

ORIGINAL RESEARCH**Assessment of spectrum of heart diseases in children****¹Dr Neeraj Agarwal, ²Dr Rishi Bansal**^{1,2}Associate Professor, Dept of Pediatrics, FH Medical College and Hospital, Etmadpur, Agra, U.P., India**Correspondence:**

Dr Rishi Bansal

Associate Professor, Dept of Pediatrics, FH Medical College and Hospital, Etmadpur, Agra, U.P., India

Received: 19 September, 2022

Accepted: 27 October, 2022

Abstract**Background:** Congenital heart defects (CHD) are the main heart diseases found in children. The present study was conducted to assess spectrum of heart diseases in children.**Materials & Methods:** 48 children of both genders underwent echocardiographic examination. Presenting complaints, Clinical signs, Different types of diagnosed heart diseases and Different types of cardiomyopathy (CMP) were recorded.**Results:** Out of 48 patients, males were 28 and females were 20. Common complaints were cough in 23, sweating in 22, anorexia in 16, chest pain in 11, cyanosis in 32, failure to thrive in 40, dyspnea in 34. Clinical signs were heart murmur in 32, dyspnea in 28, turgescient jugular veins in 30, muffled heart sounds in 23, ascites in 16 and dysrhythmia in 18. Type of cardiopathy was ASD in 20, VSD in 11, PDA in 5, AVCD in 7, TA in 3 and ToF in 2 cases. The difference was significant ($P < 0.05$).**Conclusion:** The most common congenital heart diseases observed were atrio-septal defects, ventricular septal defects, patent ductant arteriosus and tetralogy of fallot.**Key words:** Congenital heart defects, Children, Cardiopathy**Introduction**

Congenital heart defects (CHD) are the main heart diseases found in children. Heart defects have a prevalence of eight cases per 1,000 live births across the globe, representing approximately 1.35 million newborns each year with CHD.^{1,2} The impact appears to vary only slightly by region of the world, but the biggest contrast lies in the differences of priorities that are granted according to whether it is in the Western or in the southern countries.³ Rheumatic fever (RF), stimulated and promoted by precarious socio-economic conditions, has become an exceptional occurrence in high income countries. However, it remains a public health problem in low income countries.⁴

Congenital heart disease, in a definition proposed by Mitchell et al. (1), is “a gross structural abnormality of the heart or intrathoracic great vessels that is actually or potentially of functional significance.”⁵ It usually excludes congenital arrhythmias such as the long QT and the Wolf-Parkinson-White syndromes, even if the disorders are based on abnormalities present at birth.⁶ Lesions such as hypertrophic or dilated cardiomyopathy are usually not regarded as CHD. Even though the abnormal genes that cause these disorders are present at birth, the cardiomyopathy is rarely detected at this time but usually presents later in childhood or adolescence.⁷ The present study was conducted to assess spectrum of heart diseases in children.

Materials & Methods

The present study comprised of 48 children of both genders. Parental consent was obtained before starting the study.

Data such as name, age, gender etc. was recorded. All patients underwent echocardiographic examination. Presenting complaints, Clinical signs, Different types of diagnosed heart diseases and Different types of cardiomyopathy (CMP) were recorded. Results thus obtained were subjected to statistical analysis. P value less than 0.05 was considered significant.

Results

Table I Distribution of patients

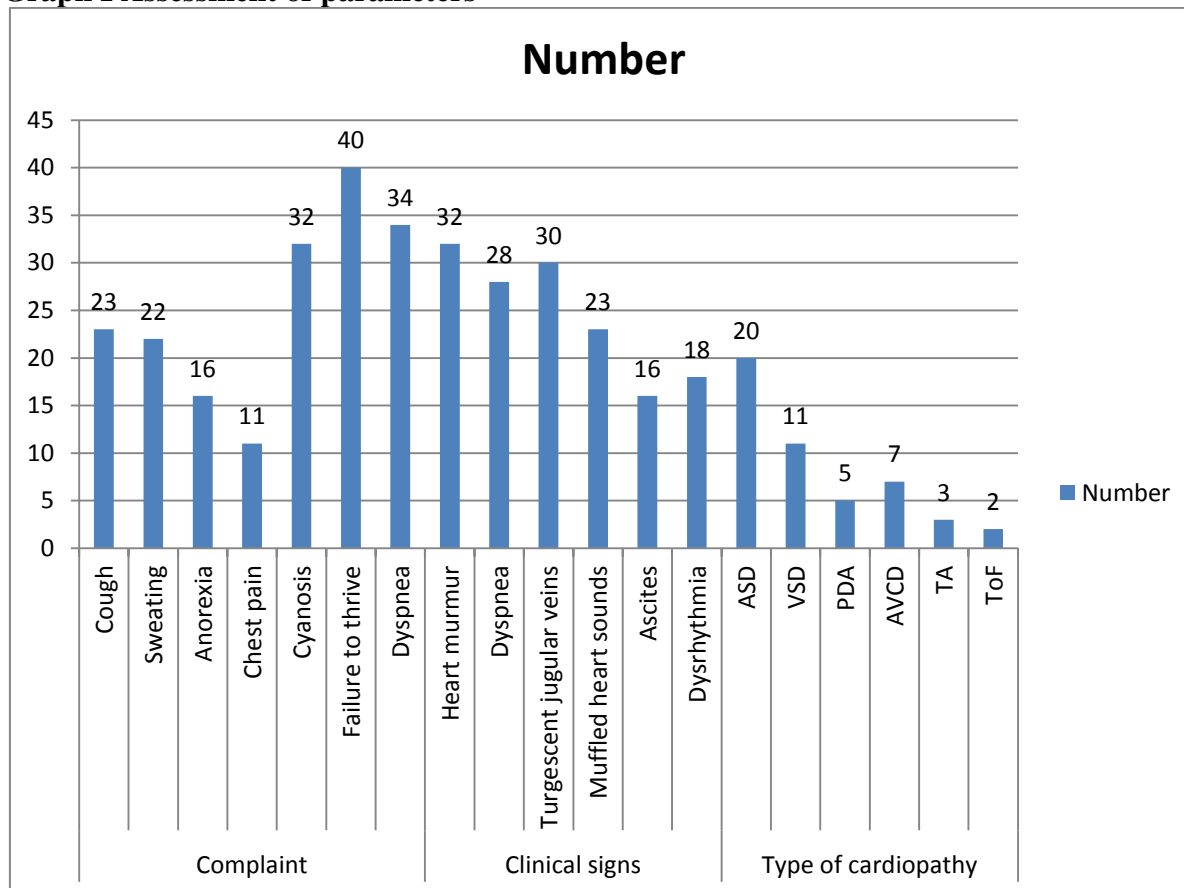
Total- 48		
Gender	Boy	Girl
Number	28	20

Table I shows that out of 48 patients, males were 28 and females were 20.

Table II Assessment of parameters

Parameters	Variables	Number	P value
Complaint	Cough	23	0.01
	Sweating	22	
	Anorexia	16	
	Chest pain	11	
	Cyanosis	32	
	Failure to thrive	40	
	Dyspnea	34	
Clinical signs	Heart murmur	32	0.02
	Dyspnea	28	
	Turgescient jugular veins	30	
	Muffled heart sounds	23	
	Ascites	16	
	Dysrhythmia	18	
Type of cardiopathy	ASD	20	0.04
	VSD	11	
	PDA	5	
	AVCD	7	
	TA	3	
	ToF	2	

Table II, graph I shows that common complaints were cough in 23, sweating in 22, anorexia in 16, chest pain in 11, cyanosis in 32, failure to thrive in 40, dyspnea in 34. Clinical signs were heart murmur in 32, dyspnea in 28, turgescient jugular veins in 30, muffled heart sounds in 23, ascites in 16 and dysrhythmia in 18. Type of cardiopathy was ASD in 20, VSD in 11, PDA in 5, AVCD in 7, TA in 3 and ToF in 2 cases. The difference was significant ($P < 0.05$).

Graph I Assessment of parameters

Discussion

The number of patients with some form of congenital heart disease (CHD) is growing rapidly as therapy becomes increasingly effective.^{8,9} Some of these patients have only mild disease with relatively little need for medical care, but others have complicated problems and require the services of an array of people with great expertise in the field.^{10,11} The present study was conducted to assess spectrum of heart diseases in children.

We found that out of 48 patients, males were 28 and females were 20. Chelo et al¹² in their study a total of 2,235 patients underwent echocardiographic examination during the study period including 1,666 subjects with heart disease. Congenital cardiopathies were found in 1,230 (73.8%) patients and acquired abnormalities in 429 (25.8%). Seven children (0.4%) had a combination of both types. Congenital heart defects (CHD) were dominated by ventricular septal defect (VSD). Acquired heart disease was mostly rheumatic valvulopathies. Dyspnea on exertion was the most frequent presenting complaint (87.6%). Discovery of a heart murmur was the principal clinical finding on physical examination (81.4%). The median age was 9 months for congenital heart disease and 132 months for acquired heart disease.

We found that common complaints were cough in 23, sweating in 22, anorexia in 16, chest pain in 11, cyanosis in 32, failure to thrive in 40, dyspnea in 34. Clinical signs were heart murmur in 32, dyspnea in 28, turgescent jugular veins in 30, muffled heart sounds in 23, ascites in 16 and dysrhythmia in 18. Type of cardiopathy was ASD in 20, VSD in 11, PDA in 5, AVCD in 7, TA in 3 and ToF in 2 cases. Hoffman et al¹³ determined the reasons for the variability of the incidence of congenital heart disease (CHD), estimate its true value and provide data about the incidence of specific major forms of CHD. The incidence of CHD in different studies varies from about 4/1,000 to 50/1,000 live births. The relative frequency of

different major forms of CHD also differs greatly from study to study. In addition, another 20/1,000 live births have bicuspid aortic valves, isolated anomalous lobar pulmonary veins or a silent patent ductus arteriosus. The incidences reported in 62 studies published after 1955 were examined. Attention was paid to the ways in which the studies were conducted, with special reference to the increased use of echocardiography in the neonatal nursery. The total incidence of CHD was related to the relative frequency of ventricular septal defects (VSDs), the most common type of CHD. The incidences of individual major forms of CHD were determined from 44 studies. The incidence of CHD depends primarily on the number of small VSDs included in the series, and this number in turn depends upon how early the diagnosis is made. If major forms of CHD are stratified into trivial, moderate and severe categories, the variation in incidence depends mainly on the number of trivial lesions included. The incidence of moderate and severe forms of CHD is about 6/1,000 live births (19/1,000 live births if the potentially serious bicuspid aortic valve is included), and of all forms increases to 75/1,000 live births if tiny muscular VSDs present at birth and other trivial lesions are included. Given the causes of variation, there is no evidence for differences in incidence in different countries or times.

The shortcoming of the study is small sample size.

Conclusion

Authors found that most common congenital heart diseases observed were atrio-septal defects, ventricular septal defects, patent ductant arteriosus and tetralogy of fallot.

References

1. Abu-Harb M, Hey E, Wren C. Death in infancy from unrecognized congenital heart disease. *Arch Dis Child* 1994;71:3–7.
2. Kuehl KS, Loffredo CA, Ferencz C. Failure to diagnose congenital heart disease in infancy. *Pediatrics* 1999;103:743–7.
3. Richards MR, Merritt KK, Samuels MH, Langmann AG. Congenital malformations of the cardiovascular system in a series of 6,053 infants. *Pediatrics* 1955;15:12–29.
4. Michae' lsson M. Congenital heart disease—some data on the relative incidence, natural history and operability. *Acta Paediatr Scand* 1965;154 Suppl 159:154–5.
5. Kerrebijn KF. Incidence in infants and mortality from congenital malformations of the circulatory system. *Acta Paediatr Scand* 1966; 55:316–20.
6. Carlgren LE. The incidence of congenital heart disease in Gothenburg. *Proc Assoc Eur Paediatr Cardiologists* 1969;5:2–8.
7. Feldt RH, Avasthey P, Yoshimasu F, Kurland LT, Titus JL. Incidence of congenital heart disease in children born to residents of Olmsted County, Minnesota 1950–1969. *Mayo Clin Proc* 1971;46: 794–9.
8. Czeizel A, Kamara's J, Balogh O, Szentpeteri J. Incidence of congenital heart defects in Budapest. *Acta Paediatr Hung* 1972;13: 191–202.
9. Sama'nek M, Slav'ik Z, Zborilova' B, Hrobonova' V, Vor'iskova' M, Skovra'nek J. Prospective study on incidence, treatment and outcome of heart diseases in 91,823 live born children. *Pediatr Cardiol* 1989;10:205–11.
10. Stoll C, Alembik Y, Roth MP, Dott B, De Geeter B. Risk factors in congenital heart disease. *Eur J Epidemiol* 1989;5:382–91.
11. Ferencz C, Czeizel A, Lys F. The problem of comparative analysis of birth prevalence of congenital cardiovascular malformations. *Acta Paediatr Hung* 1990;30:169–89.
12. Chelo D, Nguetack F, Menanga AP, Um SN, Gody JC, Tatah SA, Ndombo PO. Spectrum of heart diseases in children: an echocardiographic study of 1,666 subjects in a pediatric hospital, Yaounde, Cameroon. *Cardiovascular Diagnosis and Therapy*. 2016 Feb;6(1):10.

13. Hoffman JI, Kaplan S. The incidence of congenital heart disease. Journal of the American college of cardiology. 2002 Jun 19;39(12):1890-900.