Oxidative stress is the primary event in hypertension and chronic renal failure; evaluation of other biochemical markers

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ABSRACT

The current study was performed in the Department of Medical Biochemistry, M.G.M. Medical College, Indore (M.P.) to assess the trend of antioxidants and some important biochemical parameters in male suffering from chronic renal failure and chronic renal failure with hypertension. The blood samples were analyzed for sodium, potassium, protein, creatinine, urea, superoxide dismutase, glutathione reductase, glutathione peroxidase, catalase and malonialdehyde. Highly significant (p<0.001) increased results were observed in case of serum potassium ions, creatinine, urea and haemolysate glutathione reductase, glutathione peroxidase and catalase were decreased highly significantly (p<0.001) in morbid groups when compared with age matched male healthy control group.

Key words: Hypertension, Chronic renal failure

INTRODUCTION

Hypertension is an important public health challenge in the United States because of its high prevalence and concomitant increased in risk of cardiovascular-renal disease1, and estimated that in the United State 50 millions have hypertension², and 20 millions have chronic renal failure³. Kidneys are vital in pathogenesis of hypertension⁴. As a consequence the prevalence and incidence of hypertension also increases with age. The relationship between age and hypertension has been consistently demonstrated in cross-sectional survey as well as in longitudinal cohort studies conducted in western population^{5,6}. Estimation of chronic kidney disease are also age dependent because chronic kidney disease was present in about 8% of the framingham population, but increased to 20% in elederly7. Chronic renal failure and Chronic renal failure with hypertension are associated with the changes in biochemical parameters in the form of sodium, potassium, protein, creatinine, urea, superoxide, dismutase,

glutathione reductase, glutathione peroxidase, catalase, and malondialdehyde which are useful tool for the diagnosis of morbid groups.

MATERIAL AND METHODS

The clinical material for present study comprised 20 patients of chronic renal failure, 20 chronic renal failures with hypertension admitted in medicine ward M Y Hospital, M.G.M. Medical College, Indore and 20 ages matched healthy control group. The age range was taken from 40 to 70 years. Blood was collected from the patients at the time of admission as well as from individuals of male healthy control group. Clinical investigations were performed in the Department of Medical Biochemistry M.G.M. Medical College, Indore (M.P.) Serum protein, creatinine, urea and superoxide dismutase were estimated by biuret, jaffe's, diacetyl monoxime, and misra H.P. et al., methods respectively. Plasma malondialdehye and haemolysate glutathione reductase, glutathione peroxidase, and catalase, were estimated by Jean

S. No.	Parameters	Healthy control group (20) mean±S.D.	Chronic renal failure (20) mean±S.D.	P-values
Biocl	nemical			
1.	Protein (mg/dl)	6.68±0.30	6.26±0.12	P<0.001
2.	Creatinine (mg/dl)	0.94±0.144	3.47±1.22	P<0.001
3.	Urea (mg/dl)	24.15±5.122	63.5±9.02	P<0.001
Elect	rolytes			
4.	Sodium (m/Eq./L.)	139.49±2.45	123.95±2.96	P<0.001
5.	Potassium (m/Eq./L.)	4.4±0.49	6.7±0.57	P<0.001
Antic	xidants			
6.	Supeoxidse dismutase (EU/gm protein/ml)	11.08±0.50	8.65±0.50	P<0.001
7.	Glutahione reductase (EU/gm protein)	19.32±0.61	16.85±0.55	P<0.001
8.	Glutathione perodixase (EU/gm Hb%)	8.5±0.48	6.87±0.43	P<0.001
9.	Catalase (EU/mg protein/ml)	5.88±0.47	3.85±0.39	P<0.001
Oxid	ant product			
10.	Malondialdehyde (nano mol/ml)	3.22±0.47	5.62±0.85	P<0.001

Table 1: Mean ± S.D. values and significant test between male healthy control v/s male chronic renal failures

P<0.001; (Highly significant)

Table 2: Mean ± S.D. values and significant test between male
healthy control v/s male chronic renal failures with hypertension

S. No.	Parameters	Healthy control group (20) mean±S.D.	Chronic renal failure (20) mean±S.D.	P-values
Bioch	nemical			
1.	Protein (mg/dl)	7.17±0.30	6.44±0.16	P<0.001
2.	Creatinine (mg/dl)	0.91±0.144	3.09±1.01	P<0.001
3.	Urea (mg/dl)	24.15±5.122	58.25±6.67	P<0.001
Elect	rolytes			
4.	Sodium (m/Eq./L.)	139.49±2.45	111.65±10.7	P<0.001
5.	Potassium (m/Eq./L.)	4.4±0.49	6.6±0.56	P<0.001
Antio	xidants			
6.	Supeoxidse dismutase (EU/gm protein/ml)	11.08±0.50	8.8±0.32	P<0.001
7.	Glutahione reductase (EU/gm protein)	19.32±0.61	16.79±0.40	P<0.001
8.	Glutathione perodixase (EU/gm Hb%)	8.5±0.48	6.76±0.30	P<0.001
9. Oxida	Catalase (EU/mg protein/ml) ant product	5.88±0.47	3.71±0.24	P<0.001
10.	Malondialdehyde (nano mol/ml)	3.22±0.47	5.44±0.52	P<0.001

P<0.001; (Highly significant)

C.D., *et al* method (1983), Horn H.D. (1963), Hafeman D.G. (1974), and Asror K Sinha method (1972) respectively. Serum electrolytes were estimated by using flame photometer. Obtained data were analyzed statistically by using student "t" test.

RESULTS

- Table number 1 and 2 are showing levels of parameters in chronic renal failure and chronic renal failure with hypertensive patients and age matched male healthy control group.
- We observed; Increased levels of serum creatinine, urea, potassium ions, and plasma malondialdehye were found to be highly significant (p<0.001) in chronic renal failure and chronic renal failure with hypertension as compared to healghy male control group.
 Other biochemical parameters such as serum protein, sodium ions, superoxide dismutase and haemolysate glutathione reductase, glutathione peroxidase and catalase were decreased highly significantly (p<0.001) in chronic renal failure and chronic renal failure with hypertensive patients when compared to age matched male healthy control group.

DISCUSSIONS

Hypertension in common in individuals with renal disease. The prevalence of hypertension varies with the causes of the underlying renal disease⁸. Hypertension is associated with higher prevalence of chronic renal disease⁹⁻¹¹. Over the last 10 to 20 years, the study of hypertension and its sequel of the at the tissue level, as apposed to the level of whole-body physiology has become increasingly common¹². Hypertension with renal disease may be due in large part to reduced renal blood flow and glomerular filtration rate. Prevalence of hypertension varied inversely with glomerular filtration rate in chronic renal failure¹³ and hypertensive subjects exhibited increased renal vacuolar resistance¹⁴. Increased oxidative stress and inflammation manifest in chronic renal failure¹⁵⁻¹⁶.

Patients with history of chronic renal failure and chronic renal failure with hypertension, the serum creatinine, urea, potassium ions and plasm malondialdehye levels were found increased¹⁷⁻¹⁹. In the present study, the results were found to be same i.e. hyperkalemia, and increased creatinine, urea and plasm malondialdehyde levels. Chronic renal failure and chronic renal failure with hypertension are characterized by decreased¹⁹⁻²⁰ sodium ions, protein, superoxide dismutase, glutathione reductase, glutathione peroxidase and catalase. In the present study, all the above stated parameters in chronic renal failure and chronic renal failure with hypertensive patients were found to be decreased when compared with normal male healthy control age matched individuals. So it is concluded that chronic renal failure and chronic renal with hypertension are associated with increased in same of the parameters on one side, and decreased in some others on the other hand.

REFERENCES

- Jaing He, Paul K Whelton. Epidemiology and prevention of hypertension. *Med Clin of N Am* 81(5): 1077 (1997).
- Chobaman A.V., Bakris, G.L., Block, H.R., et al., The seventh report of the Joint National Committee on prevention, detection, evaluation and treatment of high blood pressure, JAMA 289: 2560-72 (2003).
- 3. Coresh, J., Astar, B.C., Greene, T., et al.,

Prevalance of chronic kidney desease and decreased kidney function in the adult US population. Third National Health and Nutrition Examination Survey. *Am. J. Kidney Dis.* **41**: 1-12 (2003).

4. Burt, VL., Whelton, P., Rocella EJ, *et al.*, Prevalance of hypertensionin the US adult Population; result from the third National Health and Nutrition Examination Survey, 99 (1988), *Hypertension* **25**: 305-13 (1995).

- Joffres MR, Homet P, Rabkins, SW., et al, Prevalence control and awareness of high blood pressure among Canadian adults. Can Med Assoc J. 146: 1997-2005 (1992).
- Van Leer, E.M., Scidell, J.C., Kromhout, D. Levels and trends in blood pressure and Prevalence and treatment of hypertension in the Netherlands, 1987-1991. *Am. J. Prev. Med.* 10: 194-199 (1994).
- Culleton, B.F., Larson, M.G., Evan, J.C., *et al.* Prevalence and Correlates of elevated serum Creatinine levels. *Arch Intern Med.* 159: 1785-99 (1999).
- Malhotra, D., Schrier, R.W., Bilateral Kidney disease and hypertension. In Luscher TF, Kaplan NM (eds). Renovascular and Renal Paranchymatous Hypetension. Berlin, Springerverlay, 439-65 (1992).
- Nissension, A.R., Periera, B.J., Collins, A.J., et al. Prevalence and characteristics of individuals with chronic kidney disease in a large health maintenance organization. Am. J. Kidney. Dis. 37: 1177-83 (2001).
- Culleton, B.F., Larson, M.G., Evans, J.C., et al., Prevalence and Correlates of elevated serum Creatinine levels. *Arch Intern Med.* 159: 1785-99 (1999).
- Coresh, J., Wei, G.L., Mc. Quillian G., *et al.* Prevalence of high blood pressure and elevated serum creatinine level in the adult U S population. Third National Health and Nutrition Examination Survey (1988-1994). *Arch Inter n Med.* 161: 1207-16 (2001).
- 12. Re, R.N., The rennin-angiotension system. *Med. Clin North Am* **71**: 5 (1987).

- Muhlhauser, I., Prange, K., Sawieki, P.T., *et al.* Effect of dietary sodium on blood pressure in IDDM patients with nephropathy. *Diabetologia* 39: 212 (1996).
- Hollenberg, N.K., Sandor, T., Vasomotion of renal blood flow in essential hypertension: Oscillations in xeno in transit. *Hypertension* 6: 579 (1984).
- Shipak, M.G., Fried, L.F., Crump, C., *et al.* Elevations of inlammatory and pro coagulant biomarkers in elderly persons with renal insufficiency. *Circulation.*, **107**: 87-92 (2003).
- Oberg, B.P., Mc, Menamim, E., Lucas, F.L., et al. Increased prevalence of oxidative stress and inflammation in patients with moderate to sever chronic kidney disease. *Kidney Int* 65: 1009-16 (2004).
- Price, S., Wilson, L., Pathophysiology, Clinical concepts of disease process 6th ed. St. Louis Mo: Mosbyi (2003).
- Metheney N. Fluid and electrolyte balance 4th ed. Philadelphia Pa: Lippincot Williams and Wikkins; (2000).
- Norasatala, D., Veziri, Michael Discus, Nathum D Ho, Lalch Broujerdiord Ram K Sindhu. Oxidative stress and dysregulation of superoxide dismutase and NADPH oxidase in renal insufficiency. *Kidney International* 63: 179-185 (2003).
- Sener, Goksel, Paskaloglu Kubra, Satiroglu Henden, Alican Inci, Kacmaz Ayhan, Sackarchan Abdullah, L-carnitine amellorates oxidative damage due to chronic renal failure in rats. *J. Cardiol Pharma* 43(5): 698-705 (2004).

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