



DYSLIPIDIMIA; TO DETERMINE THE VARIOUS LIPID PROFILE PATTERN IN PATIENTS OF CHRONIC RENAL FAILURE

Dr. Ashok Kumar Lohano¹, Dr. Irfan Ahmed Bhatti², Dr. Azhar Iqbal³, Dr. Salman Shah Jilani⁴

1. Assistant Professor Medicine
Peoples University Medical Hospital
Nawabshah
2. Postgraduate Student
Peoples University Medical Hospital
Nawabshah
3. Registrar Nephrology Department
Shaikh Zayed Hospital
Raheem Yar Khan
4. Senior Medical Officer
Fuji Foundation Pakistan.

Correspondence Address:
Dr Ashok Kumar Lohano
Assistant Professor Medicine
Flat # 1 Pak Plaza,
Trunk Bazar Nawabshah
drashokfcp@yahoo.com

Article received on:
21/02/2015

Accepted for publication:
24/04/2015

Received after proof reading:
09/07/2015

ABSTRACT... Objective: To determine the various lipid profile pattern of dyslipidemia in patients of chronic renal failure. **Study design:** Cross-sectional study. **Place and duration of study:** This study was conducted at Nephrology department of Isra University hospital Hyderabad and Medicine Department Civil hospital Nawabshah, from October 2013 to September 2014. **Methodology:** This study consisted of 237 patients of chronic renal failure asses on detailed history regarding chronic renal failure, patients were subject to relevant investigations i.e. fasting lipid profile and venous blood was drawn after 14 hours fasting state in the early morning and sent to laboratory for analysis of fasting lipid profile. Inclusion Criteria were all patients of chronic renal failure with 1 years history, age > 35 to 60 years and either any sex. Diagnosed on the basis of serum creatinine as define in operational definition. Exclusion Criteria were patients on lipid lowering agents, acute complication of diabetes mellitus such as diabetic keto-acidosis, patients suffering from hypothyroidism, type 1 DM and HTN, lactic-acidosis, hypoglycemia. Results were prepared with help of tables and graphs. Data was analyzed through SPSS software. **Results:** Out of 237 patients, 113(47.68%) patients were 51 to 60 years of age, 94(39.66%) were 41 to 50 years of age and 30(12.66%) were 35 to 40 years as presented in Chart No 1. The average age of the patients was 50.11 ± 6.94 years. There were 54.43% (129/237) male and 45.57% (108/237) female. Various lipid profile pattern of dyslipidemia in patients of chronic renal failure are presented in table-I. Results of fasting lipid profile are Triglycerides high in 86(36.3%), HDL low in 76(32.1%) cases, LDL in high 83(35%) cases, Cholesterol high in 105(44.3%) cases and VLDL-cholesterol high in 68(28.7%). Stratification analysis showed that lipid profile pattern of dyslipidemia was insignificant between male and female as presented in table 2. While with respect to age groups, rate of Cholesterol, triglycerides, LDL-cholesterol was high in above 40 years of age groups as compare to 30 to 40 years of age groups. Rate of low HDL was also high in above 40 years of age groups as compare to 30 to 40 years as presented in table 3. **Conclusions:** In conclusion, our study demonstrated that dyslipidemia often our patients chronic renal failure. This underscores the need for the first test patients with lipid abnormalities early treatment can prevent cardiovascular events delay chronic kidney disease.

Key words: Chronic renal failure, various lipid profile pattern, Dyslipidaemia.

Article Citation: Lohano AK, Bhatti IA, Iqbal A, Jilani SS. Dyslipidimia; to determine the various lipid profile pattern in patients of chronic renal failure. Professional Med J 2015;22(7):865-870.

INTRODUCTION

Chronic renal failure (CRF) is characterized by an irreversible loss in renal function.¹ Chronic Renal Failure (CRF) is associated with premature atherosclerosis and increased incidence of cardiovascular morbidity and mortality.^{2,3} Numerous reasons associated to atherogenesis and cardiovascular disease in patients with CRF. During the CRF-induced lipid risk of these problems, oxidative stress, inflammation, lack of exercise, anemia, high blood pressure, vascular calcification, endothelial, and the availability

of nitric oxide in distress.^{4,5} Over the past 30 years, numerous studies have been brought to the characteristics and procedures of the CRF-induced dyslipidemia. Most previous studies have focused on the effect of CRF on the concentration, the formation of, and permission to various plasma lipoproteins and their remnants.²

Chronic kidney disease (CKD) is marked by a growing worldwide and is associated with a poor prognosis.⁶ 1999-2004 National Health and Nutrition Examination Survey (NHANES), USA

15.3%, according to the prevalence of CKD.⁷ CKD is an independent causal factor for incident cardiovascular disease (CVD), Fair CKD and mortality in patients with end-stage renal disease as the leading cause of CVD point out that there is still debate about whether the accumulation of evidence over the past decade (ESRD).^{8,9} Khalid Amin’s study of 50 patients, regardless of the cause and sex, fasting lipid profile reported under CRF scheduled. There are triglycerides, fasting lipid profile results in 23 (46%), low HDL (16%) 8 cases (4%), 2 cases, high cholesterol (32%) of the initial 16 cases and 8 (16% of cases in LDL and total lipid) cases 5 () and cases (4) of 8% to 10%.¹

Aim of this study is to determine different pattern of dyslipidemia in chronic renal failure patients. There by strategy could be developed for its management and prevention of complications.

MATERIAL & METHODS

This study was conducted at Nephrology department of Isra University hospital Hyderabad and Medicine Department Civil hospital Nawabshah, from October 2013 to September 2014.

This is Cross-sectional study of 237 cases of chronic renal failure asses on detailed history regarding chronic renal failure, patients were subject to relevant investigations i.e. fasting lipid profile and venous blood was drawn after 14 hours fasting state in the early morning and sent to laboratory for analysis of fasting lipid profile. Inclusion Criteria were all patients of chronic renal failure with 1years history, age > 35 to 60 years and either any sex. Diagnosed on the basis of serum creatinine as define in operational definition. Exclusion Criteria were patients on lipid lowering agents, acute complication of diabetes mellitus such as diabetic keto-acidosis, patients suffering from hypothyroidism, type 1 DM and HTN, lactic-acidosis, hypoglycemia.

RESULTS

A total of 237 patients with chronic renal failure with 1years history, 113(47.68%) patients were 51 to 60 years of age, 94(39.66%) were 41 to 50 years of age and 30(12.66%) were 35 to 40 years

as presented in Fig-1.

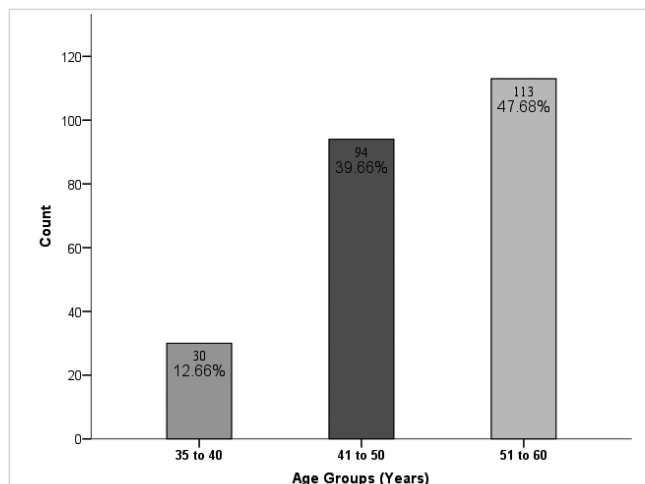


Fig. 1. Age distribution of the patients

The average age of the patients was 50.11 ± 6.94 years. There were 54.43% (129/237) male and 45.57% (108/237) female. Various lipid profile pattern of dyslipidimia in patients of chronic renal failure are presented in table-I.

Fasting lipid profile	Cutoff	Count	% age
Total cholesterol	≤ 200mg/dl		
	Yes (Normal)	132	55.7%
	No (High)	105	44.3%
Triglycerides	46-236mg/dl		
	Yes (Normal)	151	63.7%
	No (High)	86	36.3%
LDL-cholesterol	≤ 130mg/dl		
	Yes (Normal)	154	65%
	No (High)	83	35%
HDL-cholesterol	≥ 35mg/dl		
	Yes (Normal)	161	67.9%
	No (Low)	76	32.1%
VLDL-CHOLESTROLE	< 30mg/dl		
	Yes (Normal)	169	71.3%
	No (High)	68	28.7%

Table-I. Various lipid profile pattern of dyslipidimia in patients of chronic renal failure (n=237)

Results of fasting lipid profile are Triglycerides high in 86(36.3%), HDL low in 76(32.1%) cases, LDL in high 83(35%) cases, Cholesterol high in 105(44.3%) cases and VLDL-cholesterol high in 68(28.7%). Stratification analysis showed that lipid profile pattern of dyslipidemia was insignificant between male and female as presented in table-II.

FASTING LIPID PROFILE	Cutoff	n	Male n=129	Female n=108	P-Value
Total cholesterol	≤ 200mg/dl				0.203
	Yes (Normal)	132	67(51.9%)	65(60.2%)	
	No (High)	105	62(48.1%)	43(39.8%)	
	46-236mg/dl				0.38
Triglycerides	Yes (Normal)	151	79(61.2%)	72(66.7%)	
	No (High)	86	50(38.8%)	36(33.3%)	
LDL-cholesterol	≤ 130mg/dl				0.96
	Yes (Normal)	154	84(65.1%)	70(64.8%)	
	No (High)	83	45(34.9%)	38(35.2%)	
	≥ 35mg/dl				0.91
HDL-cholesterol	Yes (Normal)	161	88(68.2%)	73(67.6%)	
	No (Low)	76	41(31.8%)	35(32.4%)	
VLDL-CHOLESTROLE	< 30mg/dl				0.25
	Yes (Normal)	169	88(68.2%)	81(75%)	
	No (High)	68	41(31.8%)	27(25%)	

Table-II. Various lipid profile pattern of dyslipidimia in patients of chronic renal failure with respect to gender (n=237)

While with respect to age groups, rate of Cholesterol, triglycerides, LDL-cholesterol was high in above 40 years of age groups as compare

to 30 to 40 years of age groups. Rate of low HDL was also high in above 40 years of age groups as compare to 30 to 40 years as presented in table-III.

Fasting lipid profile	Cutoff	N	Age Groups (Years)			P-Value
			30-40 n=30	41-50 n=94	51-60 n=113	
Total cholesterol	≤ 200mg/dl					0.016
	Yes (Normal)	132	24(80%)	50(53.2%)	58(51.3%)	
	No (High)	105	6(20%)	44(46.9%)	55(48.7%)	
	46-236mg/dl					0.0005
Triglycerides	Yes (Normal)	151	30(100%)	51(54.3%)	70(61.9%)	
	No (High)	86	0(0%)	43(45.7%)	43(38.1%)	
LDL-cholesterol	≤ 130mg/dl					0.0005
	Yes (Normal)	154	29(96.7%)	72(76.6%)	53(46.9%)	
	No (High)	83	1(3.3%)	22(23.4%)	60(53.1%)	
	≥ 35mg/dl					0.0005
HDL-cholesterol	Yes (Normal)	161	30(100%)	68(72.3%)	63(55.8%)	
	No (Low)	76	0(0%)	26(27.7%)	50(44.2%)	
VLDL-CHOLESTROLE	< 30mg/dl					0.49
	Yes (Normal)	169	20(66.7%)	71(75.5%)	78(69%)	
	No (High)	68	10(33.3%)	23(24.5%)	35(31%)	

Table-III. Various lipid profile pattern of dyslipidimia in patients of chronic renal failure with respect to age groups (n=237)

DISCUSSIONS

Dyslipidemia is a frequent finding in chronic liver disease. Dyslipidemia is also seen in other illnesses like diabetes Mellitus and chronic renal failure etc. Advances in medical science have led to prolonged survival of patients with chronic diseases. Increasingly, it is known about

diseases and attempts are being made to slow the progression. Patients with chronic kidney disease are also having this advantage due to dialysis centers and are trying to protect against complications. Lipid abnormalities are common complications of CRF. They are characterized by altered levels of the individual lipids, while the total

lipids generally remaining within normal range.

In our study observed that the 47.68% upper age range was between 50 and 60 years, followed by 12.66% in between 35-40 years. One study done at Nigeria¹⁰ the reported maximum 82% percentage observed between 20 and 50 years and 18% in above 50 year, which is also supported by previous studies.^{11,12} Nigeria that cover the peak frequency of CKD between the 3rd and 4th decades. In developing countries, the frequency of CKD increases with advancing age^{13,14} and the peak incidence is found in 7th and 8th decades. The reason for disparity in peak age range among CKD patients from developed countries may be related to genetics, sociocultural factors, access to diagnostic tools, therapeutic modalities and the pattern of diseases causing CKD.^{15,16}

Our study observed 35% case in LDL, which compare with the study of Khalid Amin et al showing out of 50 patients only 2 have raised LDL while remaining of 48 patients within normal limits¹⁷. They concluded that the levels of LDL are not generally higher CRF patients. Adjustment factor, which we LDL in the CRF The change of the molecules. LDL is a heterogeneous molecule consisting of protein, fatty acid, phospholipids and cholesterol. It undergoes oxidative modifications in CRF. This results in the production of LDL-6 or a small dense LDL, atherogenic molecules. Thus, a fundamental change in LDL is not raising the same, but change the configuration and particle size.

In Triglycerides our study it was found that the higher 36.6% cases. This elevated triglycerides is consistent with other studies conducted internationally. In a study by Micheal PT Gillet and his colleagues at the Department of Biochemistry UAE University; UAE, published in 2001, found that higher triglycerides CRF¹⁸ patients be concluded J. Syrjanen colleagues.¹⁹

Our study reveals that all studied lipid profile were significantly higher for patients with chronic renal failure.^{20,21,22} These results are consistent with previous reports. He stressed the importance

of the evaluation of patients with CKD for lipid disorders experimental evidence supports the idea that the lipids directly glomerulosclerosis and tubulointerstitial injury and that the preparation of the lipid abnormalities associated with kidney disease will slow to keep chronic kidney.²³ Studies have shown strong clinical evidence that the two rise and the quality of plasma LDL is associated with accelerated atherosclerosis in CKD.²⁴ Seeing this study, as dyslipidemia occurs in all parts of the lipid profile, especially cholesterol (44.3%) contrary to previous reports where hypertriglyceridemia was normal lipid was nature.^{25,26}

CONCLUSIONS

In conclusion, our study demonstrated that dyslipidemia often our patients chronic renal failure. This underscores the need for the first test patients with lipid abnormalities early treatment can prevent cardiovascular events delay chronic kidney disease.

Copyright© 24 April, 2015.

REFERENCES

1. Amin K, Javed M, Abid M, Iqbal MN, Qayyum A. **Pattern of dyslipidemia in patients with CRF.** Professional Med J. Mar 2006;13(1):79-84.
2. Vaziri ND. **Dyslipidemia of chronic renal failure: the nature, mechanisms, and potential consequences.** Am J Physiol Renal Physiol. 2006;290:262-72.
3. Keith DS, Nichols GA, Gullion CM, Brown JB, Smith DH. **Longitudinal follow-up and outcomes among a population with chronic kidney disease in a large managed care organization.** Arch Intern Med. 2004;164:659-63.
4. McCullough PA. **Why is chronic kidney disease the "spoiler" for cardiovascular outcomes?** J Am Coll Cardiol. 2003;41:725-8.
5. Vaziri ND. **Oxidative stress in chronic renal failure. The nature, mechanism and consequences.** Semin Nephrol. 2004;24:469-73.
6. Tsimihodimos V, Dounousi E, Siamopoulos KC. **Dyslipidemia in chronic kidney disease: An approach to pathogenesis and treatment.** Am J Nephrol. 2008;28:958-73.
7. Whaley-Connell AT, Sowers JR, Stevens LA, McFarlane SI, Shlipak MG, Norris KC, et al. **Kidney early evaluation**

- program investigators: CKD in the United States: kidney early evaluation program (KEEP) and national health and nutrition examination survey (NHANES) 1999–2004.** *Am J Kidney Dis.* 2008;51(suppl 2):S13–S20.
8. Hallan SI, Coresh J, Astor BC, Asberg A, Powe NR, Romundstad S, et al. **International comparison of the relationship of chronic kidney disease prevalence and ESRD risk.** *J Am Soc Nephrol.* 2006;17:2275-84.
 9. Lewington S, Whitlock G, Clarke R, Sherliker P, Emberson J, Halsey J, et al. **Blood cholesterol and vascular mortality by age, sex, and blood pressure: a meta-analysis of individual data from 61 prospective studies with 55,000 vascular deaths.** *Lancet.* 2007;370:1829–39.
 10. Chijioke A. **Pattern of lipid profile in dialysis naive chronic kidney disease patients from ilorin, Nigeria.** *Int J Nephrol.* 2009;6(1):6.
 11. Alebiosu CO, Ayodele OO, Abbas A, Olutoyin IA. **Chronic renal failure at the Olabisi Onabanjo University Teaching Hospital, Sagamu, Nigeria.** *Afr Health Sci.* 2006;6:132-8.
 12. Akinsola A, Sanusi AA, Adekun TA, Arogundade FA. **Magnitude of the problem of chronic renal failure in Nigerians.** *African J Nephrol.* 2004;8:24-6.
 13. Feest TG, Mistry CD, Grimes DS, Mallick NP. **Incidence of advanced chronic renal failure and the need for end-stage renal replacement treatment.** *BMJ.* 1990;301:897-900.
 14. McGowan MG. **Prevalence of advanced renal failure in Northern Ireland.** *BMJ.* 1990;301:900-3.
 15. Chijioke A, Adeniyi AB. **End stage renal disease: racial differences.** *Orient J Med.* 2003;15:24-31.
 16. Adu D, Anim- Addo Y, Foli AK. **The nephrotic syndrome in Ghana: Clinical and pathological aspects.** *Quart J Med.* 1981;50:297-306.
 17. Amin K, Javaid M, Abid M. **Pattern of dyslipidemias in patients with CRF.** *Professional Med J.* Mar 2006;13(1):79-84.
 18. Micheal P. **Levels of cholesterol esters and other lipids in the plasma of patients with end-stage renal failure.** *Am J Med.* 2001;21(5):283-6.
 19. Syrjanen J, Mustonen J. **Hypertriglyceridemia and hyperuricemia are risk factors for progression of IGA nephropathy.** *Nephrol Dial Transplant.* 2000 Jan; 15(1):34-42.
 20. Mshelia DS, Buratai LB, Mamza YP. **Lipid profile in pre-dialysis chronic kidney disease patients attending University of Maiduguri Teaching Hospital, Nigeria.** *Nigerian J Clin Practice.* 2002;2:173-8.
 21. Mshelia DS, Pindiga HU. **Dyslipidaemia, Lipid oxidation and free radicals in diabetic nephropathy: an overview.** *Highland Med Res J.* 2004;2:1-7.
 22. Mshelia DS, Kadiri S, Osifo BOA. **Antioxidant vitamins in patients with chronic glomerulonephritis.** *J Life Environ Sci.* 2004;6:386-90.
 23. Keane WF, Kasiske BL, O'Dunnell MP. **The role of altered lipid metabolism in the progression of renal disease: Experimental evidence.** *AmJ Kid Dis.* 1991;17:38-42.
 24. Kennedy R, Case C, Fatti R, Johnson D, Isabel N, Marwick TH. **Does renal failure cause an Atherosclerotic milieu in patients with end-stage renal disease?** *Ann J Med.* 2001;110:198-204.
 25. Lipinska I, Curewick V. **The value of measuring percentage of high-density lipoprotein in assessing risk of cardiovascular disease.** *Arch Intern Med.* 1982;142:469-72.
 26. Atman P, Alaupovic P, Gustafson A. **Serum apolipoprotein profile of patients with chronic renal failure.** *Kid Intern.* 1997;32:368-75.

PREVIOUS RELATED STUDY

Khalid Amin, Masood Javed, Muhammad Abid, Muhammad Naeem Iqbal, Abdul Qayyam. PATTERN OF DYSLIPIDEMIA IN PATIENTS WITH CRF (Original) Prof Med Jour 13(1) 79-86 Jan, Feb, Mar, 2006.

Tufail Muhammad, Aamir Shaukat. DYSLIPIDEMIAS & CHRONIC RENAL FAILURE (Original) Prof Med Jour 9(3) 252-254 Jul, Aug, Sep, 2002.



“Patience is the art of concealing your impatience.”

Guy Kawasaki



AUTHORSHIP AND CONTRIBUTION DECLARATION

Sr. #	Author-s Full Name	Contribution to the paper	Author=s Signature
1	Dr. Ashok Kumar Lohano	Conception, design and statistical expertise,	<i>Ashok</i>
2	Dr. Irfan Ahmed Bhatti	Critical revision of the article for important intellectual content	<i>Irfan</i>
3	Dr. Azhar Iqbal	Critical revision of the article for important intellectual content	<i>Azhar</i>
4	Dr. Salman Shah Jilani	Drafting of the article	<i>Salman</i>