



# Evaluation of Heart Rate Variability in Children with Congenital Heart Diseases before and after Surgical Repair

Nashwa Ibrahim Ebied <sup>a\*</sup>, El-Atafy El-Metwally El-Atafy <sup>b</sup>,  
Walid Ahmed El-Shehaby <sup>a</sup> and Amr Mohamed Zoair <sup>a</sup>

<sup>a</sup> Pediatrics Department, Faculty of Medicine, Tanta University, Tanta, Egypt.

<sup>b</sup> Cardiothoracic Surgery Department, Faculty of Medicine, Tanta University, Tanta, Egypt.

## Authors' contributions

*This work was carried out in collaboration among all authors. All authors read and approved the final manuscript.*

## Article Information

DOI: 10.9734/AJPR/2022/v10i4200

## Open Peer Review History:

This journal follows the Advanced Open Peer Review policy. Identity of the Reviewers, Editor(s) and additional Reviewers, peer review comments, different versions of the manuscript, comments of the editors, etc are available here: <https://www.sdiarticle5.com/review-history/93114>

**Original Research Article**

**Received: 20/08/2022**

**Accepted: 27/10/2022**

**Published: 10/11/2022**

## ABSTRACT

**Background:** Surgical repair of congenital heart diseases (CHD) affects heart rate variability (HRV) in different ways. It leads to varying degrees of cardiac autonomic derangement, clinically detectable as depression of HRV parameters. The aim of this work was to evaluate HRV in children with CHDs before and after surgical repair to detect the effect of CHDs (acyanotic or cyanotic) and cardiac surgery on cardiac autonomic function.

**Methods:** This observational case-control study enrolled thirty cases with CHD. Cases were divided in two equal groups: group1 included children with CHDs (not repaired) before surgical intervention and group 2 included children after surgical repair of CHDs. Another fifteen healthy children, matched for age and sex, were enrolled as a control group. All children in this study were subjected to full history taking, clinical evaluation, Holter monitor, twenty-four-hour ambulatory electrocardiographic recordings and the seven time-domain indices of HRV.

\*Corresponding author;

**Results:** There was significant increase of HR in patients before surgical repair as compared to patients after surgical repair as compared to control group. There was significant decrease of PNN50, r MMSD, SDNN index and SDANN of children before surgical repair as compared to children after surgical repair as compared to controls. There was a negative significant correlation between HR and age before surgery, after surgery there was a negative significant correlation between HR and age, pNN50, rMMSD, SDNN index and SDANN, in control there was negative significant correlation with age, SDNN index, SDANN and SDNN.

**Conclusion:** HRV parameters increased significantly in children with CHDs after surgical repair as compared to these before surgical repair denoting effect of cardiac surgery on cardiac autonomic function.

*Keywords: Heart rate variability; congenital; heart diseases; surgical repair.*

## 1. INTRODUCTION

Congenital Heart Diseases (CHDs) remain the main cause of illness and death in children and the most prevalent human birth defect, affecting about one percent of all live births globally [1].

CHDs are divided into cyanotic and acyanotic heart diseases. In general, acyanotic CHDs are more common than cyanotic group with bicuspid aortic valve disease being the most familiar kind of CHD [2]. In order to adjust for the hemodynamic changes introduced by illness, the autonomic regulation of the heart rate (HR) and of circulation might exhibit pathologic behaviour [3].

Earlier investigations have focused on heart rate variability (HRV) in certain cardiac abnormalities like tetralogy of Fallot. The purpose of the research was to evaluate HRV in young patients diagnosed with CHDs either cyanotic or acyanotic abnormalities, comparing them to healthy control of the same age. The study's findings revealed that infants with congenital heart disease exhibit more complicated HRV and sympathetic overactivity, both of which may be an attempt to compensate for hemodynamic change [4].

Variability in heart signal of two sequential beats is called HRV. Short and long-term variability reflect autonomic nervous system function, so that increased or decreased HRV is an indicator of human health [5].

As a good indicator of cardiac autonomic nervous system activity, HRV has found widespread use. The various physiological elements that affect the heart's regular beat are accurately reflected in it. Actually, they are a great tool for studying the balance between the

sympathetic and parasympathetic branches of the nervous system [6].

Surgical repair of CHDs affects HRV in different ways. Depressed HRV values are a clinical marker of cardiac autonomic dysfunction [7].

Few studies indicate that surgical aortic valve replacement may result in HRV problems. Replacement of the aortic valve surgically has a strong depressive effect on a number of cardiac autonomic measures, while less invasive techniques allow for a better presentation of HRV [8].

Changes in cardiac autonomic regulation may influence the long-term prognosis of patients who undergo univentricular heart repair. Compared to bidirectional superior Cavo pulmonary anastomosis with antegrade pulmonary blood flow, whole Cavo pulmonary connection results in a considerable decrease in global cardiac autonomic tone [9].

The aim of this work was to assess HRV in children with CHDs before and after surgical repair to detect the effect of CHDs (acyanotic or cyanotic) and cardiac surgery on cardiac autonomic function.

## 2. PATIENTS AND METHODS

This observational case-control study enrolled thirty children with CHDs. Cases were selected from Pediatric Cardiology Unit, pediatric Department and Cardio thoracic Surgery Department, Tanta University Hospital (TUH) between June 2018 to June 2020.

**The inclusion criteria** were children in pediatric age with CHDs (acyanotic or cyanotic) before and after surgical repair.

The exclusion criteria were hepatic diseases, renal disorders or failure, pulmonary diseases, or acute or chronic disease that can affect autonomic nervous system like diabetes mellitus.

Cases were assigned into two equal groups: group1 included children with CHDs (not repaired) before surgical intervention and group 2 included children after surgical repair of CHDs. Fifteen healthy control children, matched for age and sex, were included.

**All children were subjected to the following:** providing the full history including birth, personal, feeding, developmental, family history of cardiovascular events, any complaints or drug intake, diet, and physical activity.

**Clinical Evaluation:** General, regional, and systemic examination as body weight, height, and vital signs such as HR, respiratory rate, and blood pressure. Cardiac examination including inspection, palpation, percussion, and auscultation for detection of cardiomegaly and evidence of murmur was performed.

**Investigations:** Plain X-ray heart and chest, ECG and Echocardiography and Doppler to detect and evaluate CHDs.

**Holter monitor:** Each kid in this research will undergo 24-hour Holter monitoring with ECG recording and interpretation, and heart rate variability (HRV) analysis (before and after cardiac surgery). Mathematical analysis and the assessment of HRV based on computer systems can help with the diagnosis of patient's cardiac status when coupled with 24-hour ambulatory electrocardiographic recordings and the seven time-domain (SDANN, SDNN, SDNN index, rMSSD, SD, PNN50, and mean RR) indices of HRV.

## 2.1 HRV Parameters Measured during 24 Hours with Holter Electrocardiogram [10]

Calculating the 24-hour standard deviation of sinus RR intervals (SDNN). Variation in the mean normal sinus RR intervals over all 5-mm segments (SDANN). Normal sinus RR interval standard deviation across all 5-minute intervals (SDNN index). Sum of squares of the differences in RR intervals produced by regular sinus waves (rMSSD). How often the RR gap between heartbeats is more than 50 milliseconds (pNN50).

## 2.2 Statistical Analyses

Using SPSS software, the acquired data were organised, tabulated, and statistically analysed (Statistical Package for the Social Sciences, version 21, SPSS Inc. Chicago, IL, USA). The range, mean, and standard deviation were determined for quantitative data. Boxplots were created to depict the quantitative data's median, first and third quartiles. Kruskal-Wallis ( $\chi^2$  value) was computed to compare more than two means of non-parametric data, and the Mann-Whitney U test was used to compare each pair of means if the  $\chi^2$  value was significant. Pearson's correlation coefficient was used to assess the relationship between variables ( $r$ ). The significance level of  $p < 0.05$  was selected for interpreting the findings of significance tests.

## 3. RESULTS

Regarding HR of the studied children with CHDs (before and after surgical intervention) and the control group, there was a considerable increase of HR in patients before surgical repair as compared to patients after surgical repair as compared to control group.

**Table 1. Clinical diagnosis of CHDs (before and after surgical intervention) groups**

Clinical diagnosis	CHD (n=30)	
	N	%
VSD	10	33.3
ASD	5	16.7
PDA	4	13.3
PFO	2	6.7
TOF	5	16.7
PA	4	13.3

VSD: Ventricular septal defect, ASD: Atrial septal defect, PDA: Patent ductus arteriosus, PFO: Patent foramen ovale, TOF: Tetralogy of Fallot, PA: Pulmonary atresia

**Table 2. HR of the studied children with CHDs (before and after surgical intervention) and the control group (n=45)**

<b>HR</b>			
Range	96-159	78-154	74-164
Mean±SD	133.00±25.88	120.00±21.89	94.00±28.22
$\chi^2$ values	9.185		
<b>P</b>	0.010*		
<b>Z value</b>	G1 vs G2 & G3, P=112, 0.013*		
<b>P</b>	G2 vs G3, P=0.019*		

$\chi^2$  values of Kruskal Wallis test, Z value of Mann-Whitney U test

Regarding pNN50 and rMSSD of the studied children with CHDs, there was significant decrease in children before surgical repair as compared to children after surgical repair as compared to control group.

**Table 3. pNN50 and rMSSD of the studied children with CHDs (before and after surgical intervention) and the control group (n=45)**

<b>pNN50 (%)</b>	<b>Children with CHDs (n=30)</b>		<b>Control children (n=15)</b>
	<b>Before surgical intervention (Not repaired) (n=15)</b>	<b>After surgical repair (n=15)</b>	
<b>pNN50</b>			
Range	0-2.80	0-17.50	0-214
Mean±SD	0.74±0.93	1.80±5.52	38.73±63.26
$\chi^2$ values	12.773		
<b>P</b>	0.002*		
<b>rMSSD</b>			
Range	1.00-32.00	3.00-61.00	10.00-88.00
Mean±SD	13.10±9.88	16.60±16.68	52.20±27.38
$\chi^2$ values	11.996		
<b>P</b>	0.002*		
<b>Z value</b>	G1 vs G2 & G3, P=0.820*, 0.002*		
<b>P</b>	G2 vs G3, P=0.005*		

Regarding SDNN index, SDANN and SDNN, there was significant decrease in children before surgical repair as compared to children after surgical repair as compared to control group.

**Table 4. SDNN index, SDANN, SDNN of the studied children with CHDs (before and after surgical intervention) and the control group (n=45)**

	<b>Children with CHDs (n=30)</b>		<b>Control children (n=15)</b>
	<b>Before surgical intervention (Not repaired) (n=15)</b>	<b>After surgical repair (n=15)</b>	
<b>SDNN index</b>			
Range	5.00-40.00	3.00-61.00	21.00-100.00
Mean±SD	18.10±12.02	16.60±16.68	65.90±32.21
$\chi^2$ values	13.750		
<b>P</b>	0.001*		
<b>Z value</b>	G1 vs G2 & G3, P=0.733*, 0.001*		
<b>P</b>	G2 vs G3, P=0.002*		

	Children with CHDs (n=30)		Control children (n=15)
	Before surgical intervention (Not repaired) (n=15)	After surgical repair (n=15)	
<b>SDANN</b>			
Range	14.00-38.00	0.00-87.00	34.00-159
Mean±SD	25.00±10.00	31.30±24.58	94.20±40.16
$\chi^2$ values	15.377		
P	0.0001*		
Z value	G1 vs G2 & G3, P=0.650, 0.0001*		
P	G2 vs G3, P=0.002*		
<b>SDNN:</b>			
Range	5.00-50.00	10.00-116.00	48.00-180.00
Mean±SD	30.20±15.60	34.50±29.71	117.60±47.81
$\chi^2$ values	17.503		
P	0.0001*		
Z value	G1 vs G2 & G3, P=0.733, 0.0001*		
P	G2 vs G3, P=0.001*		

Regarding correlation between HR and age and findings of Holer Monitor among the studied children, there was significant negative correlation between HR and age before and after surgery. pNN50, rMMSD, SDNN index and SDANN, in control there was negative significant correlation with age, SDNN index, SDANN and SDNN.

**Table 5. Correlation between HR and age and findings of Holer Monitor among the studied children with CHDs (before and after surgical intervention) and the control group (n=45)**

Variables	Correlation between HR and other variables among children with CHDs and control group (n=45)					
	Children with CHDs (n=30)			Control children (n=15)		
	Before surgical intervention (Not repaired) (n=15)		After surgical repair (n=15)			
	r	P	r	P	r	P
Age	-0.716	0.020*	-0.792	0.006*	-0.725	0.018*
pNN50	-0.212	0.557	-0.678	0.031*	-0.408	0.241
rMMSD	-0.160	0.659	-0.728	0.017*	-0.630	0.051
SDNN index	-0.537	0.109	-0.860	0.001*	-0.814	0.004*
SDANN	0.510	0.1032	-0.723	0.018*	-0.720	0.019*
SDNN	0.195	0.586	-0.607	0.063	-0.766	0.006*

#### 4. DISCUSSION

HRV is a physiological characteristic that distinguishes vegetative modulation of the heart rhythm. The relationship between a drop in HRV in patients with acute myocardial infarction (AMI) and an increase in mortality in this group of patients sparked an interest in HRV research more than four decades ago. Currently, HRV

analysis is extensively researched in clinical practise to identify the effect of the parasympathetic and sympathetic systems on sinus node function, as well as the possibility of cardiovascular problems and mortality [11].

In agreement with our study, Bakari et al. [12] studied 28 individuals with ASD between the ages of 4.5 and 8.7 years and 32 healthy

control children of the same age without hemodynamic problems underwent HRV analysis in the time and frequency ranges. All HRV measures showed to be lower in children with ASD compared to those in good health.

In a research including 39 patients who had had the Fontan operation, Butera et al. [13] found a substantial drop in HRV in the context of a reduced vagal tone. Davos et al. [14] shown that autonomic nervous system dysfunction lasted for an extended period of time after Fontan surgery. Extracorporeal circulation may also result in an imbalance of the autonomic nervous system (ANS) owing to the release of catecholamines, partial vagal denervation resulting from surgical injury, or ischemia. Massin and von Bernuth et al. [15] found reduction in HRV time and frequency domain in CHDs cases. This decrease was connected with the functional class of heart failure as determined by the New York Heart Association, but not with any particular hemodynamic data. Massin et al. and Finley et al. [15,16] observed HRV parameters to be considerably diminished in children with ASD. These results indicate either an increase in sympathetic heart control or a reduction in parasympathetic heart control.

In several pediatric ASD studies as mentioned by Horner and Bialkowski et al. [17,18] the HRV values are reduced as compared with those of controls, but the values increased after defect closure.

Edwards et al. [19] observed that atrial stress is a key stimulator of atrial natriuretic peptide release. Similar to neurohumoral variables, these data indicate a potential favourable impact of lower volume and pressure on HRV. When the consequences of right atrial dilatation and tension remain for an extended length of time, ASD patients develop atrial arrhythmias [20].

Attie et al. [21] mentioned that atrial tachyarrhythmias, thromboembolic events and heart failure are prevalent and contribute to cardiac morbidity and death in individuals.

In contrary, Hami and Corcia et al. [22] showed that there was no considerable variation regarding the HR average in children who exposed to surgery compared to controls.

Came in line with our findings, Özyılmaz et al. [23] reported significant decrease of HRV parameters of children with CHDs before surgical

repair as compared to these after surgical repair. In contrary, Hami and Corcia et al. [22] showed that no considerable change was recorded concerning pNN50 in children who exposed to surgery compared to controls.

In the current study, we found that rMSSD value did not differ in children with CHD after surgery, but it was lower in comparison to control group.

Bialkowski et al. [24] mentioned The SDNN value is more altered by parasympathetic and sympathetic impulses associated with vagal tonus than the rMSSD and pNN50 (%) values in the temporal domain study of HRV in humans.

Koca et al. [12] mentioned that Short-term HRV is represented by the HF component of frequency domain analysis and the pNN50 (%) and rMSSD values of time domain analysis (high frequency power).

Similarly, Massin and Von Bernuth et al. [25] Long-term HRV (low frequency power) is expressed by the LF component of frequency domain analysis and the SDNN, SDANN parameters of time domain analysis and SDNN index, and both are regulated by adrenergic and cholinergic activities.

In Özyılmaz et al. [23] study, The HRV values of patients who received transcatheter ASD closure at the sixth month resembled those of the control group.

In contrary, Hami and Corcia et al. [22] showed that no considerable variation was observed regarding rMSSD in children who exposed to surgery compared to controls.

Comparable to our results, Baevskii et al. [26] observed that In the sick group, the SDNN parameter was 103.0 - 50.6, compared to 138.7 - 36.2% in the controls.

In the study of Cansel et al. [27] pNN50 (percent) and rMSSD values were found to be lower in ASD patients, however this difference was not statistically significant. The HRV values in the time domain analysis rose after the procedure when compared to the initial levels, and they neared the levels of the control group when compared to those obtained before to the surgery. One day following the surgery, the SDNN, pNN50 (%), and SDANN values rose statistically significantly. The lack of change in

the rMSSD, NN, and SDNN index values may be related to the fact that the measurements were collected shortly after the surgery.

In contrary, Hami and Corcia et al. [22] showed that no considerable change was recorded concerning SDANN in children who exposed to surgery compared to controls.

In Özyılmaz et al. [23] study, ASD patients had lower rMSSD, SDNN, SDNN index, pNN50 (%) and SDANN values than the control subjects one day prior to ASD closure.

In contrary, Hami and Corcia et al. [23] showed statistically insignificant difference in SDNN in studied group compared to controls.

Literature revealed age-related variations in HRV as a result of ANS development, but occasionally contradictory findings. Some writers report a steady decline in sympathetic activity in the first decade of life [10], but others claim an increase in parasympathetic activity in the same period, as shown by elevated values of all time domain indices, followed by a gradual drop [28].

## 5. CONCLUSIONS

HRV parameters increased significantly in children with CHDs (acyanotic and cyanotic) after surgical repair as compared to these before surgical repair denoting effect of cardiac surgery on cardiac autonomic function.

## CONSENT AND ETHICAL APPROVAL

The cases' parents or legal guardians provided informed consent. The Ethics Committee, Faculty of Medicine, Tanta University had approved the study.

## COMPETING INTERESTS

Authors have declared that no competing interests exist.

## REFERENCES

1. Bruneau BG, Srivastava D. Congenital heart disease: Entering a new era of human genetics. *Circulation Research*. 2014;114:598-9.
2. Ratti C, Veronesi B, Grassi L, Bompani B. [Congenital heart diseases in clinical practice]. *Recenti Prog Med*. 2012;103: 213-7.

3. Aletti F, Ferrario M, de Jesus TB, Stirbulov R, Silva AB, Cerutti S, et al. Heart rate variability in children with cyanotic and acyanotic congenital heart disease: analysis by spectral and non linear indices. *Annu Int Conf IEEE Eng Med Biol Soc*. 2012;4189-92.
4. Acharya R, Kannathal N, Krishnan S. Comprehensive analysis of cardiac health using heart rate signals. *Physiological Measurement*. 2004;25:1139.
5. Acharya R, Lim C, Joseph P. Heart rate variability analysis using correlation dimension and detrended fluctuation analysis. *Itbm-Rbm*. 2002;23:333-9.
6. Silvilairat S, Wongsathikun J, Sittiwangkul R, Pongprot Y, Chattipakorn N. Heart rate variability and exercise capacity of patients with repaired tetralogy of Fallot. *Pediatr Cardiol*. 2011;32:1158-63.
7. Compostella L, Russo N, Compostella C, Setzu T, D'Onofrio A, Isabella G, et al. Impact of type of intervention for aortic valve replacement on heart rate variability. *International Journal of Cardiology*. 2015; 197:11-5.
8. Compostella L, Russo N, D'Onofrio A, Setzu T, Compostella C, Bottio T, et al. Abnormal heart rate variability and atrial fibrillation after aortic surgery. *Rev Bras Cir Cardiovasc*. 2015;30:55-62.
9. Madan K, Garg P, Deepak KK, Talwar S, Airan B, Choudhary SK. Heart rate variability in patients undergoing univentricular heart repair. *Asian Cardiovasc Thorac Ann*. 2014;22: 402-8.
10. Rajendra Acharya U, Paul Joseph K, Kannathal N, Lim CM, Suri JS. Heart rate variability: a review. *Medical and biological engineering and computing*. 2006;44: 1031-51.
11. Shvartz VA, Kiselev AR, Bockeria OL. Heart rate variability in atrial septal defect both before and after operation. *Cor et Vasa*; 2018.
12. Koca B, Bakari S, Öztunç F. Correlation among the heart rate variability indices in healthy children and those with atrial septal defect. *Turk Kardiyol Dern Ars*. 2013;41: 193-8.
13. Butera G, Bonnet D, Sidi D, Kachaner J, Chessa M, Bossone E, et al. Patients operated for tetralogy of Fallot and with non-sustained ventricular tachycardia have reduced heart rate variability. *Herz*. 2004; 29:304-9.

14. Davos CH, Francis DP, Leenarts MF, Yap SC, Li W, Davlouros PA, et al. Global impairment of cardiac autonomic nervous activity late after the Fontan operation. *Circulation*. 2003;108(Suppl 1):li180-5.
15. Massin M, Von Bernuth G. Clinical and haemodynamic correlates of heart rate variability in children with congenital heart disease. *European Journal of Pediatrics*. 1998;157:967-71.
16. Finley JP, Nugent S, Hellenbrand W, Craig M, Gillis D. Sinus arrhythmia in children with atrial septal defect: an analysis of heart rate variability before and after surgical repair. *Heart*. 1989;61:280-4.
17. Białkowski J, Karwot B, Szkutnik M, Sredniawa B, Chodor B, Zeifert B, et al. Comparison of heart rate variability between surgical and interventional closure of atrial septal defect in children. *Am J Cardiol*. 2003;92:356-8.
18. Horner SM, Murphy CF, Coen B, Dick DJ, Harrison FG, Vespalcova Z, et al. Contribution to heart rate variability by mechanoelectric feedback. Stretch of the sinoatrial node reduces heart rate variability. *Circulation*. 1996;94:1762-7.
19. Edwards BS, Zimmerman RS, Schwab TR, Heublein DM, Burnett JC, Jr. Atrial stretch, not pressure, is the principal determinant controlling the acute release of atrial natriuretic factor. *Circ Res*. 1988;62:191-5.
20. Webb G, Gatzoulis MA. Atrial septal defects in the adult: recent progress and overview. *Circulation*. 2006;114:1645-53.
21. Attie F, Rosas M, Granados N, Zabal C, Buendía A, Calderón J. Surgical treatment for secundum atrial septal defects in patients >40 years old. A randomized clinical trial. *J Am Coll Cardiol*. 2001;38:2035-42.
22. Hami K, Corcia M. Heart Rate Variability Modifications after Surgery for Congenital Heart Disease in Young Patients. *Evid Based Med Pract*. 2017;3:1-3.
23. Özyılmaz İ, Ergül Y, Tola HT, Saygı M, Öztürk E, Tanıdır İ C, et al. Heart rate variability improvement in children using transcatheter atrial septal defect closure. *Anatol J Cardiol*. 2016;16:290-5.
24. Białkowski J, Karwot B, Szkutnik M, Sredniawa B, Chodor B, Zeifert B, et al. Comparison of heart rate variability between surgical and interventional closure of atrial septal defect in children. *The American Journal of Cardiology*. 2003;92:356-8.
25. Massin M, von Bernuth G. Normal ranges of heart rate variability during infancy and childhood. *Pediatr Cardiol*. 1997;18:297-302.
26. Baevsky R, Kirillov O, Kletskin S. Mathematical analysis of heart rate changes under stress. Moscow: Nauka. 1984.
27. Cansel M, Yagmur J, Ermis N, Acikgoz N, Taşolar H, Atas H, et al. Effects of transcatheter closure of atrial septal defects on heart rate variability. *J Int Med Res*. 2011;39:654-61.
28. Silvetti MS, Drago F, Ragonese P. Heart rate variability in healthy children and adolescents is partially related to age and gender. *International journal of cardiology*. 2001;81:169-74.

© 2022 Ebied et al.; This is an Open Access article distributed under the terms of the Creative Commons Attribution License (<http://creativecommons.org/licenses/by/4.0>), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

*Peer-review history:*

*The peer review history for this paper can be accessed here:*  
<https://www.sdiarticle5.com/review-history/93114>