

Cystatin C





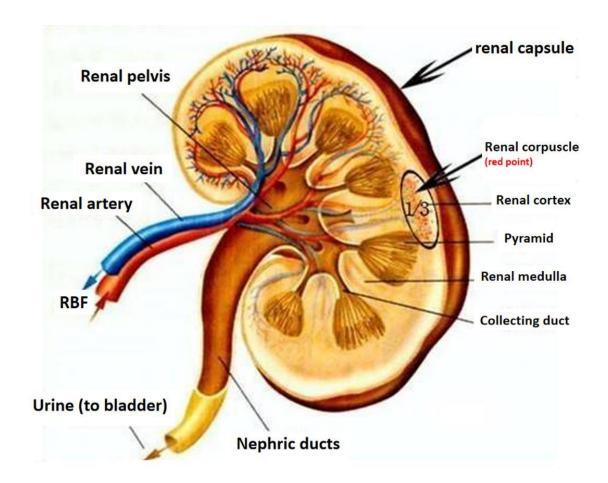
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a. Renal Structure

The kidneys are a pair of bean shaped organs and are a reddish-brown colour. Each kidney is made up of more than a million nephrons. Each nephron consists of glomerulus, renal vesicles and tubules, and glomerulus and renal vesicles form renal corpuscles.





a. Renal Function

Water

Keep the water balance in the body.

Bones

The kidney is an important activation site for vitamin D and regulates calcium absorption.

Heart

The balance of electrolytes (potassium, sodium and calcium) depends on normal kidney function, imbalance can lead to arrhythmias.

pH-balanced

Regulate pH in the body.

Blood Pressure

Maintain the stability of blood pressure.

Metabolic Waste

Remove uric acid, urea, toxins and other metabolic wastes.

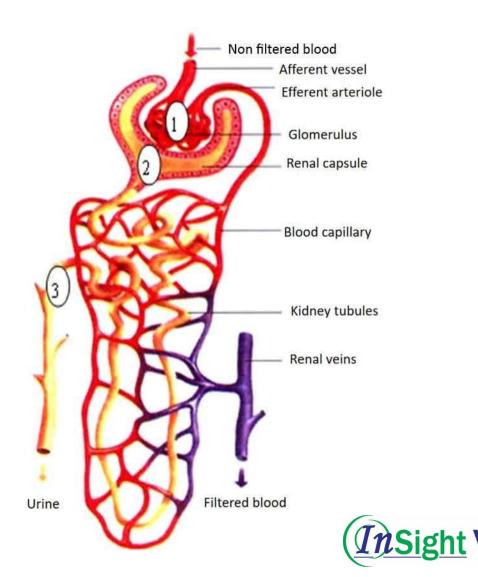
Blood

The release of erythropoietin (EPO) encourages the bone marrow to produce more red blood cells.

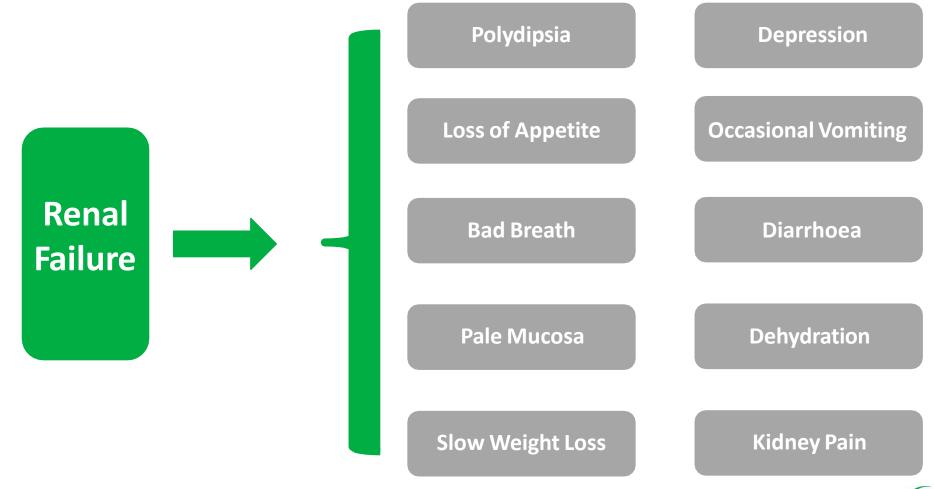


a. Schematic Diagram of Urine Formation

Urine production depends on glomerular filtration and reabsorption and secretion of renal tubules and collecting ducts.



b. Clinical Symptoms of Renal Failure





b. Aetiology of Renal Failure

Infection

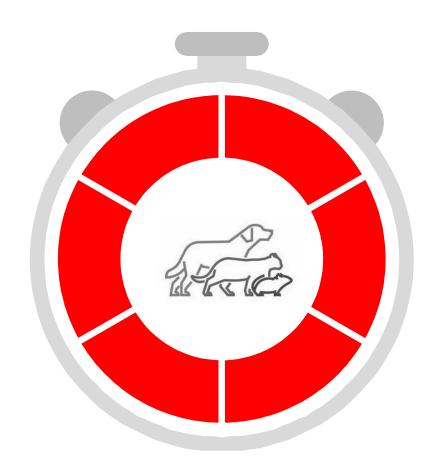
pyosis of uterus, pyelonephritis, babesiosis, urinary tract infection

Trauma

bladder splitting and kidney damage caused by injury

Poisoning

ingesting rat poison, heavy metals etc.



Tumours

urinary system tumors, renal lymphoma etc.

Metabolism

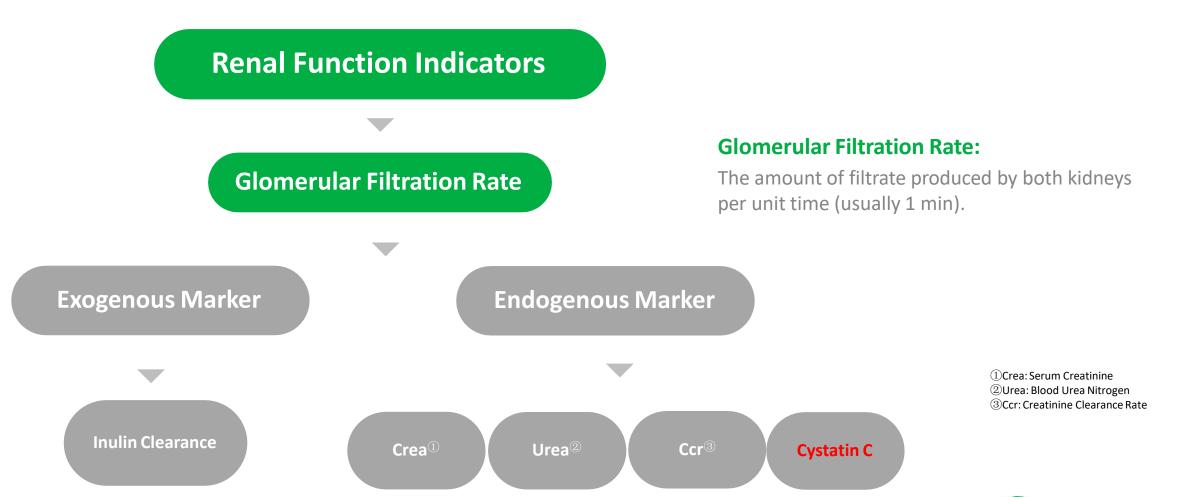
adrenal cortical dysfunction, diabetes etc.

Diet

poor eating habits and poor quality cat/dog food



c. Traditional Indicators of Renal Function





c. Traditional Indicators of Renal Function

shortcoming

The determination of exogenous marker renal clearance is regarded as the "gold standard" for GFR evaluation.

High material cost

Isotopically labelled substances are involved in radiation exposure

Specimen collection and experimental operation are complicated

The timeliness of critical patient testing cannot be achieved



c. Traditional Indicators of Renal Function

Ideal endogenous markers should have:

- ☐ Stable generation rate
- Stable blood concentration, not affected by other pathological changes, does not bind with protein
- ☐ Glomerular free filtration
- ☐ Renal tubules do not secrete or reabsorb
- ☐ No external renal clearance



It cannot meet the requirement of endogenous GFR marker and is greatly affected by opportunistic disease conditions. There is obvious passive reabsorption in renal tubules.



Inconvenient 24h urine collection leads to inaccurate urine volume. When GFR was low, the renal tubules compensatively secreted more creatinine, leading to greater overestimation of GFR in Ccr.



Creatinine levels are influenced by many factors, including age, sex, body shape, height, muscle mass and diet. Renal tubules secrete creatinine at different times in the same individual and at different rates.



Cystatin C

Also known as y2 trace basic protein or post-gamma globulin, is one of the cysteine protease inhibitor proteins.





Can be continuously transcribed and expressed at a constant rate in all nucleated cells without tissue specificity.

It is not affected by age, sex, weight, inflammation and other factors.









Can filter freely from the glomerulus and is completely reabsorbed by the epithelial cells.

At the same time, renal tubular epithelial cells do not secrete Cystatin C into the lumen.

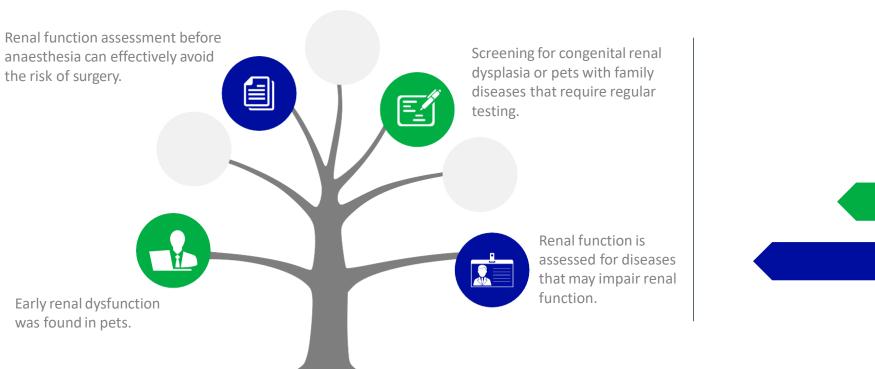




Cystatin C can vary greatly within an individual but little between individuals.



Clinical Application of Cystatin C



Health examination

Genetic screening

Pre-operative examination

Renal function assessment



In J.-P. BRAUN's research, the coefficients of correlation between Pl-Cystatin C and Pl-Urea or Pl-Creatinine were high in dogs with kidney diseases, and in all (diseased & healthy) dogs, ranging from r =0.78 to r = 0.88 (Fig. 1). In dogs with kidney diseases, there was a linear relationship between Pl-Cystatin C and log Pl-Creatinine or log Pl-Urea.

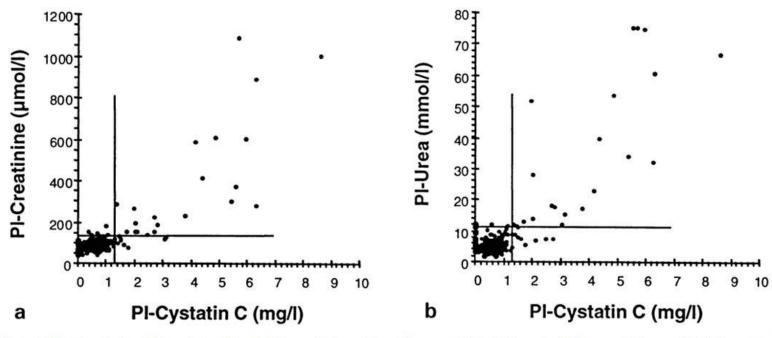


Fig. 4. Scatterplots of Pl-cystatin C and Pl-creatinine (a) or Pl-urea (b) in 179 control dogs and dogs with kidney diseases. (Lines are upper limits of the reference intervals.)



In Naoki IWASA's study, all the dogs were divided into two groups; one group, which accepted the offer to start regular follow-up visits every 2 to 4 weeks (Clinical pathology-based monitoring group; CPBM), and another group, which refused the regular follow-up visits (symptom-based monitoring group; SBM).

The survival period in the CPBM group (median 441 days) was significantly longer than the survival period of the SBM group (median 262 days) (P<0.05) (Fig. 2). These dogs were expected to have a significant decrease in renal function because high serum Cystatin C concentration reflects a reduced GFR, even though serum Crea concentration levels could not detect the reduced GFR.

Results from the present study suggest that the early treatment intervention by CPBM of asymptomatic dogs with normal serum Crea concentration levels could extend the prognosis of chronic kidney disease if serum Cystatin C concentrations are high.

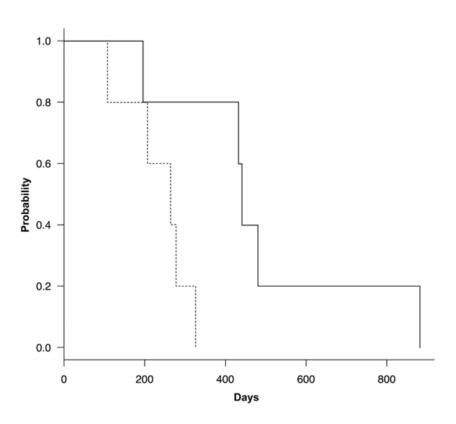


Fig. 2. Kaplan–Meier survival curves for the Clinical pathology-based monitoring (CPBM) group and symptom-based monitoring (SBM) group. Solid line: the CPBM group. Dashed line: the SBM group.



In IWONA POŚWIATOWSKA–KASZCZYSZYN's study, a plasma tracer elimination curve was generated and clearance was calculated by dividing the injected dose by the area under the curve (AUC), estimated using a one-compartment pharmacological model. The obtained values of Pcio were standardised according to body weight. GFR, estimated basing on Pcio, was applied to compare Cystatin C, Creatinine, and Urea data as exponents of renal function in cats. In data analysis by ROC method (Fig. 3), the area below the curve (AUC) for particular parameters amounted: 0.854 for plasma clearance of iohexol (Pcio) within 95% credibility interval; 0.819 for Cystatin C in 95% credibility interval; 0.744 for Creatinine in 95% credibility interval; 0.689 for Urea in 95% credibility interval.

Analysis of the obtained results, based on ROC curve, proved that Cystatin C seems to be a more advantageous parameter than Creatinine and Urea in the assessment of renal function diagnosis in cats.

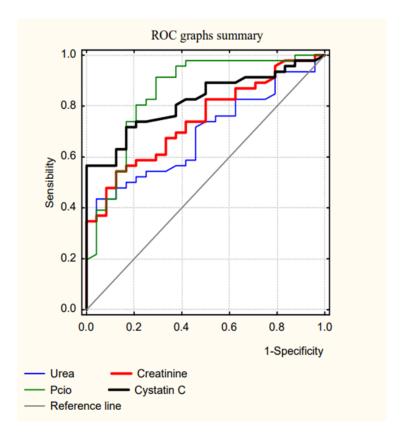


Fig. 3. Graphical expression of sensibility and specificity of four examined methods. Diagram presents the measurement of plasma clearance of iohexol (Pcio) and serum concentrations of urea, creatinine, and cystatin C.

e. SDMA

Some manufacturers have launched SDMA tests as an indicator for kidney function in animals. However, there is an argument to suggest that if SDMA is tested alone, without measuring the concentration of ADMA and amino acids, then the result is not useful to the veterinarian. This is because metharginine, whether in free form or in proteins, must be detected by hydrolysis. Only on the premise of understanding the synthesis and metabolism of the individual, combined with the concentration of ADMA and amino acids can the concentration of SDMA be analysed and the meaning behind the result understood.

For this reason, some papers show that Cystatin C is a more effective renal function indictor than SDMA.

Reference: J. Kim et al (2020) Biomarkers for Chronic Kidney Disease in Dogs: A Comparison Study J. Vt med sci, 82(8) 1130-1137.



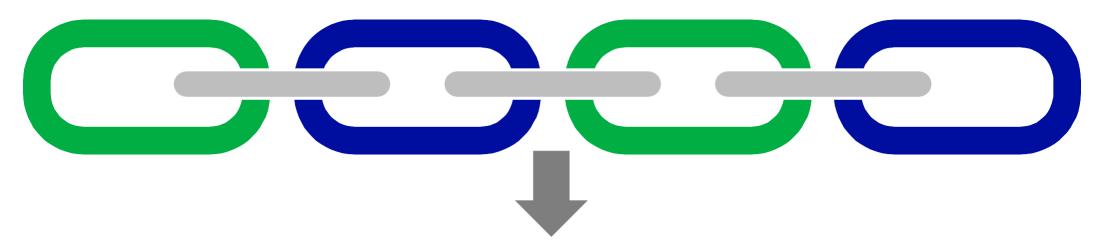
f. Conclusion

A. Endogenous marker is better than Exogenous marker in operation and cost.

B. In terms of disease identification ability, Cystatin C is superior to Urea, Crea and Ccr.

C. It is more suitable for highly specific immunochromatographic kit for Cystatin C in dogs and cats.

D. The market is more accepting of Cystatin C tests for kidney disease.



Cystatin C is the best choice!



(InSight V-IA

Canine Cystatin C (cCys C) Rapid Quantitative Test

Woodley have developed a rapid, accurate and reliable, highly sensitive detection method for Canine Cystatin C.

The InSight V-IA Canine Cystatin C (cCys C) Rapid Quantitative Test is a fluorescence immunoassay used with the InSight V-IA Veterinary Immunoassay Analyser for quantitative determination of cCys C concentration in canine serum or plasma.

The test is used as an aid to evaluate kidney injury.

It can be stored at room temperature.









Insight V-IA

Feline Cystatin C (fCys C) Rapid Quantitative Test

Woodley have developed a rapid, accurate and reliable, highly sensitive detection method for Feline Cystatin C.

The InSight V-IA Feline Cystatin C (fCys C) Rapid Quantitative Test is a fluorescence immunoassay used with the InSight V-IA Veterinary Immunoassay Analyser for quantitative determination of fCys C concentration in feline serum or plasma.

The test is used as an aid to evaluate kidney injury.

It can be stored at room temperature.









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Thank You



