ABNORMAL LEFT VENTRICULAR MASS INDEX-A SURROGATE END POINT IN VITAMIN D DEFICIENT CHRONIC KIDNEY DISEASE

Pinaki Mukhopadhyay¹, Kartik Patar², Nandini Chattejree², Kajal Ganguly³

Dept. of Nephrology¹, Dept. of Medicine², Dept. of Cardiology³ NRS Medical College & Hospital, Kolkata, West Bengal, India.

Cardiovascular morbidity and mortality is prevalent in established chronic kidney disease(CKD) vitamin D deficiency.

AIMS

To evaluate the effect of vitamin D deficiency on cardiac structure and functions in predialysis diabetic and non-diabetic CKD patients.

METHOD

All diabetic and nondiabetic CKD stage 3 onwards predialysis patients not on any form of vit D therapy were includedprospectively. Demographical, clinical and laboratory data were analysed.Vit D was measured in all cases and echocardiogram was done and all the parameters was examined. Left Ventricular Mass index (LVMI) was calculated by Devereux's(Penn) formula.More than 134 g/m2in men and 110 g/m2 in female was considered abnormal.

Vitamin D Deficiency in CKD Renin Angiotensin System/Insulin Resistance Inflammation Oxidative stress Immune dysfunction Thrombosis Hyperparathyroidism VSM proliferation Hypocalcemia Hypertension Vascular dysfunction & Atherogenesis CKD and CVD Progression

Conceptual model of major pathways through which vitamin D deficiency in patients with chronic kidney disease (CKD) may lead to CKD progression and complications such as premature cardiovascular disease (CVD). VSM – vascular smooth muscle. Adapted from ref 22.

STATISTICAL ANALYSIS

Appropriate parametric and non-parametric statistical tests were performed to check any significant difference, correlation and association of parameters with the disease state. Chi-Squir test used to analysed nonparametric data and calculation of 'p' value, data are also plotted in percentage form wherever needed.

Multiple regression analysis was carried out using back ward selection method, regression criteria being p=>0.05 vitamin D ,eGFR, ejection fraction and haemoglobin as predictor with LVMI as regressor.

Showing correlation between left ventricular mass index and vitamin deficiency

Vitamin D Level	Standard Error	95% CI		
¢30ng/ml	8.012	99.75-104.94		
10-30 ng/ml	6.191	26.34-152.16		
<10 ng/ml	6.081	42.29-167.54		

	Pearson Correlation sig. (2-tailed), N = 150						
	Age	eGFR	LVMI	Vit D	EF	Hb	
LVMI	.409	.021*		.011*	.231	.447	
VitD	.101	.011*	.011*		.532	.645	
EF	.246	.561	.231	.532		.899	
Hb	.195	.010*	.447	.645	.899		

^{*}Correlation is significant at the 0.05 level (2-tailed).

RESULTS

About 150 pts were enrolled in one year of them M:F were 1.6:1. Mostly belonged to the age group of 41-60yrs(58%). About 96(64%) were diabetic and rest of 54(36%) were non diabetic. CKD III,IV,V comprises of 26%,14%,24% and 4%,12% and 20% respectively. Diabetic CKD patients are more severely deficient of vitamin D than non-diabetic CKD (38% vs 8%). Higher stages of CKD showed severe grade of Vit D deficiency. Among Stage V CKD 14% were grade II and 28% were grade III vitamin D deficient (p=0.07). Among all CKD patients with severe vitamin D deficiency (stage III) 15 (10 %) had moderatelyabnormal(Group II) LVMI and 51(34%) had severely abnormal (Group III) LVMI and among moderate vitamin D deficient (stage II) patients 36(24%) had group II LVMI and 21(14%) had group III LVMI. This correlation is extremely significant (p=0.0163).There is significant correlation between lowVitamin D level and LVMI related inversely with standard error 6.08 with 95% CI (142.2-167.5).

CONCLUSION

Chronic kidney disease with vitamin D deficiency associated with abnormal LVMI which results in widespread risk for CVD in CKD.

DR. PINAKI MUKHOPADHYAY

MD. DCH. DM. (PGI, CHANDIGARH) Associate Professor & H.O.D. Dept. of Nephrology, NRS Medical College, Kolkata, India C/51, East Rajapur, Santoshpur, Jadavpur, Kolkata-700 075, India Phone: 91-033-9231978078

E-mail : drpinaki71@yahoo.com





