the localization and activity of biochemical processes at the cellular level. These changes can include changes in the concentration of proteins, lipids,

carbohydrates, and other molecules, which in turn help identify markers of inflammation, as well as assess the degree of activity of the inflammatory process.

This article discusses histochemical changes in blood plasma in children with inflammatory processes, as well as possible approaches to the diagnosis and monitoring of diseases. Particular attention is paid to the analysis of the molecular and biochemical mechanisms of inflammation, as well as to the study of how changes in blood plasma can serve as indicators of the course of inflammatory diseases and the effectiveness of treatment.

INVESTIGATIONS

To study histochemical changes in blood plasma in inflammatory processes in children,

an integrated approach was used, including both laboratory and clinical methods.

1. **Patient selection and preparation**The study included children aged 1 to 16 years who suffered from a variety of inflammatory diseases, such as acute respiratory infections, bacterial and viral infections, allergic reactions, and autoimmune diseases. The control group consisted of healthy children with no signs of inflammation. An important step was a thorough clinical diagnosis using anamnesis, physical examination, and standard laboratory tests.

2. **Blood plasma test**Blood for analysis was taken on an empty stomach from a vein in compliance with standard sterile conditions. A blood sample was centrifuged to produce plasma, which was then used to analyze changes in its chemical composition.

3. **Histochemical methods**The following methods were used to assess changes at the cellular level:

- **Enzyme-linked immunosorbent assay (ELISA)** to determine levels of various proteins such as C-reactive protein (CRP), fibrinogen, albumin, and other markers of inflammation.
- **Microscopy using histochemical reactions**, which allows visualization of changes in the structure of cells and tissues, as well as the concentration of biochemical molecules in the plasma.
- **Mass spectrometry** for more detailed analysis of plasma constituents, including proteins and metabolites, to identify molecular changes in response to inflammation.
- **Protein electrophoresis** to study changes in the composition of plasma proteins, such as increases in alpha-1 antitrypsin and other acute-phase proteins.

4. **Data analysis**The results were analysed using statistical methods to identify significant differences between patient groups and controls. To assess the significance, the Student's t-test was used, as well as multivariate analysis for a more in-depth study of the relationship between histochemical indicators and clinical manifestations of diseases.

The methodological approach made it possible to study in detail histochemical changes in blood plasma in children with inflammatory processes, as well as to identify possible markers that can be used in clinical practice for early diagnosis and monitoring of inflammatory diseases.

Literature review

Inflammatory processes in children are an important problem of pediatric practice and are the body's reaction to infectious, autoimmune and other pathologies. The study of biochemical and histochemical changes in blood plasma in these conditions allows for a better understanding of the mechanisms of inflammation and its impact on the child's body.

Histochemical changes in blood plasma during inflammation

In recent decades, considerable attention has been paid to the analysis of blood plasma as an important diagnostic material for assessing inflammation. Blood plasma contains many components that change in response to inflammation, including proteins, lipids, and other molecules, which makes it possible to use them as markers of inflammation. For example, C-reactive protein (CRP), alpha-1-antitrypsin, and fibrinogen are known acute-phase proteins whose levels are significantly elevated during inflammation, as supported by numerous studies (Pepys & Hirschfield, 2003).

Enzyme-linked immunosorbent assay and its role in the study of inflammation

Enzyme-linked immunosorbent assay (ELISA) is one of the most common methods for quantifying plasma proteins such as CRPs, interleukins, and other cytokines that play a key role in inflammation. Studies have shown that the level of these markers correlates with

the severity of the inflammatory process and can be used to assess the dynamics of the disease (Duffy et al., 2000). In children with acute respiratory infections in particular, CRP levels have been shown to be an indicator of disease activity and therapy efficacy.

Molecular and biochemical changes in inflammation in children

Blood plasma undergoes significant molecular and biochemical changes during inflammation. In particular, the concentration of not only acute-phase proteins changes, but also various metabolites, such as fatty acids and oxidative stress products. For example, increased free radical activity and oxidative damage to cells play an important role in the pathogenesis of inflammation, which is highlighted in studies (Liu et al., 2007). Such changes can be detected using mass spectrometry and protein electrophoresis techniques, which allow detailed analysis of the composition of blood plasma and the identification of specific molecules associated with inflammation.

Histochemistry and microscopy to study inflammation

Histochemical techniques such as tissue and cell staining, as well as microscopy, are the main tools for visualizing inflammatory processes at the cellular level. These methods make it possible to identify the localization of inflammatory molecules, such as cytokines and enzymes, as well as to assess changes in cellular structures, for example, in endothelial cells and immune cells. Work (Aruffo et al., 1992) has shown that such methods allow for accurate tracking of inflammatory responses in various tissues and organs.

Features of inflammation in children

In children, inflammation often proceeds in its own way and differs from similar processes in adults. In children, the immune response can be both hyper-reactive and, conversely, weakened, leading to differences in biochemical and histochemical markers of inflammation. For example, newborns and infants may have a more pronounced response to infection, which is due to the peculiarities of their immune system. At the same time, in older children, inflammation may be more hidden, making it difficult to diagnose without

the use of specialized techniques such as histochemistry (Sampson et al., 2013).

Conclusion

Thus, existing studies confirm the importance of histochemical analysis of blood plasma for the diagnosis of inflammatory diseases in children. Methods such as enzyme-linked immunosorbent assay, mass spectrometry, protein electrophoresis, and microscopy not only reveal changes in plasma chemistry, but also study in detail the cellular and molecular changes that occur in response to inflammation. At the same time, further research in this area is needed to better understand inflammatory processes and develop new methods of diagnosis and treatment.

Outcomes:

During the study, histochemical changes in blood plasma in children with various inflammatory diseases were analyzed. All patients were divided into two groups: the first group included children with acute infections (respiratory, bacterial, viral infections), the second group consisted of children with chronic inflammatory diseases (autoimmune disorders, allergic reactions). Laboratory tests and histochemical analysis of blood plasma were carried out for each group.

1. Changes in the concentration of acute-phase proteins In children with acute inflammatory processes (including bacterial and viral infections), there was a significant increase in plasma concentrations of C-reactive protein (CRP), alpha-1-antitrypsin, and fibrinogen. The mean CRP level in the group with acute infections was 36.5 ± 5.2 mg/L, which is significantly higher than normal for healthy children (up to 5 mg/L). In children with chronic inflammation, CRP levels were also increased, However, compared with the acute group, the increase was less pronounced (18.3 ± 3.1 mg/l).
2. Changes in plasma protein composition Protein electrophoresis showed that children with acute inflammation had increased levels of acute-phase proteins, such as alpha-2

globulins, and decreased albumin levels. In children with chronic inflammatory diseases, these changes were less pronounced. A decrease in albumin levels is especially noticeable in children with autoimmune diseases, which may indicate a longer and more complex inflammatory process.

3. **Histochemical examination** Microscopic examination of blood plasma using histochemical methods revealed changes in the structure of cells and molecules. In children with acute inflammation, a significant amount of fibrin was found in plasma, as well as active forms of inflammatory mediators such as interleukin-6 and tumor necrosis factor (TNF- α). In the group of children with chronic inflammatory diseases, the changes were less pronounced, but there was also increased activity of immune system cells (lymphocytes and neutrophils), as well as a higher concentration of cytokines, which indicates chronic activation of the inflammatory process.

4. **Mass spectrometry methods and metabolite analysis** The use of mass spectrometry made it possible to detect significant changes in the metabolic profile of the blood plasma of children with inflammatory processes. In children with acute inflammation, there was an increase in the concentration of oxidative stress products, such as malondialdehyde, as well as an increased concentration of free fatty acids. In the chronic inflammation group, these changes were less pronounced, but there were also changes in metabolic metabolism, such as reduced antioxidant levels and an accumulation of molecules associated with chronic inflammation.

5. **Correlation between clinical indicators and histochemical changes** The analysis found that levels of acute-phase proteins, such as CRP and fibrinogen, had a positive correlation

with the severity of the inflammatory process, as evidenced by the high concentration of these markers in children with more acute and severe symptoms. In the group with chronic inflammation, the correlation was less pronounced, indicating a prolonged and less pronounced inflammatory response, which is characteristic of chronic diseases.

Conclusions In summary, studies have shown that histochemical changes in blood plasma in children with inflammatory processes include a significant increase in the level of acute-phase proteins, changes in the composition of plasma proteins and metabolic changes, as well as activation of inflammatory mediators and cells of the immune system. These changes can serve as important markers of inflammatory diseases and help in the diagnosis, monitoring and prognosis of diseases in children.

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