

Sclerostin and Survival Among Patients Undergoing Haemodialysis

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OBJECTIVES

Sclerostin is a protein expressed by osteocytes and has recently been shown to be a good predictor for bone formation in patients with chronic kidney disease. Serum sclerostin levels are increased in patients with end-stage renal disease and has recently been shown to be associated with aortic valve calcification in these patients. Besides, the effect of the level of sclerostin on survival is unknown.

METHODS

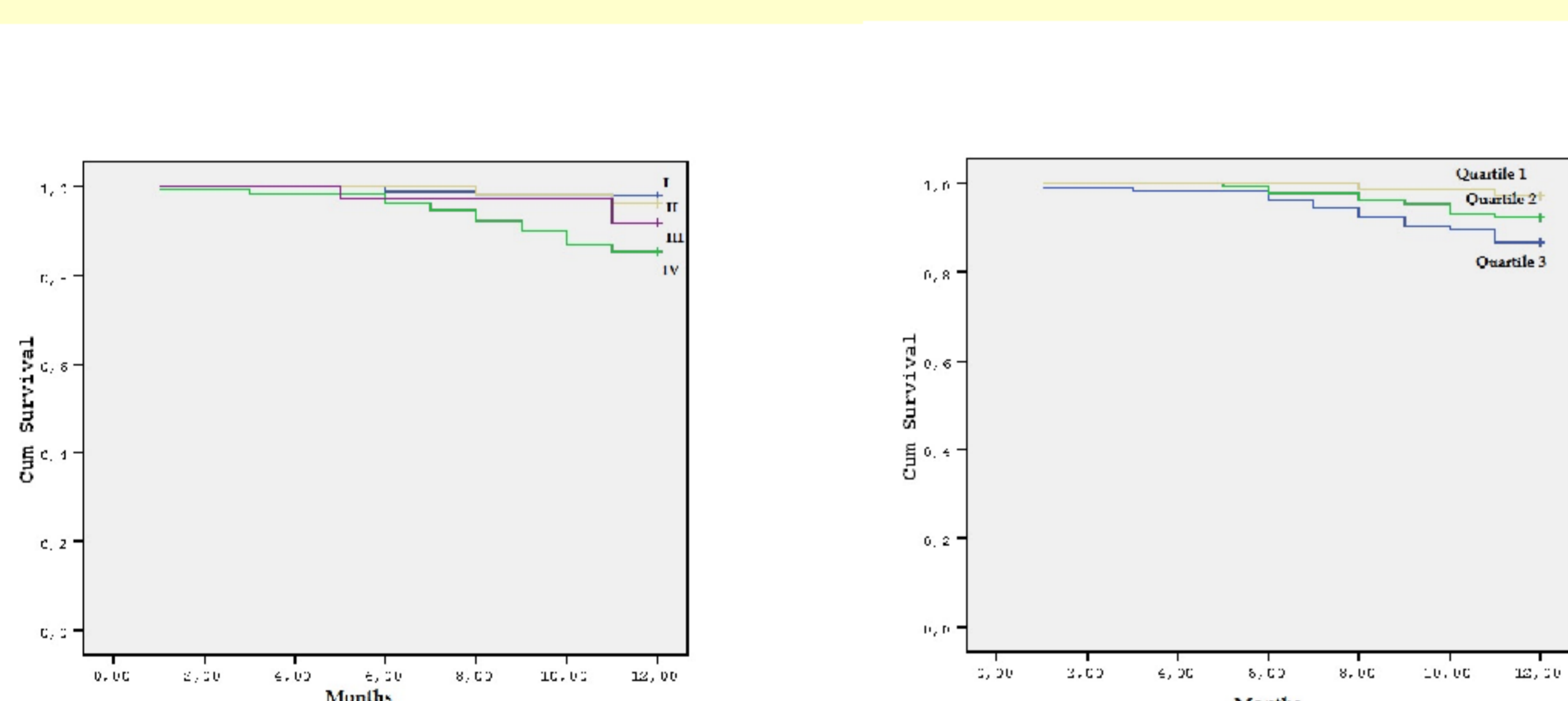
We examined survival according to serum sclerostin levels in a prospective cohort of 350 (164 males, 186 females, mean age: 57±13 years, mean hemodialysis vintage: 58±32 months) patients who were on maintenance haemodialysis treatment.

RESULTS

During follow-up of one year, 26 hemodialysis patients (7,42%) died. Patients who died were older (67,8±9 vs 56,5±13 years, p=0.013), had lower 25-hydroxy vitamin D3 (19,6±9,1 vs 29,8±11 mcg/L, p=0.024) and higher sclerostin levels (2143±1327 vs 1469±1373 pg/ml, p=0.017). Patients with 25-hydroxy vitamin D3 levels greater than median value (21,6 mcg/L; Group 1) were associated with an increase in survival when compared to patients with 25-hydroxy vitamin D3 levels greater than median value and receiving calcitriol therapy (Group 2), patients with 25-hydroxy vitamin D3 levels lower than median value and receiving calcitriol (Group 3) and finally patients with 25-hydroxy vitamin D3 levels lower than median value and not receiving calcitriol therapy (Group 4) (Log-rank: p=0.004 for the trend, Figure 1). Increased serum sclerostin quartiles are associated with decreased patient survival (Log-rank: p=0.025; Figure 2). Serum sclerostin levels in the highest quartile (>2282 pg/ml) were associated with a 22% increase in the multivariable adjusted risk of death, as compared with the lowest quartile (<370 pg/ml; adjusted also for both calcitriol therapy and serum 25-hydroxy vitamin D3 levels).

Figure 1

Figure 2



CONCLUSIONS

Increased sclerostin levels seem to be independently associated with mortality among patients who are on maintenance haemodialysis treatment.

REFERENCES:

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