1. Staging of CKD (based on plasma creatinine concentration)

Staging is undertaken **following diagnosis of CKD** in order to facilitate appropriate treatment and monitoring of the patient. There are separate but related algorithms for staging CKD in cats and dogs.

Staging is based initially on fasting plasma creatinine, assessed on at least two occasions in the stable patient. The patient is then substaged based on proteinuria and blood pressure.

Based on these categories, some empirical recommendations can be made about the type of treatment it would be logical to use for these cases. In addition, predictions based on clinical experience might be made about the likely response to treatment.

Stage	Plasma creatinine		Comments	
	μmol/l			
	mg/dl			
	Dogs	Cats		
-	<125	<140	At risk of CKD	
	<1.4	<1.6	For patients identified as 'at risk' consider regular	
			screening and taking steps to reduce risk factors	
			Non-azotemic	
			Some other renal abnormality present e.g.	
1	<125	<140	inadequate concentrating ability without identifiable	
	<1.4	<1.6	non-renal cause; abnormal renal palpation and/or	
			abnormal renal imaging findings; persistent	
			proteinuria of renal origin; abnormal renal biopsy	
			results, progressively elevating creatinine levels	
			Mild renal azotemia [lower end of the range lies	
	405 470	440 040	within the reference range for many labs but the	
2	125 - 179	140 - 249	insensitivity of creatinine as a screening test means	
	1.4 - 2.0	1.6 - 2.8	that animals with creatinine values close to the	
			upper limit of normality often have excretory failure]	
			Clinical signs usually mild or absent	
2	100 120	250 420	Madayata yanglarataamia	
3	180 - 439	250 - 439	Moderate renal azotaemia	
	2.1 - 5.0	2.9 – 5.0	Systemic clinical signs may be present	
4	>440	>440	Severe renal azotaemia	
•	>5.0	>5.0	Systemic clinical signs are usually present	
			cyclec cca. e.g.ic a.c acaa, p. cooin	

Note these plasma creatinine levels apply to average size dogs – those of extreme size may vary

2a. Substaging by Proteinuria

The goal is to identify renal proteinuria having ruled out post-renal and pre-renal causes.

Standard urine dipsticks can give rise to false positives therefore practitioners should consider using a more specific screening test such as the sulphosalicylic acid turbidometric test or the ERD® test.

The urine protein to creatinine (UP/C) ratio should be measured in all cases, provided there is no evidence of urinary tract inflammation or hemorrhage and the routine measurement plasma proteins has ruled out dysproteinemias. Ideally staging should be done on the basis of at least three urine samples collected over a period of at least 2 weeks.

UPC	value	Substage	
Dogs	Cats		
<0.2	<0.2	Non-proteinuric (NP)	
0.2 to 0.5	0.2 to 0.4	Borderline proteinuric (BP)	
>0.5	>0.4	Proteinuric (P)	

Patients with persistent proteinuria in the BP subcategory should be re-evaluated within 2 months and re-classified as appropriate.

UP/Cs in the NP or BP range may be categorized as 'microalbuminuric' on the ERD® test. The significance of micoralbuminuria in predicting future renal health is not understood at present. IRIS recommendation is to continue to monitor this level of proteinuria.

Proteinuria may decline as renal dysfunction worsens and so may be less frequent in animals in stages 3 and 4.

Response to any treatment given to reduce glomerular hypertension, filtration pressure, and proteinuria, should be monitored at intervals using the UP/C ratio.

2b. Substaging by Arterial Blood Pressure

Patients should be acclimatized to the measurement conditions and multiple measurements taken. The final classification should rely upon multiple pressure determinations (preferably multiple patient visits to the clinic on separate days but acceptable if during the same visit with at least 2 hours separating determinations).

Patients are substaged by blood pressure according to the degree of risk of end organ damage, and whether there is evidence of end-organ damage or complications.

Systolic BP mm Hg	Diastolic BP mm Hg	Adaptation when breed- specific reference range is available *	Arterial Pressure Substage (AP)
<150	<95	<10 mm Hg above reference range	0 Minimal Risk
150 – 159	95 - 99	10 – 20 mm Hg above reference range	1 Low Risk
160 – 179	100 - 119	20 – 40 mm Hg above reference range	2 Moderate Risk
≥ 180	≥ 120	≥ 40 mm Hg above reference range	3 High Risk
No eviden	ce of end organ	No complications (nc)	
	of end organ o	Complications (c)	
E	Blood pressure	Risk not determined (RND)	

As with proteinuria, in the absence of evidence of end organ damage, demonstration of persistence of blood pressure readings within a particular category is important.

'Persistence' of elevation should be judged on multiple blood pressure measurements made over the following timescales:

- 2 months (if at moderate risk 160 to 179 mm Hg systolic BP)
- 1 to 2 weeks (if at severe risk ≥180 mmHg).

*If available, it is preferable to use breed specific ranges for normal values and compare the measurement to the upper limit of the normal range for the breed being evaluated. Sight hounds, in particular, have a higher reference range than most breeds of dog.

3 Therapy Effect

If antihypertensive/antiproteinuric therapy is instigated, subsequent staging of hypertension/proteinuria should be based on the **current actual** blood pressure/UPC with **(T)** to indicate that this level reflects the effects of therapy,

For example:

Cat A before treatment

Creatinine 260 UPC 0.3 Systolic blood pressure 200 IRIS stage 3, BP, AP3

Cat A after treatment

Creatinine 300 UPC 0.3 Systolic blood pressure 155 IRIS stage 3, BP, AP1 (T)

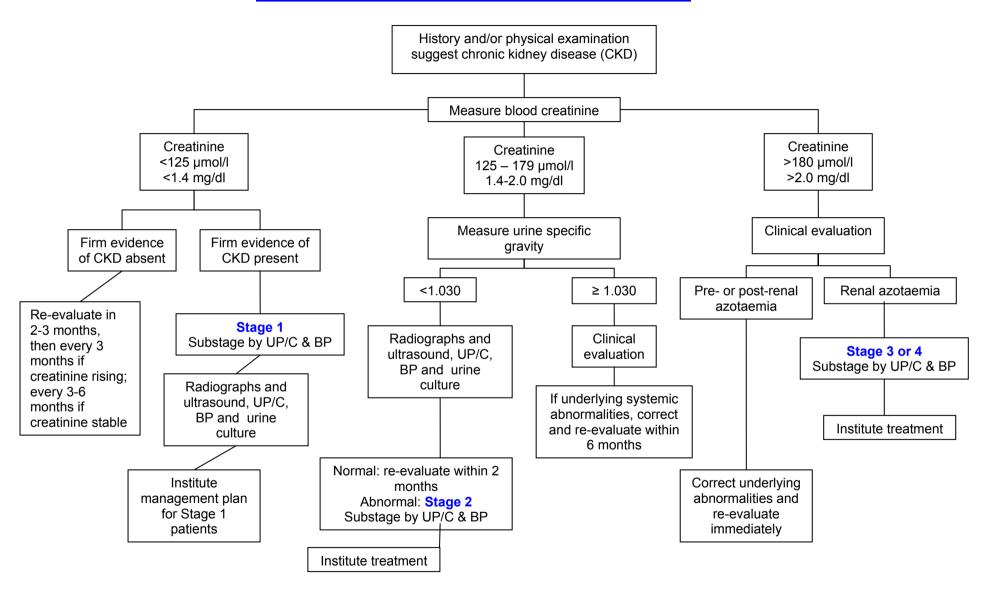
Dog B before treatment

Creatinine 160 UPC 0.8 Systolic blood pressure 155 IRIS stage 2, P, AP1

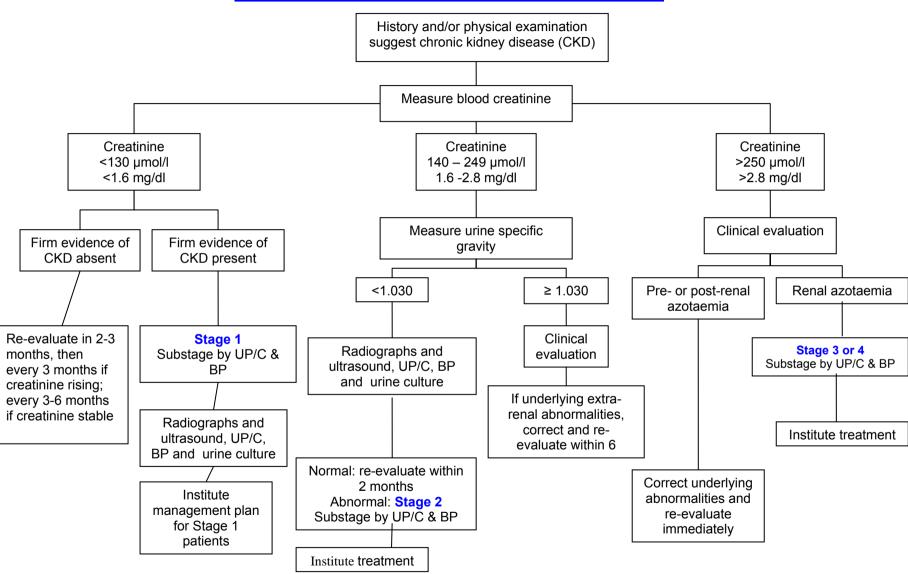
Dog B after treatment

Creatinine 170 UPC 0.4 Systolic blood pressure 155 IRIS stage 2, BP (T), AP1

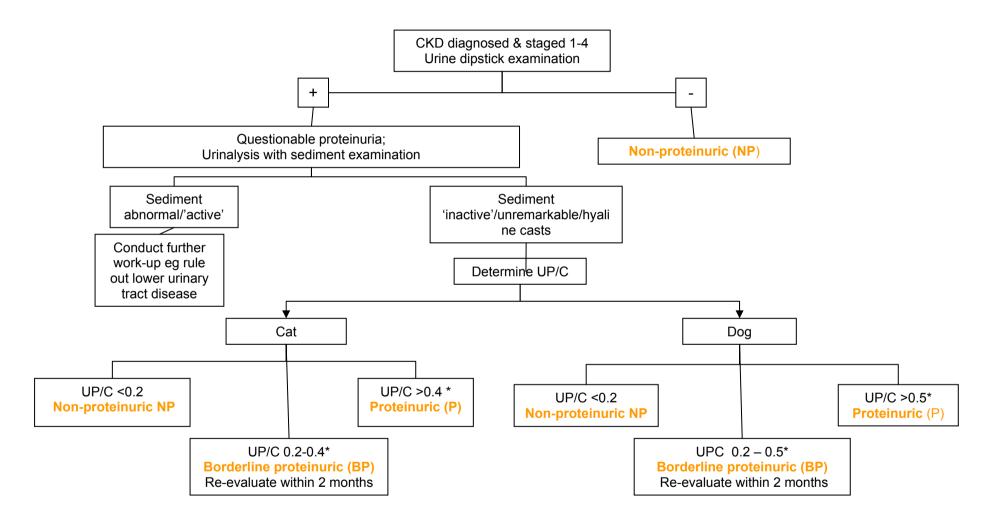
Algorithm for Staging of Chronic Kidney Disease in Dogs



Algorithm for Staging of Chronic Kidney Disease in Cats



Algorithm for Substaging by Proteinuria



*demonstrate persistence by re-evaluating:

if Borderline Proteinuric in 2 weeks to 2 months

if Proteinuric in 2-4 weeks

if UPC>2 no need to demonstrate persistence prior to initiating therapy (severe proteinuria)

Algorithm for Substaging by Blood Pressure (risk of end organ damage from hypertension)

