

Dynamics of Morphometric and Echocardiographic Changes in Children Aged 1 to 3 Years with Congenital Heart Defects

Sadoqat Saidova

Bukhara State Medical Institute

Abstract: This article in the literature explains the causes and complications of congenital heart disease in children, comparative diagnosis. This article presents information on the frequency and risk of congestive heart failure in the sympathetic nervous system in children.

Keywords: heart, EXOKG examination, anthropometric indicators.

Introduction

For every 1,000 live births in Uzbekistan, there are between 5.5 and 15.7 children with congenital heart defects. In 50% of cases, congenital heart defects cause disability among all congenital defects, and thus are among the problems of social significance [1].

The aim of our research is to compare anthropometric parameters and echocardiographic examinations in children with congenital heart disease[2]. According to the European International Register of Birth Defects, congenital heart defects (CHD) are the most common group of developmental anomalies in children and remain the leading cause of death in newborns [3,4]. Currently, there is a tendency to increase the number and weight of CHD [6]. Currently, there is a growing trend in the number and severity of registered CHD [5].

This concept has several drawbacks. First, it does not take into account the time of onset of the defect, and second, a number of anomalies of the intrathoracic vessels do not belong to CHD (e.g., permanent superior vena cava), which is important during surgery [6]. Third, diseases such as cardiomyopathies and abnormalities of the cardiac conduction system ("Wolf-Parkinson-White phenomenon, long QT interval syndrome"), which lead to "structural abnormalities" and "functional disorders" [7], do not apply to CHD. Furthermore, the term CHD refers only to congenital defects of the thoracic regions of the great vessels (e.g., coarctation of the abdominal aorta). Although the concept of CHD is currently defined, it refers to anatomical deformity of the heart or large vessels developing in the uterus. 'can be rifled [8].

At present, CHD is the leading cause of death compared to other malformations in children and remains the leading cause of death [9]. In addition, a further increase in the prevalence of CHD is expected. The main reason may be related to the improvement of diagnostic techniques associated with the improvement of the skills of ultrasound diagnostics specialists and the improvement of modern imaging techniques [10]. The most intensive development of a child is observed in the first year of his life. During this period, a significant increase in body weight and height is observed, and the functional activity of the central nervous system improves. Proper growth and weight gain are closely related to the development of functional abilities of organs and systems. Due to hemodynamic disturbances, CHD has a direct negative impact on the development of the child. Some types of CHD are associated with a sharp decline in quality of life, an increase in the number of chronic diseases, and the formation of delayed neuropsychic

development (CHD). The authors also consider congenital defects of the heart to be a cessation of development at a certain stage of ontogeny, which corresponds to a particular stage of phylogeny. Within these theories, only atavistic heart defects (female and neutral) are compatible, and the whole group of male defects cannot be explained because none of the male components of congenital heart defects are compatible with normal embryonic or similar formation.

At present, CHD is the leading cause of death compared to other malformations in children and remains the leading cause of death. The main reason may be related to the improvement of diagnostic techniques associated with the improvement of the skills of ultrasound diagnostics specialists and the improvement of modern imaging techniques. Congenital defects are explained by the cessation of cardiac development at different stages of ontogeny; he interprets them as a return to one of the stages of phylogeny. The authors synthesize the previous two views, considering congenital heart defects as cessation of development at a certain stage of ontogeny, which corresponds to this or that stage of phylogeny. Dividing congenital heart defects and large vessels into males, females, and neutrals allows the patient's gender to be used as a diagnostic symptom. However, the male and female types of defects have a very large value of the coefficient of diagnostic value. For example, given the patient field data, the probability of diagnosis in a patent ductus arteriosus is 1.32 times higher.

The study of the effects of external factors on the cardiovascular system is a current problem of applied medicine, which is confirmed by many modern studies aimed at studying the mechanisms of development of congenital heart defects, but the specificity of the types of response to external factors does not provide extensive confirmation of the findings obtained in clinical studies. As a result of our research, we divided patients into age groups and performed anthropometric changes and echocardiographic comparisons in children with congenital heart defects. Congenital heart defect - a permanent defect, deficiency and change in the anatomical structure of the heart; interferes with normal blood flow. Congenital and acquired heartworm are different. Congenital heart disease occurs as a result of malformation of the fetal heart and large heart vessels during embryonic development. Poisoning of the mother's body in the early stages of pregnancy, suffering from certain diseases, biological effects of ionizing radiation, hereditary diseases, etc. k. causes. In infancy (up to 1 year of age), incomplete development of the cardiovascular system (e.g., incomplete opening of arterial pathways or oval foramen) is also considered a heart defect. The most common types of congenital heart defect are: abnormal pathways in various combinations between large and small circulatory circles, as well as the presence of narrowed or clogged areas in the major arteries of the heart (e.g., pulmonary artery and aorta) or misalignment of these vessels; mixed powders; defects in the number and structure of the heart chambers. Depending on the degree to which the arterial and venous blood is mixed, some congenital heart defects pass with cyanosis (blue powders), some without cyanosis (white powders). It depends on which direction the blood flows (in the direction of the shunt), the degree of pressure rise in the pulmonary artery, and the condition of the heart muscle through the improper holes that connect the large and small circulatory circuits. Symptoms of congenital heart disease include physical abnormalities, paleness or bruising, shortness of breath, changes in heart size and condition, heart murmurs, and more The aim of the study: To study the anthropometric parameters and comparative features of echocardiographic changes in the heart in children born with congenital heart disease (1-3 years).

MATERIALS AND METHODS

The study was conducted at the Bukhara Regional Multidisciplinary Children's Hospital. It was conducted on the basis of bilateral agreements of Bukhara State Medical Institute. Children were divided into 3 groups: group of children aged 1 years (n = 20); The results of the examination of the group of children aged 2 years (n = 20) and the group of children aged 3 years (n = 10) were studied. The methodology of anthropometric study of children was used to conduct anthropometric measurements (Methodological recommendations on the morphometric features

of the assessment of physical development of children and adolescents //N.H. Shomirzaev, S.A. Ten and I.Tukhtanazarova, 1998). Anthropometric research included height, body weight, body length, and chest circumference measurements. Echocardiographic examinations obtained the results of ultrasound anatomy of the heart. The study was conducted on a SONOACE R3-RUS device with linear (7.5 MHz) and convex (3.5 MHz) transducers. In this study, the linear dimensions of each part of the heart, the thickness and volume of the heart were studied using the formula of J. Brunn and co-authors (1981): $V = K \cdot [(L1 \cdot W1 \cdot T1) + (L2 \cdot W2 \cdot T2)]$, where V - gland volume index (cm³), K - coefficient equal to 0.479; L, W, T - length, width and thickness of each piece of cloth. Mathematical processing was performed directly from the Excel 7.0 general data matrix using the capabilities of STTGRAPH 5.1, standard deviation indicators and representation error were detected.

Results

Studies have shown that in children 1 year of age, height ranges from 65.2 sm to 77.5 sm, with an average of 70.1 ± 0.9 sm, and in children 2 years of age from 70.4 sm to 78.3 sm, on average 75.2 ± 0.4 sm, children 3 years were found to have an average height of 82.1 ± 0.2 sm from 76.0 sm to 87.4 sm.

In children 1 year of age, body weight ranged from 3.4 kg to 7.2 kg, with an average of 4.1 ± 0.9 kg, and in children 2 years of age ranged from 7.5 kg to 10.2 kg, with an average of 8.5 ± 0.6 kg, 3 years of age averaged 10.3 ± 0.4 to 9.0 kg to 12.1 kg.

Table № 1. Indicators of physical development of children from 1 to 3 years in the study

№	Indicators	Children 0-3 years (n = 30)		
		Children 1 years old (n = 20)	2-years old children (n = 20)	3 -years old (n = 20)
1	Height, sm	$70,1 \pm 0,9$	$75,2 \pm 0,4$	$82,1 \pm 0,2$
2	Body weight, kg	$4,1 \pm 0,9$ кг	$8,5 \pm 0,6$	$10,0 \pm 0,4$
3	Chest circumference, sm	$40,2 \pm 0,6$	$43,4 \pm 0,5$	$45,2 \pm 0,8$
4	Abdominal circumference, sm	$39,0 \pm 0,4$	$42,3 \pm 0,5$	$45,4 \pm 0,6$

Note: * - reliability level $p \leq 0.05$ compared to the previous group

Echocardiographic parameters of the heart in children from 1 to 3 years Right atrium from 8.1 mm to 10.2 mm in children 1 year, average - 9.15 ± 0.1 mm, pulmonary artery width in children of the same age from 9.2 mm to 11.0 mm, average 10.2 ± 0.1 to 2 years of age to right atrium 8.5 mm 15.1 ± 0.1 mm on average 16.05 mm, pulmonary artery width 11.2 mm to 13.3 mm 12 in children of the same age 3 ± 0.3 mm. In children aged 2 to 3 years, the right atrium 10.1 from 16.2 mm to , average 18.2 ± 0.3 mm, and the pulmonary artery width averaged 13.2 ± 0.3 to 12.1 mm to 14.5 mm reaches.

Table №2. Comparative features of echocardiographic parameters in children from birth to 3 years

№	Indicators	Children 1-3 years (n = 75)		
		Children 1 years old (n = 20)	Children 2 years old (n = 20)	Children 3 years old (n = 20)
1	Right atrium , mm	$9,5 \pm 0,1$	$10,2 \pm 0,1$	$18,2 \pm 0,3$
2	Pulmonary artery width, mm	$10,2 \pm 0,1$	$12,3 \pm 0,3$	$13,2 \pm 0,3$

Note: * - reliability level $p \leq 0.05$ compared to the previous group

Anthropometric studies among children born with congenital heart defects 1 year of age showed that children born with interventricular septal defect had a height gain of 0.9 sm higher than children born with interventricular septal defect, and children weighed 0.63 kg more than children of the same age. detected. In children aged 2 years, children born with interventricular septal defect were found to have a height of 0.4 sm compared to children born with interventricular septal defect. Children of the same age had a body weight of 0.6 kg. Children born with interventricular septal defect in children aged 3 years were found to be 0.3 sm taller than children born with interventricular septal defect, and to have a body weight of 0.3 kg more than children of the same age. In children born with congenital heart disease 1 year of age, the circumference of the thoracic circumference was 0.6 sm higher than in children born with interventricular septal defect, and in children of the same age the abdominal circumference was 0.40 sm higher. Children born with congenital heart disease with congenital heart failure Children with congenital heart disease 0.5 cm higher than in newborns and 0.5 sm higher in children of the same age. Children born with congenital heart disease between the ages of 3 years, 8.0 sm, and in children of the same age, the abdominal circumference was found to be 0.6 sm higher.

Conclusion

According to the data obtained, children born with congenital heart disease 1 year, 2 years, 3 years with congenital heart failure, compared with children born with interventricular septal defect (height, body weight, chest circumference, abdominal circumference) was found to be high. Echocardiographic examination revealed that children born with congenital heart defects 1-st years of age, 2 years of age, 3 years of age, children with congenital heart failure, children born with interventricular septal defect (aortic width, pulmonary artery width) had lower body weight and height. found to be consistent with growth rates.

References

1. Kurbanovna, S. I. Functioning of the Immune System in Children, After Surgical Correction of Congenital Heart Defects. *European Journal of Life Safety and Stability* (2660-9630), 12, 439-446. (2021).
2. Abdurakhmanov, Z.M., Yemets, I.N. Long-term outcomes of aortic valve repair in children with congenital heart disease and their predictors. *Russian Journal of Cardiology*, 25 (8), (2020) DOI: 10.15829/1560-4071-2020-3971.
3. Saidova, S. Y. (2021). A study regarding revealing echocardiographic and anthropometric changes in children from birth to 3 years old with congenital heart defects. *ACADEMICIA: An International Multidisciplinary Research Journal*, 11(10), 395-399.
4. Saidova, S. Y. Revealing echocardiographic and anthropometric changes in children from birth to 3 years old with congenital heart defects. *ACADEMICIA: An International Multidisciplinary Research Journal*, 11(9), 1071-1075. (2021).
5. Abdurakhmanov, Z.M., Yemets, I.N. Long-term outcomes of aortic valve repair in children with congenital heart disease and their predictors. *Russian Journal of Cardiology*, 25 (8), DOI: 10.15829/1560-4071-2020-3971. (2020)
6. Muzaffarovn, K. S., Radjabovich, B. R., & Pulatovna, A. N. Characteristics of anthropometric parameters of the upper extremities in children and adolescents with scoliosis. *Open Access Repository*, 4(3), 1211-1215. (2023).
7. Malik, A., Yoshida, Y., Erkin, T., Salim, D., & Hamajima, N. Hypertension-related knowledge, practice and drug adherence among inpatients of a hospital in Samarkand, Uzbekistan. *Nagoya Journal of Medical Science*, 76(3-4), 255-263 (2014).

8. Yo`ldoshovna, S. S. Exocardiographic Parameters in Boys Born With Congenital Heart Defect and Healthy Under 3 Years of Age. JOURNAL OF HEALTHCARE AND LIFE-SCIENCE RESEARCH, 2(3), 22-25. (2023).