Studies on the clinical significance of serum fibroblast growth factor-23 concentration in dogs and cats with chronic kidney disease

Abstract of Doctoral Thesis

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Fibroblast growth factor (FGF)-23 is a phosphaturic hormone used as an early marker of mineral metabolic disorders in chronic kidney disease (CKD) in humans. In veterinary medicine, some studies on FGF-23 have been recently reported. However, studies evaluating the clinical importance of FGF-23 in dogs and cats are limited. The present study investigated serum FGF-23 concentrations in dogs with CKD and assessed the clinical significance of increased serum FGF-23. Furthermore, this study evaluated serum FGF-23 concentrations in young and adult cats and the association between FGF-23 and hypercalcemia in cats with CKD and upper urolithiasis.

Firstly, the association between serum FGF-23 concentrations and CKD stage in dogs was investigated and compared with other phosphate metabolic markers. The results showed that serum FGF-23 concentrations in dogs with CKD increased in an earlier CKD stage compared with the serum intact parathyroid hormone and phosphorus concentration. Therefore, FGF-23 is a potential earlier marker of mineral metabolic disorders in canine CKD.

Secondly, the clinical significance of increased serum FGF-23 concentration in dogs with CKD without hyperphosphatemia was investigated. Increased serum FGF-23 concentrations were found to be significantly associated with the subsequent development of hyperphosphatemia and CKD progression. The results indicate that reducing serum FGF-23 concentrations can prevent hyperphosphatemia and CKD progression.

Thirdly, this study evaluated serum FGF-23 concentrations in young and adult cats with CKD and found that serum FGF-23 concentration increased with elevated CKD stages. Furthermore, increased serum FGF-23 concentrations were observed in an

earlier stage than serum phosphorus concentrations. Therefore, FGF-23 is also a potential early marker of mineral metabolic disorders in CKD in young and adult cats.

Finally, this study investigated whether blood calcium concentrations were related to serum FGF-23 concentrations in cats with CKD and upper urolithiasis. Increased serum FGF-23 concentrations were significantly associated with hypercalcemia independently of serum creatinine and phosphorus concentrations. Therefore, hypercalcemia is a potential cause of increased serum FGF-23 in cats.

This study demonstrated that increased serum FGF-23 concentrations in dogs with CKD occurred earlier than secondary hyperthyroidism and hyperphosphatemia and presented a risk for the subsequent development of hyperphosphatemia and CKD progression. In young and adult cats, FGF-23 was identified as an early marker of mineral metabolic disorders in CKD. Additionally, hypercalcemia was associated with increased serum FGF-23 concentrations in cats with CKD and upper urolithiasis.