

Diagnostic update

IDEXX FGF-23 Test

FGF-23 is a renal management biomarker. In cats with early CKD (IRIS* CKD Stages 1 and 2), the IDEXX FGF-23 Test provides an evidence-based approach to recommend phosphate reduction therapy.

Background

Chronic kidney disease (CKD) affects an increasing percentage of cats as they age, reported as 0.1% of cats less than 9 years old but 30%-40% of cats above age 10, and as high as 80% of cats over the age of 15.1-3 CKD causes significant morbidity and mortality in the older cat population.⁴ Kidneys are essential in phosphate homeostasis. As CKD develops and there is a decline in glomerular filtration rate (GFR), phosphorus concentrations increase, causing an imbalance in phosphate-calcium homeostasis. 5 This is labeled chronic kidney disease-metabolic bone disease (CKD-MBD; also referred to as mineral bone disorder) and describes a complex syndrome that involves fibroblast growth factor 23 (FGF-23), parathyroid hormone (PTH), 1,25-dihydroxy D₃ (1,25 vitamin D₃ calcitriol), calcium, and phosphorus (figure 1).6 CKD-MBD leads to chronically elevated FGF-23 levels in most patients. There is strong clinical evidence from both human and veterinary literature that FGF-23 often identifies mineral disruption and phosphorus overload (CKD-MBD) earlier than total serum phosphorus and is a valuable tool in the management of CKD in cats.⁷⁻¹⁰

FGF-23 has been shown to increase with CKD severity in humans and cats.⁹⁻¹¹ However, FGF-23 does not consistently precede persistent increases in kidney biomarkers (SDMA, creatinine, BUN).

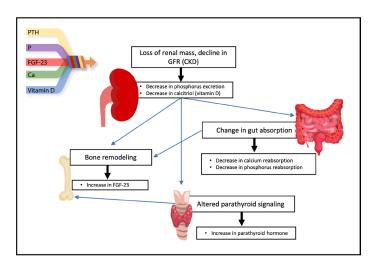


Figure 1. Simplified diagram of physiology of FGF-23 in CKD. Loss of GFR leads to a decline in phosphorus excretion, α-Klotho expression, and calcitriol production, leading to bone remodeling and increases in circulating FGF-23. These mineral imbalances, specifically calcium and phosphorus, alter gut metabolism and reabsorption, further propagating the metabolic bone disease. Due to the effect on signaling directly related to calcium reduction, a secondary increase is eventually seen in PTH, and labeled secondary renal hyperparathyroidism.

FGF-23 is not intended to be used as a core diagnostic tool to diagnose cats with CKD, but rather it serves as a useful indicator of the need for therapeutic intervention and potentially in prognosis. 10,12,13 The variability in FGF-23 levels in early CKD in cats is likely due to a combination of elements that include etiology of CKD and the complexity of MBD beyond phosphate metabolism. Phosphate metabolism has been shown to be exceptionally important in cats with all stages of CKD. Prescription renal care diets have been shown to improve quality of life and increase lifespan when fed consistently and started at early stages of CKD in cats. 3,4,14 Veterinarians are in the midst of an extensive conversation regarding the timing of diet initiation, the nature of composition of diets, and inclusion or restriction of ingredients for cats diagnosed with CKD. Further research is required to better define the specific pros and cons of therapeutic kidney diets. 8,21,29 FGF-23 offers some clarity by signaling CKD-MBD and potential phosphate overload in early-stage CKD.¹⁵ An increased FGF-23 after diagnosis of CKD supports the use of phosphate reduction therapy, the most accessible being dietary phosphate restriction. Plasma FGF-23 has been correlated to prognosis in human CKD patients, and research from veterinary literature suggests that a higher initial FGF-23 level may be a negative prognostic indicator for cats with CKD.8,16 It is likely that FGF-23 will provide similar diagnostic information about renal management in dogs, but further research is needed to confirm this. 17-19

Biology of FGF-23

FGF-23 is a phosphatonin and is likely the most important element in the control of phosphate metabolism. Primarily produced by osteocytes and osteoblasts and directed by α-Klotho expression at the level of the kidney, FGF-23 progressively rises with loss of GFR and before total serum or plasma phosphorus.²³ The relationship between FGF-23 and indirect markers, such as SDMA and creatinine (CREA), is less clear. CKD-MBD is likely related to etiology of CKD, comorbidities, and current therapies, and onset varies from cat to cat (figure 2). As phosphorus levels rise due to decreased GFR, FGF-23 increases to maintain phosphorus balance. With its coreceptor α-Klotho, FGF-23 decreases phosphorus and calcitriol in three ways: (1) downregulating sodium-phosphorus cotransporters, (2) inhibiting renal 1α-hydroxylase activity, and finally, (3) increasing 24-hydroxylase activity.²⁴⁻²⁶ In earlier-stage human CKD, FGF-23 promotes a reduction in PTH (figure 2), but in later stage disease, FGF-23 appears to contribute to renal secondary hyperparathyroidism (increasing PTH) due to decreasing levels of calcitriol and perhaps additional mechanisms yet recognized.^{27,28} This same pattern seems to be represented in studies in cats.7,9

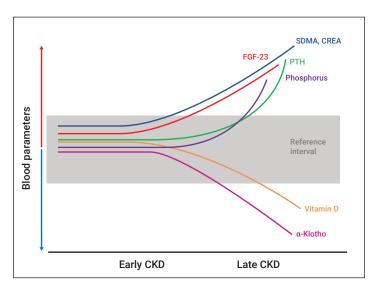


Figure 2. Diagram of relationship between FGF-23 and other important biomarkers and hormones in CKD in cats.

Clinical utility

FGF-23 should be measured in feline patients who have diagnosed, or have strongly suggested, CKD and are in earlier IRIS CKD stages (1 and 2) (figure 2).²⁰ FGF-23 is most helpful in delineating which IRIS CKD Stage 1 and Stage 2 cats will potentially benefit from phosphate reduction therapy, such as diet.¹⁵ In later-stage kidney disease, FGF-23 may not prove as beneficial as a renal management tool as diet change is always recommended unless contraindicated due to other comorbidities. Additionally, at this level of disease FGF-23 levels are likely markedly elevated.⁹

FGF-23 is the only globally available renal management marker that often identifies phosphate overload (CKD-MBD) earlier than total serum phosphorus in cats with early-stage CKD. Knowing a cat's FGF-23 level gives an evidence-based approach to several clinician and client questions: When should diet be initiated? Is this lifelong diet change needed? While diet is often suggested for patients with IRIS CKD Stage 2, it can be dependent on patient needs and the client's ability to invest in a lifelong diet change. FGF-23 supports this decision with a value indicating the medical need and value of instituting and maintaining a therapeutic renal diet. For the client, it gives an objective representation of the disease they may not yet recognize in their cat and validates a decision to commit to care that may prolong their pet's lifespan. Although the timing and nature of therapy for cats with early CKD is complex, awareness that a cat has MBD encourages early therapy, such as diet change. 21 After all, MBD is known to contribute to kidney damage though mechanisms that include vascular calcification, secondary hyperparathyroidism, and derangement of the renin-angiotensinaldosterone system (RAAS). Second, FGF-23 in early-stage CKD is a tangible piece of evidence to the owner that a diet or other therapeutic change is warranted.

FGF-23 and IDEXX SDMA

The IRIS staging guidelines for CKD include IDEXX SDMA testing and define IRIS CKD Stage 1 as SDMA levels between 15-18 µg/dL, and IRIS CKD Stage 2 as SDMA levels between 19-25 µg/dL. In cats where persistent increases in SDMA and/or other evidence (creatinine, urine specific gravity [USG]) suggest CKD, increases in FGF-23 levels indicated the presence of CKD-MBD.²² Using SDMA can allow for earlier diagnosis of CKD than traditional kidney biomarkers alone. FGF-23 as a renal management marker following earlier diagnosis can guide evidence-based decisions and validate care decisions for cats. It is possible that cats in early IRIS CKD stages will not yet have CKD-MBD and will have normal or borderline FGF-23 levels. In these cases, repeated monitoring of FGF-23, chemistry profile with IDEXX SDMA testing, and urinalysis is warranted every 3-6 months to understand when phosphate overload (CKD-MBD) has reached a level of clinical influence and phosphate reduction therapy is warranted.

IDEXX test option and when to test

IDEXX now offers an FGF-23 competitive ELISA. FGF-23 testing is indicated after diagnosis (or strong suspicion) of early CKD in cats, including IRIS CKD Stages 1 and 2. There are some comorbidities that research would suggest impact FGF-23 levels including uncontrolled hyperthyroidism, cardiac disease, moderate to severe systemic inflammation and/or neoplasia, lytic bone lesions, and profound anemia.^{3,30-32} Currently avoiding testing FGF-23 in patients with these disease patterns is recommended. Measuring FGF-23 is redundant in cats who already have total serum phosphorus above 4.6 mg/dL.

Interpretation of results

FGF-23 ≤ 299 pg/mL is within normal levels: No evidence of CKD-MBD. This is not an indication that CKD is not present, but only that FGF-23 has not increased out of the expected range for cats without CKD-MBD and that therapy to address phosphorus levels is likely not indicated at this time. If other indications exist, such as significant, stable increases in kidney biomarkers, proteinuria, or acid-base disturbance, intervention to support disease is likely still appropriate.

FGF-23 300–399 pg/mL is borderline: FGF-23 is higher than expected but not at a level that clearly indicates the need for targeted therapy. It is appropriate to institute CKD therapies indicated by other diagnostics or clinical context. Repeating FGF-23 testing in 3–6 months, alongside a chemistry profile with IDEXX SDMA testing and urinalysis, is recommended to monitor for progression and development of CKD-MBD, which would warrant intervention.

FGF-23 ≥ 400 pg/mL is increased: Targeted therapy to reduce phosphorus overload is warranted alongside all other indicated CKD therapies.



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For cats with early-stage CKD and normal or borderline FGF-23 levels, repeating FGF-23 testing with semiannual to annual lab work is indicated. CKD-MBD is not always linearly related to functional kidney biomarkers, such as SDMA and creatinine, and increases in FGF-23 may occur without changes to kidney function. Therefore, it is important to monitor onset of phosphorus overload and treat accordingly.

Follow up

Research would suggest that FGF-23 decreases after initiation of kidney appropriate diets and/or reduction in phosphorus intake. Including monitoring of FGF-23 on chemistry profiles and kidney recheck profiles of animals under therapy may prove advantageous in understanding response to therapy.⁹

Ordering information

| Test code | Test name and contents | Specimen requirements |
|------------------|---|---|
| 8988 | IDEXX FGF-23 Test—Feline | 1 mL serum |
| 89909999
8990 | Renal Monitoring Profile with IDEXX FGF-23 Test—Feline Renal Panel, IDEXX CBC*, IDEXX FGF-23 Test With IDEXX CBC-Select® | 2 mL serum, 1 mL LTT |
| 89919999
8991 | Renal Monitoring Profile, Comprehensive with IDEXX FGF-23 Test, UA, and Reflex UPC (if Indicated)—Feline Renal Panel, IDEXX CBC®, IDEXX FGF-23 Test, urinallysis with reflex urine protein:creatinine (UPC) ratio if indicated With IDEXX CBC-Select® | 2 mL serum, 1 mL LTT,
5 mL urine in a sterile
container |

Note: Renal Panel includes the following: Albumin, albumin:globulin ratio, BUN, BUN:creatinine ratio, calcium, chloride, cholesterol, creatinine, globulin, phosphorus, potassium, IDEXX SDMA® Test, sodium; potassium ratio, total protein, anion gap, TCO₂ (bicarbonate).

Additional specimen requirements

Only domestic feline[†] serum specimens will be accepted for FGF-23 measurement. Specimen refrigeration is optimal; freezing is not necessary.

Turnaround time: 2-4 days

Contacting IDEXX

Laboratory Customer Support

If you have any questions regarding test codes, turnaround time, or pricing, please contact our Laboratory Customer Support Team at 1-888-433-9987.

Expert feedback when you need it

Our medical specialty consulting service is available for expert and complimentary consultation. Please call 1-888-433-9987 if you have questions.



References

- 1. Conroy M, Brodbelt DC, O'Neill D, Chang YM, Elliott J. Chronic kidney disease in cats attending primary care practice in the UK: a VetCompass study. *Vet Rec.* 2019;184(17):526. doi:10.1136/vr.105100
- 2. Marino CL, Lascelles BD, Vaden SL, Gruen ME, Marks SL. Prevalence and classification of chronic kidney disease in cats randomly selected from four age groups and in cats recruited for degenerative joint disease studies. *J Feline Med Surg.* 2014;16(6):465–472. doi:10.1177/1098612X13511446
- 3. Sparkes AH, Caney S, Chalhoub S, et al. ISFM consensus guidelines on the diagnosis and management of feline chronic kidney disease. *J Feline Med Surg.* 2016;18(3):219–239. doi:10.1177/1098612X16631234
- 4. White JD, Malik R, Norris JM. Feline chronic kidney disease: can we move from treatment to prevention? *Vet J.* 2011;190(3): 317–322. doi:10.1016/j.tvjl.2010.12.011
- 5. Slatopolsky E. The intact nephron hypothesis: the concept and its implications for phosphate management in CKD-related mineral and bone disorder. *Kidney Int Suppl.* 2011;79(121):S3–S8. doi:10.1038/ki.2011.23
- 6. Moe S, Drüeke T, Cunningham J, et al. Definition, evaluation, and classification of renal osteodystrophy: a position statement from Kidney Disease: Improving Global Outcomes (KDIGO). *Kidney Int*. 2006;69(11):1945–1953. doi:10.1038/sj.ki.5000414
- 7. Finch NC, Geddes RF, Syme HM, Elliott J. Fibroblast growth factor 23 (FGF-23) concentrations in cats with early nonazotemic chronic kidney disease (CKD) and in healthy geriatric cats. *J Vet Intern Med.* 2013;27(2):227–233. doi:10.1111/jvim.12036
- 8. Geddes RF, Elliott J, Syme HM. Relationship between plasma fibroblast growth factor-23 concentration and survival time in cats with chronic kidney disease. *J Vet Intern Med*. 2015;29(6):1494–1501. doi:10.1111/jvim.13625
- 9. Geddes RF, Finch NC, Elliott J, Syme HM. Fibroblast growth factor 23 in feline chronic kidney disease. *J Vet Intern Med.* 2013;27(2):234–241. doi:10.1111/jvim.12044
- 10. Seiler S, Heine GH, Fliser D. Clinical relevance of FGF-23 in chronic kidney disease. *Kidney Int Suppl.* 2009;76(114):S34-S42. doi:10.1038/ki.2009.405
- 11. Liao YL, Chou CC, Lee YJ. The association of indoxyl sulfate with fibroblast growth factor-23 in cats with chronic kidney disease. *J Vet Intern Med.* 2019;33(2):686–693. doi:10.1111/jvim.15457
- 12. Lin J, Lin L, Chen S, Yu L, Chen S, Xia Z. Serum fibroblast growth factor 23 (FGF-23): associations with hyperphosphatemia and clinical staging of feline chronic kidney disease. *J Vet Diagn Invest*. 2021;33(2):288–293. doi:10.1177/1040638720985563
- 13. Nakata J, Nakahari A, Kato Y, et al. Molecular cloning and expression analysis of feline α1-microglobulin. *Vet Immunol Immunopathol*. 2011;139(1):79–82. doi:10.1016/j.vetimm.2010.08.002
- 14. 14. Elliott J, Rawlings JM, Markwell PJ, Barber PJ. Survival of cats with naturally occurring chronic renal failure: effect of dietary management. *J Small Anim Pract*. 2000;41(6):235–242. doi:10.1111/j.1748-5827.2000.tb03932.x
- 15. Geddes RF, Elliott J, Syme HM. The effect of feeding a renal diet on plasma fibroblast growth factor 23 concentrations in cats with stable azotemic chronic kidney disease. *J Vet Intern Med*. 2013;27(6):1354–1361. doi:10.1111/jvim.12187
- 16. Chang YH, Wu CH, Chou NK, et al. High plasma C-terminal FGF-23 levels predict poor outcomes in patients with chronic kidney disease superimposed with acute kidney injury. *Ther Adv Chronic Dis.* 2020;11:2040622320964161. doi:10.1177/2040622320964161
- 17. Harjes LM, Parker VJ, Dembek K, et al. Fibroblast growth factor-23 concentration in dogs with chronic kidney disease. *J Vet Intern Med*. 2017;31(3):784–790. doi:10.1111/jvim.14707
- 18. Miyakawa H, Hsu HH, Ogawa M, Akabane R, Miyagawa Y, Takemura N. Association between serum fibroblast growth factor-23 concentration and development of hyperphosphatemia in normophosphatemic dogs with chronic kidney disease. *J Vet Intern Med.* 2021;35(5):2296–2305. doi:10.1111/jvim.16237
- 19. Rudinsky AJ, Harjes LM, Byron J, et al. Factors associated with survival in dogs with chronic kidney disease. *J Vet Intern Med.* 2018;32(6):1977–1982. doi:10.1111/jvim.15322
- 20. International Renal Interest Society. Guidelines: IRIS staging of CKD. www.iris-kidney.com/guidelines/staging.html. Accessed August 29, 2022.
- 21. Foster JD. Update on mineral and bone disorders in chronic kidney disease. *Vet Clin North Am Small Anim Pract*. 2016;46(6): 1131–1149. doi:10.1016/j.cvsm.2016.06.003
- 22. Sargent HJ, Jepson RE, Chang YM, Biourge VC, Bijsmans ES, Elliott J. Fibroblast growth factor 23 and symmetric dimethylarginine concentrations in geriatric cats. *J Vet Intern Med.* 2019;33(6):2657–2664. doi:10.1111/jvim.15590



- 23. Drüeke TB. Hyperparathyroidism in chronic kidney disease. In: Feingold KR, Anawalt B, Boyce A, et al, eds. *Endotext* [online textbook]. MDText.com, Inc. Updated October 18, 2021. Accessed August 29, 2022. Available from: www.ncbi.nlm.nih.gov/books/NBK278975
- 24. Gattineni J, Baum M. Regulation of phosphate transport by fibroblast growth factor 23 (FGF23): implications for disorders of phosphate metabolism. *Pediatr Nephrol.* 2010;25(4):591–601. doi:10.1007/s00467-009-1273-z
- 25. Marks J, Debnam ES, Unwin RJ. Phosphate homeostasis and the renal-gastrointestinal axis. *Am J Physiol Renal Physiol*. 2010;299(2):F285–F296. doi:10.1152/ajprenal.00508.2009
- 26. Ramon I, Kleynen P, Body JJ, Karmali R. Fibroblast growth factor 23 and its role in phosphate homeostasis. *Eur J Endocrinol*. 2010;162(1):1–10. doi:10.1530/EJE-09-0597
- 27. Isakova T, Wahl P, Vargas GS, et al. Fibroblast growth factor 23 is elevated before parathyroid hormone and phosphate in chronic kidney disease [published correction appears in *Kidney Int.* 2012;82(4):498]. Kidney Int. 2011;79(12):1370–1378. doi:10.1038/ki.2011.47
- 28. Silver J, Naveh-Many T. FGF-23 and secondary hyperparathyroidism in chronic kidney disease. *Nat Rev Nephrol.* 2013;9(11): 641–649. doi:10.1038/nrneph.2013.147
- 29. Laflamme D, Backus R, Brown S, et al. A review of phosphorus homeostasis and the impact of different types and amounts of dietary phosphate on metabolism and renal health in cats. *J Vet Intern Med*. 2020;34(6):2187–2196. doi:10.1111/jvim.15961
- 30. Williams TL, Elliott J, Syme HM. Calcium and phosphate homeostasis in hyperthyroid cats: associations with development of azotaemia and survival time. *J Small Anim Pract*. 2012;53(10):561–571. doi:10.1111/j.1748-5827.2012.01253.x
- 31. Edmonston D, Wolf M. FGF23 at the crossroads of phosphate, iron economy and erythropoiesis. *Nat Rev Nephrol.* 2020;16(1):7–19. doi:10.1038/s41581-019-0189-5
- 32. Song T, Fu Y, Wang Y, et al. FGF-23 correlates with endocrine and metabolism dysregulation, worse cardiac and renal function, inflammation level, stenosis degree, and independently predicts in-stent restenosis risk in coronary heart disease patients underwent drug-eluting-stent PCI. *BMC Cardiovasc Disord*. 2021;21(1):24. doi:10.1186/s12872-020-01839-w