

Assessment of cardiac dysfunction by M mode echocardiography in patients of CKD: A Cross- sectional study¹Dr. Preety Motiyani, Senior Resident, Department of General Medicine, Gandhi Medical College, Bhopal (M.P.)²Dr. RR Barde, Associate Prof, Department of General Medicine, Gandhi Medical College, Bhopal (M.P.)³Dr. Mamta Meena, Assistant Prof., Department of Microbiology, Gandhi Medical College, Bhopal (M.P.)⁴Dr. RS Meena, Associate Prof., Department of Cardiology, Gandhi Medical College, Bhopal (M.P.)⁵Dr. Jitendra Kumar Rai, Senior Resident, Department of General Medicine, Gandhi Medical College, Bhopal (M.P.)**Corresponding Author:** Dr. Jitendra Rai, Senior Resident, Department of General Medicine, Gandhi Medical College, Bhopal (M.P.)**How to citation this article:** Dr. Preety Motiyani, Dr. RR Barde, Dr. Mamta Meena, Dr. RS Meena, Dr. Jitendra Kumar Rai, “Assessment of cardiac dysfunction by M mode echocardiography in patients of CKD: A Cross- sectional study”, IJMACR- March – April - 2021, Vol – 4, Issue -2, P. No. 24 – 31.**Copyright:** © 2021, Dr. Preety Motiyani, et al. This is an open access journal and article distributed under the terms of the creative commons attribution noncommercial License 4.0. Which allows others to remix, tweak, and build upon the work non-commercially, as long as appropriate credit is given and the new creations are licensed under the identical terms.**Type of Publication:** Original Research Article**Conflicts of Interest:** Nil**Abstract****Background:** CVD mortality rates are much greater in dialysis patients despite stratification for sex, race, or age group. Younger dialysis patients have an approximately 500-fold increased CVD mortality rate compared with their counterparts in the general population, and rates remain approximately five times higher, even among the oldest patients.**Objectives:** To assess the prevalence of cardiac dysfunction in patients of CKD.**Materials and Methods:** 150 patients were sampled for study at Department of Medicine Gandhi Medical College and associated Hamidia Hospital Bhopal from September 2016 to June 2017. After detailed clinical evaluation in patients with feature suggestive of CKD, all the patients had undergone urine (PH, Specific gravity, Protein, Sugar, Microscopy), blood (Hb%, FBS/PPBS, Urea, Creatinine,

Electrolyte, Calcium, Phosphorous) and ultrasound abdomen. M mode electrocardiography (12 lead ECG) was done on patient with chronic kidney disease.

Results: Majority of the patients belong to age group of 60-70 years. Majority of the patients were male [90 (60.7%)]. ECHO findings revealed that diastolic dysfunction were present in 90.67% patients, LVH in 86.67, RWMA in 57.33 and pericardial effusion in 46.67% patients. Most common valve abnormality was MR/TR in 36.2% patients, among most common chamber abnormality DCMP (36.6%) was most common. Hypertensive patients had significantly more LVH (74.66%) as compared to non hypertensive (12%) (P=0.012). Patients with hypertension had higher percentage of diastolic dysfunction (88.67%), RWMA (55.33%) and pericardial Effusion (46%). Patients with anemia had higher percentage of diastolic dysfunction

(85.33%), LVH (82%), RWMA (54%) and pericardial Effusion (44.67%). Non diabetes patients had higher percentage of diastolic dysfunction (29.33%) and LVH (60%) as compared to diabetes whereas RWMA (40.67%) and Pericardial Effusion (32.67%) were higher among diabetes.

Conclusion: Patients with chronic kidney disease have higher prevalence of diastolic dysfunction and diastolic dysfunction appears to occur earlier than systolic dysfunction and left ventricular dysfunction was the commonest cardiovascular abnormality.

Keywords: Cardiovascular dysfunction, M mode echocardiography, CKD

Introduction

Patients with chronic kidney disease (CKD) are at significantly increased risk for both morbidity and mortality from cardiovascular disease (CVD). Cardiac disease is the single most important cause of death among patients receiving long-term dialysis, accounting for 44% of overall mortality^[1,2]. It is important to emphasize that the prevalence of CVD is increased among all patients with CKD, not only those with end-stage renal disease (ESRD). That is, the prevalence of LVH increases as glomerular filtration declines, and as many as 30% of patients reaching ESRD already have clinical evidence of ischemic heart disease or heart failure. Furthermore, it is important to note that patients with a reduced glomerular filtration rate (GFR) are more likely to die of CVD than they are to develop ESRD^[3]. In cardiovascular system, left ventricular dysfunction is most common finding but their proportion is less clear^[4]. Heart and kidney are inextricably linked in terms of hemodynamic and regulatory functions. Communication between these two organs occurs at multiple levels including the sympathetic nervous system, the renin-angiotensinaldosterone system (RAAS), antidiuretic hormone, endothelin, and the

natriuretic peptides. Cardiac disease is the major cause of death in dialysis population accounting for 40% of deaths in international registries^[5]. In 1997, National Kidney Foundation (NKF) convened a Task Force, which considered coronary artery disease (CAD) and left ventricular hypertrophy (LVH) as the two target conditions in their recommendations for decreasing the cardiovascular mortality in end-stage renal disease (ESRD) patients^[6]. Cardiac disease is frequently noted in individuals around the time of commencement of dialysis^[7]. There is scanty information on the prevalence and natural history of LVH in patients with milder degrees of chronic renal failure from India. The present study was aimed to assessing the prevalence of left ventricular hypertrophy or cardiac dysfunction by M mode echocardiography in patients of chronic kidney disease.

Material and methods

Study was carried out among 150 patients at tertiary care teaching hospital affiliated with Gandhi Medical College Bhopal for the duration September 2016 to August 2017. Patients with azotemia for more than 3 months, symptoms or signs of uremia, reduced Kidney size bilaterally, broad casts in urinary sediment and symptoms or signs of renal osteodystrophy were enrolled. After detailed clinical evaluation in patients with feature suggestive of CKD, all the patients had undergone urine (PH, Specific gravity, Protein, Sugar, Microscopy), blood (Hb%, FBS/PPBS, Urea, Creatinine, Electrolyte, Calcium, Phosphorous) and ultrasound abdomen. M mode electrocardiography (12 lead ECG) was done on patient with chronic kidney disease cases with CKD without considering the etiology, patient with chronic kidney disease on routine dialysis and age more than 18 years and less than 55 years were included whereas patients with documented Ischaemic heart disease, congenital heart disease, valvular heart disease and age less than 18 years and age more than 55

years were excluded from the present study. This study is approved by the Institutional ethics committee. All the data analysis was done by using a statistical software SPSS ver. 20. Frequency distribution and cross tabulation was used to prepare the tables. Quantitative data was expressed as Mean ± SD whereas categorical data was expressed as percentage. Chi square test was used to compare the categorical data. Level of significance was assessed at 5%.

Results

In present study mean Age, serum calcium, phosphate, potassium, sodium, urea, creatinine and hemoglobin was 56.60±16.71, 9.23±0.65, 6.09±1.17, 5.39±0.75, 139.89±2.37, 203.78±93.51, 9.93±5.13 and 6.67±2.14 respectively. A Total of 150 cases of CKD were screened out of them maximum were of age group in between 60-70 years. Majority of the patients were male [90 (60.7%)]. A total 101 (68%) patients were having diabetes. ECG findings revealed that majority of the patients had Hyper acute T wave (34.7%) followed by LV strain (26%) and

Low Voltage Complexes (16.7%). In present study ECHO findings revealed that diastolic dysfunction were present in 90.67% patients, LVH in 86.67, RWMA in 57.33 and pericardial effusion in 46.67% patients. Most common valve abnormality was MR/TR in 36.2% patients, among most common chamber abnormality DCMP (36.6%) was most common. RVSP (right ventricular systolic pressure) is the estimate of pulmonary artery pressure to assess for pulmonary artery hypertension. Majority (30%) of the patients had RVSP in range of 40-50mmhg and 25.3% patients had RVSP in range of 30-40 mmHg and 3.33% patients had RVSP in range of 20-25mmHg whereas 21.3% patients had RVSP in range of 25-30mmHg (46.6% patients had RVSP in between 20-30mmhg) which suggests that 46.6% patients had mild pulmonary artery hypertension whereas 30% patients having moderate pulmonary artery hypertension and 3.3% patients having severe pulmonary artery hypertension. That means out of 150 CKD patients 80% having pulmonary artery hypertension.

Table 1: Clinical characteristics of CKD patients in terms of age, serum calcium, phosphate, blood urea, serum creatinine, serum sodium/ potassium and hemoglobin.

Variable	Minimum	Maximum	Mean	Std. Deviation
AGE	23	87	56.60	16.71
Serum calcium	7.00	10.40	9.23	0.65
Serum phosphate	1.60	8.00	6.09	1.17
Serum potassium	3.90	6.70	5.39	0.75
Serum sodium	136	145	139.89	2.37
Urea	54	544	203.78	93.51
Creatinine	3.00	43.00	9.93	5.13
Hb (%)	2.3	11	6.67	2.14

Table 2: ECG findings

ECG findings	Frequency	Percent
Hyper acute T wave	52	34.7
Low Voltage Complexes	25	16.7
LV Strain	39	26
Normal	34	22.7
Total	150	100.0

Table 3: Prevalence of cardiac dysfunction with ECHO (Echocardiography)

Variable	No of patients (n=150)	Percentage
Diastolic dysfunction	Present	136
	Absent	14
LVH	Present	130
	Absent	20
RWMA	Present	86
	Absent	64
Valve abnormality	Mild to Moderate MR	37
	MR/TR	54
	Mild TR	30
	Normal	29
Chambers	DCMP	54
	LA/LV enlarged	28
	RA/RV enlarged	53
	Normal	15
Pericardial Effusion	Present	70
	Absent	80

Table 4: Comparing Hypertension Status With ECHO (Echocardiography)

Variable	Status	HTN (>140/80mmhg) n (%)	Non HTN (<140/80mmhg); n (%)	P value
Diastolic dysfunction	Present	133 (88.67)	3 (2)	0.242
	Absent	13 (8.67)	1 (0.67)	
LVH	Present	112 (74.66)	18 (12)	0.012
	Absent	19 (12.67)	1 (0.67)	
RWMA	Present	83 (55.33)	3 (2)	0.478
	Absent	63 (42)	1 (0.67)	

Valve abnormality	Mild to Moderate MR	35 (23.33)	3 (2)	0.413
	MR/TR	52 (34.67)	3 (2)	
	Mild TR	30 (20)	0 (0)	
	Normal	29 (19.33)	0 (0)	
Chambers	DCMP	84 (56)	3 (2)	0.713
	LA/LV enlarged	14 (9.33)	2 (3)	
	RA/RV enlarged	48 (32)	1 (0.67)	
Pericardial Effusion	Present	69 (46)	1 (0.67)	0.372
	Absent	77 (51.33)	3 (2)	

Table 5: Prevalence of LVH and diastolic dysfunction in CKD patients

	No.	Percentage
LVH with systolic dysfunction	14	9.3
LVH with diastolic dysfunction	136	90.7
Total	150	100

Discussion

Premature cardiovascular disease is a significant cause of morbidity and mortality among patients with CKD. Four main structural abnormalities of the heart have been described in patients with CKD: LV hypertrophy, expansion of the nonvascular cardiac interstitium leading to inter-myocardiocytic fibrosis, changes in vascular architecture, and myocardial calcification. All these abnormalities promote systolic as well as diastolic LV dysfunction which predisposes to symptomatic heart failure, which in turn is a risk factor for premature death. Various diagnostic modalities, both invasive and noninvasive such as electrocardiography, echocardiography and radionuclide scans are utilized for diagnosing left ventricular hypertrophy and dysfunction. Cardiac assessment by echocardiography is non-invasive,

inexpensive to perform and generates detailed information about gross cardiac anatomy, objective quantification of LVM and the geometry of left ventricle hypertrophy (LVH), along with measures of function during systole and diastole. Prospective studies have shown the presence of LVH in most (70–80%) patients with end-stage renal failure, and this confers a poor prognosis^[8]. In present study mean Age, serum calcium, phosphate, potassium, sodium, urea, creatinine and hemoglobin was 56.60±16.71, 9.23±0.65, 6.09± 1.17, 5.39 ±0.75, 139.89±2.37, 203.78±93.51, 9.93±5.13 and 6.67±2.14 respectively. A similar study by Laddha *et al.* found that maximum number of patients belonged to age group of 51-60 years. Mean age of ESRD patients was 53.3 ± 12.8^[9]. Study done by Franczyk-Skóra *et al.* reported that the mean age of patients with CKD II–IV was 63.97 ±12.5

years and 67.78 ± 12.0 in dialysis group^[10]. Reports of Laddha *et al.* reported that mean haemoglobin percentage was 7.78 ± 1.84 gm%. Mean blood urea level was 151.7 ± 51.37 mg%. Mean serum creatinine level was 10.35 ± 5.56 gm%^[9]. In present study majority of the patients were male 90 (60.7%) followed by 59 (39.3%) females. Reports of Laddha *et al. et al.* showed that out of 70 patients, there were 53 males (75.7%) and 17 females (24.3)^[9]. Franczyk-Skóra *et al.* showed that there were more males in dialysis group than in other CKD group (68.57% vs. 30.12%)^[10]. In present study ECG findings revealed that majority of the patients had Hyper acute T wave (34.7%) followed by LV strain (26%) and Low Voltage Complexes (16.7%). In a similar study by Laddha *et al.* showed that ECG changes in decreasing order of frequency were sinus tachycardia in 48.6%, LVH in 45.7%, ST –T changes in 30%, ventricular ectopics and Tall ‘T’ wave in 7.1%, QT prolongation and low voltage pattern in 5.7%, ventricular tachycardia in 2.9% and complete heart block in 1.4% was noted^[9].

Echocardiography provides an excellent non-invasive method to study the details of anatomy of cardiac chambers, wall dimensions, valve movements and is also used to assess the cardiac performance. In present study ECHO findings revealed that diastolic dysfunction were present in 90.67% patients, LVH in 86.67, RWMA in 57.33 and pericardial effusion in 46.67% patients. Most common valve abnormality was MR/TR in 36.2% patients, among most common chamber abnormality DCMP (57.7%) was most common. In the study of Laddha *et al.*^[9], LVH was present in 74.3%, systolic dysfunction was present in 24.3% of patients as suggested by reduced LVEF measurement and diastolic dysfunction was observed in 61.4% by abnormal E/A ratio of ESRD patients. Mild pericardial effusion (less than 10 mm thickness) was present in 14.3% patients. Mitral and aortic

valve calcification and mitral regurgitation was noted in 7.1% patients^[9]. Robert N. Foley *et al.* (1995) had found abnormalities of left ventricular structure and functions were very frequent on baseline echocardiography: 73.9% had left ventricular hypertrophy, 35.5% had left ventricular dilatation and 14.8% had systolic dysfunction in ESRD patients^[11]. NP Singh *et al.* (2000) had found LVH in 76.92%, diastolic dysfunction in 72% but did not find systolic dysfunction in CKD patients^[12]. Zoccali *et al.* (2000) had found 77% LVH, 22% systolic dysfunction by LVEF measurement in haemodialysis patients^[13]. S. Agarwal *et al.* (2003) had observed diastolic dysfunction in 60% and systolic dysfunction in 15% of patients^[14]. The above findings were consistent with our study. Franczyk-Skóra *et al.* assessed indices of LV diastolic dysfunction in 118 CKD patients and reported that the analysis of echocardiographic parameters showed that in CKD patients the stage of renal failure was associated with the significant increase in LV mass, IVSd, IVSs, systolic LV and diastolic LV^[10]. Parfrey *et al.*^[15] study, it has been observed that shortly after the dialysis session, a reduction in diastolic diameter of the LV and an increase in the thickness of the LV wall occur which is associated with volume depletion by ultrafiltration. Also, in this study, the RV diameter was found to be much greater in CKD patients stage V/ dialysis (29.9 ± 2.9) than in stages II–IV. Left ventricular muscle mass was over 1.5-times higher in dialysis patients than in CKD stage II subjects in the study done by Franczyk-Skóra *et al.* [6] According to Zoccali *et al.*^[13] the increase in mass of 1 g/m 2.7/ month was associated with a 62% increase in the incident risk of fatal and non-fatal cardiovascular events in dialysis patients. They also suggested that changes in LV mass index represent a stronger predictor of mortality and cardiovascular complications than LV mass itself. Study done by Agarwal *et al.* is in agreement to the present study

findings as in present study maximum patients had diastolic dysfunction (90.67%)^[14]. Left ventricular diastolic dysfunction is an important cause of cardiac morbidity in ESRD patients.

In present study there was statistically significant association between the findings of 2D – Echo in patients having hypertension as compared to normotensive group for LVH. Juan M. *et al.* (1998) had found statistically significant difference in E/A ratio, fractional shortening, and LVEF among hypertensive and normotensive patients^[16]. Patrick S *et al.* (1999) had found that rise in mean arterial blood pressure was associated with increase in LVH in ESRD patients^[17]. SA Kale *et al.* (2001) had found that hypertension was identified as important risk factor for all three LV disorders LVH, diastolic dysfunction and systolic dysfunction^[18]. In agreement to present study Laddha *et al.* assessed the prevalence of systolic and diastolic dysfunction in patients of end stage renal disease (ESRD) on haemodialysis and reported that LVH was more common in hypertensive (87.5%) patients as compared to non hypertensive (45.5%) patients ($p=0.01$), whereas RWMA (16.7% vs.4.5%) and pericardial effusion (14.6% vs.13.6%) was similar among both the groups in hypertensive and non-hypertensive patients respectively^[9]. Franczyk-Skóra *et al.* showed that subjects with hypertension were more likely to have atrial fibrillation, increased diameter of LA, increased atrial volume before dialysis and decreased after it as well as reduced early diastolic velocity (E') and ejection fraction^[6].

Conclusion

Based on the study findings we conclude that cardiovascular abnormalities in chronic kidney disease were observed in maximum no of patients and left ventricular dysfunction was the commonest cardiovascular abnormality. LVH was the most common

echocardiographic abnormality in chronic kidney disease cases. Diastolic function was deranged in more number of patients. Major contributing factors for left ventricular hypertrophy and diastolic dysfunction were diabetes, hypertension and anaemia. Echocardiography was more sensitive for detecting LVH and minimal pericardial effusion prior to clinical detection. Early identification of factors involved is necessary to prevent this devastating process.

Limitations

Cross sectional nature of the study was the main drawback of the study so, present study findings cannot be applied to larger populations, and a large randomized clinical trial is needed to strengthen the present study results. Another limitation was that we were not included hyperthyroidism, hyper homocysteine and other invasive methods in present study.

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