

ORIGINAL RESEARCH

A correlative study of serum beta 2 microglobulin with serum creatinine in chronic kidney disease patients

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ABSTRACT

Background: Chronic kidney disease (CKD) is a worldwide health problem, affecting millions of people. Causes of chronic kidney disease include diabetes, high blood pressure, glomerulonephritis, polycystic kidney disease, lupus, and other forms of cardiovascular diseases. β_2 microglobulin is a non-glycosylated single chain protein present in light chain of HLA-class I Antigen. It is synthesized in the body and excreted in the urine. When the glomeruli in the kidneys are damaged, they are unable to filter out β_2 microglobulin, so the level in the blood rises. Our study was aimed to assess and compare the status of β_2 microglobulin level in chronic kidney disease subjects and healthy controls. **Materials and Methods:** This was a case-control study, conducted on 220 CKD patients. Cases were selected from the Medicine OPD of Jawahar Lal Nehru Medical College and Associated Group of Hospitals, Ajmer. Age and sex-matched healthy controls (n = 100) were selected from the Medicine OPD. The study was approved by the Institutional Ethical Committee. All samples were collected under aseptic conditions from the antecubital vein. **Results:** The mean activity of serum β_2 microglobulin was significantly higher in CKD subjects as compared to healthy controls ($p < 0.0001$). Positive Pearson correlation between serum creatinine and serum β_2 microglobulin was found ($r=0.62$). **Conclusion:** Serum β_2 microglobulin level was elevated in patients with CKD and this level progressively increased with increase in creatinine. Serum β_2 microglobulin can be used as a biomarker for the early detection of CKD in the general population to prevent the morbidity and mortality which are associated with CKD. If CKD is detected early and managed appropriately the deterioration in kidney functions can be slowed and the risk of cardiovascular diseases in renal patients can be reduced.

Keywords: Chronic kidney disease (CKD), Glomerular filtration rate (GFR), Serum creatinine (SCr), Beta-2-microglobulin (β_2 M).

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INTRODUCTION

Chronic kidney disease (CKD) is a worldwide health problem, affecting millions of people.^[1] CKD represents a progressive, irreversible decline in glomerular filtration rate (GFR).^[2] CKD is a type of kidney disease in which there is a gradual loss of kidney function over a period of months or years. Causes of chronic kidney disease include diabetes, high blood pressure, glomerulonephritis, and polycystic kidney disease. Loss of renal function is common in renal failure, irrespective of the underlying cause of the kidney disease.^[3,4,5,6]

Serum Creatinine (SCr) the most routinely used marker for kidney dysfunction is often detected in normal range in the early stage of kidney dysfunction thus making diagnosis difficult. Effect of age, gender, ethnicity, nutritional status, and body muscle mass on the serum levels of creatinine are proven beyond

doubt.^[7] β_2 microglobulin is a non-glycosylated single chain protein present in light chain of HLA-class I Antigen. In the kidneys, β_2 microglobulin passes through blood-filtering units called the glomeruli and is then reabsorbed by the renal proximal tubules. Normally, only small amounts of β_2 microglobulin are present in the urine, but when the renal tubules become damaged or diseased β_2 microglobulin concentrations increase in urine due to the decreased ability to reabsorb this protein. When the glomeruli in the kidneys are damaged, they are unable to filter out β_2 microglobulin, so the level in the blood rises. β_2 microglobulin is a potentially amyloidogenic low molecular weight protein. Its serum level increases with the progression of chronic kidney disease, to reach very high concentrations in patients with chronic kidney disease.

Serum creatinine is widely accepted as the most

common parameter to assess renal functions. The prevalence of CKD in general population was 16% in India.^[8] If CKD is detected early and managed appropriately the deterioration in kidney functions can be slowed and the risk of cardiovascular diseases in renal patients can be reduced. Our study was aimed to assess and compare the status of serum β 2microglobulin in chronic kidney disease patients and healthy controls.

MATERIALS AND METHODOLOGY

This study was a case-control study, conducted on 220 chronic kidney disease patients. Patients were diagnosed as chronic kidney disease based on clinical history, physical examination & serum creatinine level. Cases were selected from the Medicine OPD of Jawahar Lal Nehru Medical College and Associated Group of Hospital, Ajmer. Age and sex-matched healthy controls (n=100) were selected from the Medicine OPD of Jawahar Lal Nehru Medical College and Associated Group of Hospital, Ajmer. The results of patients were compared with healthy controls (n=100). The study was approved by the Institutional Ethical Committee.

Inclusion criteria for the study group: Established cases of chronic kidney disease were taken.

For the control group: Age and sex-matched healthy individuals were included.

Exclusion criteria: Patients with acute kidney injury, acute infection, acute myocardial infarction, active

liver disease or liver dysfunction, hematological and malignant overt diseases. Patients with autoimmune diseases. Kidney transplant patients. Pregnant and lactating women. Patient with thyroid disorder. Patients who were unwilling to participate in the study were excluded.

Blood samples were collected after an overnight fast (12-14hrs) under aseptic conditions from all the study participants. All samples were centrifuged and analyzed for serum creatinine, and serum β 2microglobulin. Serum creatinine was measured by Jaffe’s colorimetric kinetic method.^[9] Serum β 2microglobulin was measured by Turbidimetric method.^[10]

STATISTICAL ANALYSIS

All data were analyzed by SPSS and p < 0.0001 was considered highly significant. Pearson correlation between serum creatinine and serum β 2microglobulin was performed.

RESULTS

Total 320 subject were studied. Table 1 figure 1 shows the Mean \pm SD levels of serum creatinine, β 2microglobulin are 0.88 \pm 0.42 mg/dL, 1.80 \pm 1.10 mg/L, in control group respectively. It also shows the Mean \pm SD levels of Serum creatinine, β 2microglobulin are 7.26 \pm 2.72 mg/dL, 8.30 \pm 3.12 mg/L in CKD cases respectively. Figure 2 shows positive pearson correlation between serum creatinine and serum β 2 microglobulin (r=0.62).

Table 1: Biochemical Parameters of Healthy Control Subjects v/s Chronic Kidney Disease Subjects

| Parameters | Group I Healthy subjects Mean \pm SD (n=100) | Group II Chronic Kidney Disease Patients Mean \pm SD (n=220) | P Value |
|-----------------------------------|---|---|---------------|
| Serum creatinine (mg/dL) | 0.88 \pm 0.42 | 7.26 \pm 2.72 | P<0.0001 (HS) |
| Serum Beta-2-microglobulin (mg/L) | 1.80 \pm 1.10 | 8.30 \pm 3.12 | P<0.0001 (HS) |

Figure1: Comparison of Biochemical Parameters of Healthy Control Subjects and Chronic Kidney Disease Subjects

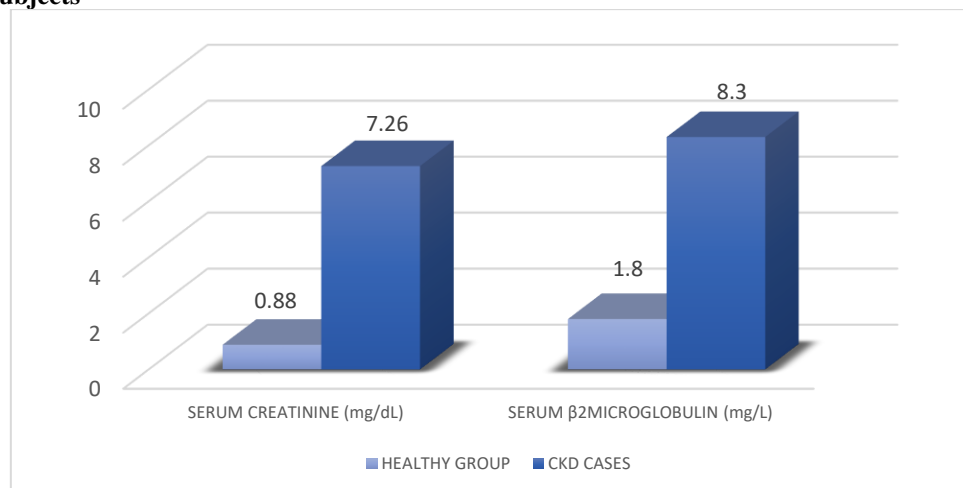
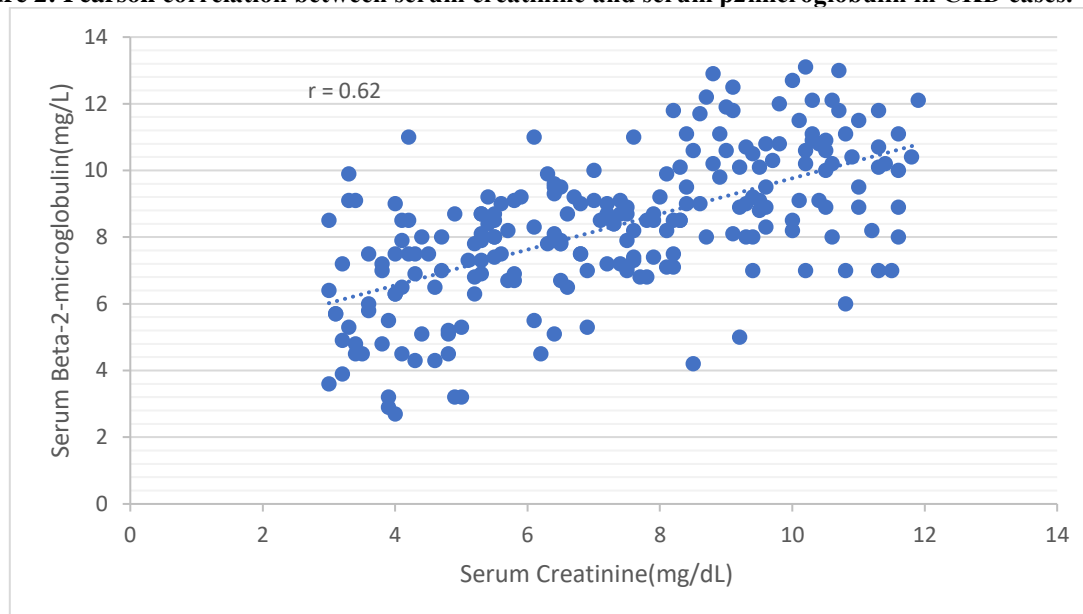


Figure 2: Pearson correlation between serum creatinine and serum β 2microglobulin in CKD cases.

DISCUSSION

Chronic kidney disease (CKD) is a worldwide health problem, affecting millions of people. Causes of chronic kidney disease include diabetes, high blood pressure, glomerulonephritis, polycystic kidney disease, lupus, and other forms of cardiovascular diseases.

Serum β 2microglobulin level was elevated in patients with CKD and this level progressively increased with increase in serum creatinine. Kidneys eliminate β 2microglobulin via glomerular filtration and tubular catabolism and hence, serum level of β 2microglobulin is highly correlated with chronic kidney disease.

In the present study the Mean \pm SD of Serum creatinine, β 2microglobulin for control was found to be 0.88 ± 0.42 mg/dL, 1.80 ± 1.10 mg/L respectively and the Mean \pm SD of Serum creatinine, β 2microglobulin for case group was found to be 7.26 ± 2.72 mg/dL, 8.30 ± 3.12 mg/L respectively. Serum creatinine and serum β 2microglobulin was significantly higher for CKD cases as compared to control group. Positive Pearson correlation between serum creatinine and serum β 2microglobulin was found ($r=0.62$)

In the kidneys, β 2microglobulin passes through blood-filtering units called the glomeruli and is then reabsorbed by the renal proximal tubules, structures that reclaim water, proteins, vitamins, minerals, and other vital substances. Normally, only small amounts of β 2M are present in the urine, but when the renal tubules become damaged or diseased, β 2M concentrations increase due to the decreased ability to reabsorb this protein. When the glomeruli in the kidneys are damaged, they are unable to filter out β 2M, so the level in the blood rises.

Our findings are in agreement with Sedighi et al who found that the mean activity of serum β 2microglobulin was significantly higher in chronic

kidney disease patients compared to the healthy subjects.^[11]

CONCLUSION

In the present study there was a significant increase in the level of serum β 2-microglobulin in CKD patients when compared to healthy controls. Present study has also revealed a positive correlation between serum creatinine and serum β 2-microglobulin in chronic kidney disease patients ($r=0.62$).

Serum β 2-microglobulin can be used as a biomarker for early detection of chronic kidney disease in general population to prevent morbidity and mortality which are associated with CKD.

REFERENCES

1. Di Angelantonio, E.; Danesh, J.; Eiriksdottir, G.; Gudnason, V. Renal Function and Risk of Coronary Heart Disease in General Populations: New Prospective Study and Systematic Review. *PLOS Med.* 2007, 4 (9), e270.
2. Brenner, B. M.; Anderson, S. The Inter Relationships Among Filtration Surface Area, Blood Pressure, and Chronic Renal Disease. *J. Cardiovasc. Pharmacol.* 1992, 19 (Suppl. 6), S1–7.
3. Liao, M.-T.; Sung, Hung, K. C.; Wu, C. C.; Lo, L.; Lu, K. C.; Hung Insulin Resistance in Patients with Chronic Kidney Disease. *J. Biomed. Biotechnol.* 2012, 2012, article ID 691369, 12 pages [doi:10.1155/2012/691369].
4. Clatworthy. Kidney Failure, Also Known as End-Stage Kidney Disease, Is a Medical Condition in Which the up to: "What Is Renal Failure?". *Johns Hopkins Medicine.* Archived from the Original on 18 June 2017. (retrieved Dec 18 2017).
5. GBD Mortality and Causes of Death, Collaborators. Global, Regional, and National Life Expectancy, All-Cause Mortality, and Cause-Specific Mortality for 249 Causes of Death, 1980–2015: A Systematic Analysis for the Global Burden of Disease Study 2015". (8 October 2016). *Lancet* 2015, 388 (1005 3), 1459–1544.

6. Ots, M.; Pechter, U.; Tamm, A. Characteristics of Progressive Renal Disease. *Clin. Chim. Acta* 2000, 297 (1–2), 29–41.
7. Woo KS, Choi JL, Kim BR, Kim JE, Han JY. Clinical usefulness of serum cystatin C as a marker of renal function. *Diabetes Metab J* 2014; 38(4): 278-284.
8. Shrestha N, Gautam S, Mishra SR, Virani SS, Dhungana RR. Burden of chronic kidney disease in the general population and high-risk groups in South Asia: A systematic review and meta-analysis. *PloS one*.2021 Oct 14;16 (10): e0258494.
9. Marakala V, Avinash SS, Shivashankar AR, Malathi M, Kumar A. Serum creatinine assay: enzymatic vs kinetic Jaffe's method. *J Evol Med Dent Sciences*, 2012; vol 1(4):328-34.
10. Sun X, Chu J, Chen P, Yang H. Comparative study on determining the level of Beta-2 microglobulin dissimilarity technique, Labelled Immunoassays and Clinical Medicine, 2001; 8(3):160-2.
11. Sedighi O, Abediankenari S; Omranifar B. Association Between PlasmaBeta-2 Microglobulin Level and Cardiac Performance in Patients WithChronicKidneyDisease. *NephroUrolMon*.2015Jan. ;7(1):e23563.