

CARDIAC EVALUATION IN NEWBORNS: AN ECHOCARDIOGRAPHY BASED STUDY

Muhammad Sohail Arshad¹, Waqas Shakir¹, Hafiz Muhammad Anwar-ul-Haq¹, Mudasser Adnan¹,
Munir Ahmad¹

1 Children's Hospital & The Institute of Child Health, Multan

Address for Correspondence:

Munir Ahmad
Department of Neonatology, Children's Hospital & The Institute of Child Health, Multan, Pakistan
Emails: dr.munir.ahmad2@gmail.com

Contribution

MSA conceived the idea and designed the study. Data collection and manuscript writing was done by MSA, WS, HMAH, MA, and MA. All the authors contributed equally to the submitted manuscript.

All authors declare no conflict of interest.

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ABSTRACT

Objective: To evaluate the cardiac status of newborns in terms of structural and functional heart diseases.

Methodology: This descriptive study was conducted at Paediatric Cardiology Department of "The Children's Hospital and The Institute of Child Health", Multan, Pakistan, from January 2017 to December 2019. Neonates aged 0-28 days and referred for echocardiographic evaluation were enrolled. Diagnostic findings were confirmed with the assistance of echocardiography. We were focused on analyzing total number and types of heart diseases along with gender distribution during the study period.

Results: During the study period, a total of 2729 newborns were evaluated. We noted 1523 (55.8%) newborns as structurally and functionally normal, 866 (31.7%) had congenital heart disease, 69 (2.5%) duct dependent lesions, 248 (9.1%) persistent pulmonary hypertension and 23 (0.8%) left ventricular (LV) dysfunction but with structurally normal heart. Newborns with LV dysfunction might have hypoxic myocardial damage, metabolic derangements or acute myocarditis. Ventricular Septal Defect (VSD) followed by Patent Ductus Arteriosus (PDA) and Atrial Septal Defect (ASD) were the most frequent types of acyanotic CHD observed in 248 (28.2%), 171 (19.7%) and 100 (11.5%) neonates respectively while Tetralogy of Fallot (TOF) was the most common type of cyanotic CHD 74 (8.5%).

Conclusion: VSD followed by PDA, ASD and TOF were the most common types of CHD among neonates. Diagnosis of congenital heart defects in the early age is pointing towards improvement in healthcare facilities.

Keywords: Congenital heart disease, echocardiography, ventricular septal defect.

INTRODUCTION

“A gross structural abnormality of the heart or intra-thoracic great vessels that is actually or potentially of functional significance” was labeled as congenital heart disease (CHD) by Mitchell SC and Colleagues.¹ Congenital heart defects are considered to be a global health issue and the most frequent cause of congenital anomalies contributing around 28.0% of all congenital anomalies while incidence of congenital heart defects is estimated to be between 3 to 10/1000 live births, however, some researchers estimated this incidence to be as high as 17.5/1000 live births.^{2,3}

Among developing countries like Pakistan, burden of congenital heart defects is mounting every passing day due to rise in risk factors and causative factors. In some regional community based findings, it has been revealed that more than 30% of congenital heart defects cases require critical interventions during 1st year of life and majority come to cardiac healthcare units with complications.⁴ Data from developed countries have shown an increase in the survival rates from 20-80% in adult CHD cases because of early detection and appropriate treatment modalities.^{5,6}

Recent data from Pakistan have found ventricular septal defect (VSD) to be the most prevalent type of acyanotic heart lesion (27%) among children with CHD whereas Tetralogy of Fallot (TOF) was seen to be the commonest types of cyanotic heart defect noted in 10.9% cases.⁷ In the last 5 years, to the best of our knowledge, no study has been conducted specifically to find out the pattern or types of CHD in neonates while data from 2010 suggested VSD in 31.0% followed by atrial septal defect (ASD) in 22.9% and patent ductus arteriosus (PDA) in 15.0% of neonates with CHD.⁸

In the last few decades, significant progress has been made in improving the screening and treatment of congenital heart defects especially in the developed countries. Understanding the patterns of congenital heart defects is of great interest for the researchers and policymakers worldwide as knowing about the disease types and its burden is of huge importance. Acknowledging differences in different patterns and trends in congenital heart defects can lead to better insights to enhance the current practices in centers catering these cases. As no

recent data elaborates frequency or patterns of congenital heart defects in neonates in Pakistan, this study was aimed at to evaluate the cardiac status of newborns in terms of structural and functional heart diseases.

METHODOLOGY

This descriptive study was conducted at Paediatric Cardiology Department of “The Children’s Hospital and The Institute of Child Health”, Multan, Pakistan, from January 2017 to December 2019. Approval from “Institutional Ethical Committee” was taken for this study (No.203, dated: 15/01/2018).

All newborns from birth to 28 days of age, referred for echocardiographic evaluation were enrolled. Reason for referral included O₂ dependency, cyanosis, murmur, any other associated anomalies (e.g. VACTER, VACTRAL, cleft lip and palate, TEF, TORCH) and family history of CHD or maternal diabetes mellitus. Newborns who could not be sent for echocardiographic evaluation were excluded from this study.

Echocardiography among all cases was done by consultant Paediatric Cardiologist. Newborns with patent foramen ovale and small PDA were considered normal as these are age related findings. Likewise, babies with bicuspid aortic valve without any aortic stenosis or aortic regurgitation and babies with mild to moderate pulmonary hypertension were also considered as normal. In case of a combination or multiple cardiac lesions in a patient, the major lesion was taken as the main diagnosis. We were focused on analyzing total number and types of heart diseases along with gender distribution of cases. All the study data was recorded on a predesigned proforma. Data was handled and analyzed using SPSS version 26.0. Data was represented in terms of frequency and percentage.

RESULTS

During the study period from 2017 to 2019, a total of 2729 newborns were evaluated. Table number 1 is showing diagnostic findings of newborns evaluated during the study period. Out of these 2729 cases, 1523 (55.8%) were structurally and functionally normal, 866 (31.7%) had congenital heart disease,

69 (2.5%) duct dependent lesions, 248 (9.1%) persistent pulmonary hypertension (these patients did not have congenital heart disease) and 23 (0.8%) left ventricular (LV) dysfunction but with structurally normal heart. Newborns with LV dysfunction might have hypoxic myocardial damage or metabolic derangements or acute myocarditis.

Table 1: Diagnostic Findings of Newborns Evaluated During the Study Period (n=2729)

Diagnostic Findings	Number (%)
Normal	1523 (55.8%)
Structural Heart Disease	
Duct Dependent CHD	69 (2.5%)
Non-Duct Dependent CHD	866 (31.7%)
Functional Heart Disease	
Persistent Pulmonary Hypertension	248 (9.1%)
Dysfunctioning LV	23 (0.8%)

We observed CHDs among 866 newborns. VSD followed by PDA and ASD were the most frequent types of acyanotic CHD observed in 248 (28.2%), 171 (19.7%) and 100 (11.5%) neonates respectively while TOF was the most common type of cyanotic CHD found among 74 (8.5%) neonates. Table number 2 is showing distribution of various types of CHDs noted among studied neonates with regards to gender. Among 866 CHDs cases, 666 were having acyanotic CHD while 200 had cyanotic CHDs.

Table 2: Types of Congenital Heart Disease with Regards to Gender Distribution

Types of CHD	Number (%)	Male (%)	Female (%)
Tetralogy of Fallot	74 (8.5%)	38 (51.3%)	36 (48.7%)
TGA with VSD with PS	19 (2.2%)	11 (57.9%)	8 (42.1%)
TGA with VSD with PH	12 (1.4%)	7 (58.3%)	5 (41.7%)
VSD	248 (28.2%)	135 (54.4%)	113 (45.6%)
ASD	100 (11.5%)	52 (52.0%)	48 (48.0%)
Complete AVSD	34 (3.9%)	17 (50.0%)	17 (50.0%)
PDA	171 (19.7%)	91 (53.2%)	80 (46.8%)

Pulmonary atresia	19 (2.2%)	10 (52.6%)	9 (47.4%)
Tricuspid atresia	5 (0.6%)	3 (60.0%)	2 (40.0%)
Univentricular Heart + Complex CHD	31 (3.6%)	17 (54.8%)	14 (45.2%)
Aortic Stenosis	31 (3.6%)	16 (51.6%)	15 (48.4%)
Pulmonary Stenosis	35 (4.0%)	19 (54.3%)	16 (45.7%)
Coarctation of Aorta	47 (5.4%)	28 (59.6%)	19 (40.4%)
Cardiac TAPVC	1 (0.1%)	0 (0%)	1 (100%)
Supra cardiac TAPVC	1 (0.1%)	1 (100%)	0 (0%)
Ebstein Anomaly	14 (1.6%)	8 (57.1%)	6 (42.9%)
Others	24 (2.8%)	14 (58.3%)	10 (41.7%)
Total	866 (100%)	467 (53.9%)	399 (46.1%)

TGA with intact septum were the most common duct dependent lesion noted among 24 (34.8%) neonates followed by critical coarctation of aorta 17 (24.6%). Table number 3 shows distribution of various types of duct dependent lesions found among neonates.

Table 3: Distribution of Duct Dependent Lesions

Duct Dependent Lesions	Total	Male	Female
TGA with intact septum	24 (34.8%)	13 (54.2%)	11 (45.8%)
Critical Coarctation of Aorta	17 (24.6%)	10 (58.8%)	7 (41.2%)
Critical Aortic stenosis	6 (8.7%)	3 (50.0%)	3 (50.0%)
Pulmonary atresia with intact septum	15 (21.7%)	7 (46.7%)	8 (53.3%)
Hypoplastic Left Heart	7 (10.1%)	3 (42.9%)	4 (57.1%)
Total	69 (100%)	36 (52.2%)	33 (47.8%)

DISCUSSION

The exact prevalence of CHD at the time of birth has not yet been estimated globally which could further elaborate the influence of various environmental as well as genetic risk factors on the prevalence of

CHDs among different geographies of the world.⁹CHDs can involve the chamber or valves inside the heart or the blood vessels which carry blood to and from the heart. The defects can be in isolation or in combination with different defects. Physiologically they can be divided broadly into cyanotic and acyanotic heart defects.¹⁰Acyanotic heart defects can remain asymptomatic throughout life but some of them warrant early intervention. CHDs vary from simple defects without any symptoms to more complex lesions presenting as severe, life-threatening conditions. In contrast to developed countries, in parts of the developing world, access to timely treatment remains largely unavailable for CHDs. Better diagnostic facilities, early intervention and improved surgical outcomes have led to a change in spectrum of patients with congenital heart defects.¹¹ This improved survival as also led to development of new specialties like Grown up with congenital heart defects (GUCHD).

We notice a male predominance (53.9%) among neonates with CHDs. Previous local data has revealed 55.3% of the cases with heart defects to be male children.¹²Others have also found similar findings revealing male predominance among CHD cases⁷ while data from India also recorded 66% of CHD cases to be male.¹³We noted that VSD followed by PDA and ASD were the most frequent types of acyanotic CHD observed in 28.2%, 19.7% and 11.5% neonates respectively while TOF was the most frequent type of cyanotic CHD found among 8.5% neonates. Hussain S et al in a local study evaluating neonates with CHD noted VSD to be the commonest type of CHD (31.0%) followed by ASD (22.9%), PDA (14.9%) and TOF (6.9%).¹⁴ Farooqui R et al also noted VSD and PDA to be the most frequent forms of CHDs among neonates which is correlating well with the current data. VSD has been found the most frequent CHD type in Pakistan.⁸ Meshram RM and Gajimwar VS from India in a recent research noted VSD (30.1%) to form majority of CHD cases while ASD (20.7%), TOF (16.1%) and PDA (10.2%) were the other most frequent types of CHDs.¹⁵ Most of the other regional studies also report similar patterns of CHDs.¹⁶⁻¹⁸Differences in patterns and frequencies of CHD types may be credited to differences in detection methods, standards of healthcare care and variations in genetic and environmental factors.^{19,21} We were unable to record any complications of CHD among enrolled study participants but growth failure and congestive cardiac failure are some of the most common ones. Inpatient case fatality rate among neonates with CHD is around 19% while majority of the deaths have been attributed to complex CDH,

refractory congestive cardiac failure and sepsis.^{15,22} All murmurs should be carefully screened unless thought to be physiological. CHD requires regular monitoring to allow optimum growth and development.

Being the 1st study from a Pediatric Cardiology Unit from South Punjab is one of the major significance of this study but there is a need to form National Registry for these patients. We need more and more cardiac surgery facilities capable of performing surgeries in infancy among CHD cases requiring correction/intervention.

Our study had few limitations as well. We were unable to record typical patterns of presentation and comorbidities in our cases. We could not gather any follow up data about functional and growth limitations of these patients.

CONCLUSION

VSD followed by PDA, ASD and TOF were the most common types of CHD among neonates. Diagnosis of congenital heart defects in the early age is pointing towards improvement in healthcare facilities. Population based studies on a large scale are needed to estimate the burden and pattern of congenital heart defects in Pakistan.

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