# Electrolyte, blood gas and acid-base imbalance in canine chronic renal disease in dogs

Katyayini Sharma \*a, Ajay Katoch a, Ankur Sharma a, Devina Sharma b and Adarsh Kumar c

- <sup>a</sup> Department of Veterinary Medicine, DGCN College of Veterinary and Animal Sciences, Palampur, Himachal Pradesh, India.
- b Department of Veterinary Parasitology, DGCN College of Veterinary and Animal Sciences, Palampur, Himachal Pradesh, India.
- c Department of Veterinary Surgery and Radiology, DGCN College of Veterinary and Animal Sciences, Palampur, Himachal Pradesh, India.

#### **Abstract**

Chronic kidney disease (CKD) in dogs is a progressive and often irreversible condition where the kidneys gradually lose their ability to filter waste, regulate fluid balance and maintain essential electrolyte levels. Acid-base disturbances are commonly identified in critically ill veterinary patients. So, the present investigation was aimed to study blood gas, acid base and electrolyte alteration in canine chronic kidney disease. Study was conducted on 20 dogs suffering from stage IV chronic kidney disease out of which 10 cases were found to be positive for haemoprotozoan infection through microscopic examination or rapid diagnostic kits. Haematologically mean values of TEC, PCV and Platelets were significantly reduced. Biochemically the mean values of Creatinine, SGPT and SGOT were significantly increased. Mean values of HCO3, pCO2, BE and BB were significantly decreased as compared to healthy dogs on venous blood gas and acid-base analysis. Electrolyte and mineral estimation revealed significant decrease in the values of sodium, chloride, calcium and magnesium whereas phosphorus were increased. The assessment of blood gas, acid-base balance, electrolyte levels and timely presentation of clinical cases played a crucial role in managing canine chronic kidney disease.

Keywords: Chronic kidney disease, Acid-base, Electrolytes

### Introduction

Electrolyte and acid-base balance are essential components of physiological homeostasis in dogs, influencing vital processes such as neuromuscular function, fluid regulation, and cellular metabolism. Acid-base disturbances in dogs can occur due to a variety of conditions, including renal insufficiency, diabetic ketoacidosis (DKA), prolonged vomiting or diarrhoea, shock, and respiratory diseases that impair gas exchange (Tvedten *et al.*, 2016). The present study describes on electrolyte, blood gas and acid-base imbalances in dogs with chronic kidney disease.

#### **Materials and Methods**

The present study was carried out on dogs presented during the period of February, 2024 to August, 2025 in the Department of Veterinary Medicine, Veterinary Clinical Complex, College of Veterinary

and Animal Sciences, CSKHPKV Palampur (H.P). Preliminary screening was done on the basis of patient's history and presenting clinical signs as depression, inappetence, dehydration, pyrexia, enlarged lymph nodes, vomiting, diarrhoea, polydipsia, polyuria, halitosis, dental tartar and seizures. In addition to haemato-biochemical estimation, electrolyte, mineral analysis, blood gas and acid-base analysis were performed. Blood smear examination (Fig. 1) and serological detection kits (Fig. 2) were used for the detection of haemoprotozoan infection. Dogs in stage IV of chronic kidney disease as per IRIS staging were selected. On this basis, a total of 20 dogs suffering from stage IV CKD were included in the present study. 10 apparently healthy dogs independent of sex and breed presented for routine health check-ups and regular vaccination were considered as healthy control group.

Corresponding author: katyayinisharma@gmail.com

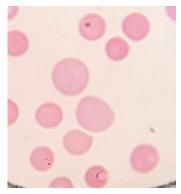


Fig. 1: Babesia gibsoni observed in a Giemsa-stained peripheral blood smear



Fig. 2: Qualitative immunochromatographic assay kit

## **Results and Discussion**

Out of 20 dogs in stage IV of chronic kidney disease, 10 were found to be positive for haemoprotozoan infection representing 50% of the cases. Most common clinical signs observed were depression, dehydration, inappetence, vomiting and pale mucous membrane. These findings were in accordance with Nakang *et al.* (2019) and Kumar *et al.* (2020), The mean values of rectal temperature, heart rate and respiration rate varied non significantly as compared to healthy animals. The mean values of haematological parameters of dog with stage IV CKD are presented above in Table 1.

Table 1: Mean values of haematological parameters in dogs with stage IV chronic kidney disease

Parameters	Healthy Control (n=10)	CKD (n=20)
Hb (g/dl)	13.47±0.34	10.93±0.92
PCV (%)	38.87±1.04	28.5±2.3**
TEC (×10 <sup>12</sup> /L)	6.49±0.21	4.45±0.38 **
TLC (10 <sup>9</sup> /L)	11.13±0.6	19.71±4.13
N (%)	79.68±0.85	80.747±2.82
L (%)	15.3±0.62	10.221±2.127
M (%)	4.04±0.27	7.463±0.818**
E (%)	1.35±0.18	1.205±0.2686
MCV (fl)	61.57±0.47	64.68±1.42
MCH (pg)	21.88±0.4	24.69±0.35***
MCHC (g/dl)	35.18±0.58	252.41±41.5**
Platelets (10 <sup>9</sup> /L)	299.2±23.72	121.88±15.58***

<sup>\*</sup> Significant at 5% (P<0.05); \*\* Significant at 1% (P<0.01); \*\*\*Significant at 0.1% (P<0.001)

The mean values of PCV, TEC and platelets were significantly decreased whereas mean values of MCH and MCHC were significantly increased. These findings are in accordance with Sharma *et al.* (2015), Devpriya *et al.* (2018) and Eashwar *et al.* (2024). Anaemia in canine chronic kidney disease is multifactorial and associated with decreased erythroid precursor cells,

gastrointestinal bleeding, and systemic inflammation as noted by Crivellenti *et al.* (2023). Decreased platelet count was also reported by Sharma *et al.* (2015) which can be attributed to the reduced thrombopoietic activity in uremic dogs. The mean values of biochemical parameters in dogs with stage IV CKD are presented above in Table 2

Parameters	Healthy (n=10)	CKD (n=20)
Glucose (mg/dl)	102.67±2.66	126.152±10.498
Bilirubin (mg/dl)	0.22±0.04	1.62±1.215
AST (U/L)	38.05±3.98	86.06±15.04*
ALT (U/L)	33.92±2.4	88.35±11.402**
ALP(U/L)	79.25±8.72	179.43±51.606
TP (g/dl)	6.6±0.35	6.632±1.167
Creatinine (mg/dl)	1.01±0.08	15.08±1.48***
BUN (mg/dl)	25.14±4.54	515.31±174.18

Table 2: Mean values of biochemical parameters in dogs with Stage IV Chronic Kidney Disease

The mean values of AST, ALT and creatinine were significantly increased as compared to healthy animals. These findings were in accordance with Bradea *et al* (2013), Puri *et al*. (2015), and Devpriya *et al*. (2018). Reason for increased creatinine was its diminished renal excretion in CRF and at least 75% loss of functional nephrons as mentioned by Lefebvre

(2011). Elevations in liver enzymes like ALT and AST are not exclusive to primary liver diseases. They can also occur in extrahepatic conditions including circulatory disturbances and systemic illnesses, which are common in advanced CKD as stated by Alvarez *et al.* (2009). The mean values of electrolyte estimation in dogs with stage IV CKD are presented above in Table 3.

Parameters	Healthy (n=10)	CKD (n=20)
Sodium (mmol/L)	151.8±1.71	141.3±2.8*
Potassium (mmol/L)	4.7±0.16	5.09±0.4
Chloride (mmol/L)	110.05±1.85	102.96±1.89*
Calcium (mg/dl)	10.56±0.27	9.737±0.219*
Phosphorus (mg/dl)	5.07±0.12	6.774±0.478*
Magnesium (mg/dl)	1.78±0.07	1.39±0.084**

<sup>\*</sup> Significant at 5% (P<0.05); Significant at 1% (P<0.01); \*\*\*Significant at 0.1% (P<0.001)

The mean values of electrolytes in dogs with stage IV CKD showed a significant decrease in the values of sodium, chloride, calcium and magnesium whereas the mean values of phosphorus was increased as compared to healthy dogs. These findings are in accordance with Kandula and Karalpudi (2015), Puri *et al.* (2015) and Sumit *et al.* (2018). Suggestive reason for hyperphosphatemia was declining kidney function because the kidneys are the primary route of phosphorus excretion and reduced kidney function results in phosphorus retention (Sonu *et al.*, 2019). Sakaguchi (2022) stated that tubular dysfunction or interstitial fibrosis may play a pivotal role in the development

of hypomagnesemia in CKD, which impairs tubular magnesium reabsorption. Hyponatremia observed could be due to hypotension, pain, and renal injury which activates sympathetic nervous system, renin angiotensinal dosterone system and antidiuretic hormone release (Prowle *et al.*, 2010) leading to increased absorption of sodium and water. Hypocalcaemia may be due to increase in GFR leading to abrupt increase in serum phosphorus concentration, causing a decrease calcium concentration due to chelation of phosphorus. This was in agreement with Schenck et al. (2005). The mean values of blood gas and acid-base analysis in dogs with stage IV CKD are presented above in Table 4.

<sup>\*</sup> Significant at 5% (P<0.05); \*\* Significant at 1% (P<0.01); \*\*\* Significant at 0.1% (P<0.001)

Parameters	Healthy (n=6)	CKD (n=20)
pH (mmHg)	$7.35 \pm 0.02$	7.2705±0.025
pCO2 (mmol/L)	38.15± 2.02	27±0.202***
HCO3 (mmol/L)	20.16± 0.52	11.484±0.771***
AnGap (mmol/L)	30.26± 1.25	30.266±2.278
tCO2 (mmol/L)	20.8± 0.65	12.273±0.785***
BE (mmol/L)	-6.08± 0.84	-12.773±1.189**
BEact (mmol/L)	-6.26± 1.12	-14.173±1.201**
BEecf (mmol/L)	-5.98± 1.34	-13.805±1.037***
BB (mmol/L)	42.92± 0.46	33.184±1.309***
stHCO3 (mmol/L)	20.16± 0.42	13.926±0.8522***
st pH	$7.34 \pm 0.01$	7.1507±0.027**
cH+ (nmol/L)	49.63± 0.81	55.015±3.364

Table 4: Mean values of blood gas and acid-base analysis in dogs with stage IV chronic kidney disease

The mean values of HCO<sub>3</sub> pCO<sub>5</sub> BE and BB were significantly decreased as compared to healthy dogs. These findings were in accordance with Koenhemsi and Gonul (2019) and Lippi et al. (2023). The decreased HCO<sub>3</sub> concentration in CKD patients possibly was due to reduced reabsorption of filtered bicarbonate as number of functioning renal tubules become limited as CKD progresses (Buduk et al., 2020). In CKD a common finding is metabolic acidosis which stimulates compensatory hyperventilation leading to a decrease in arterial pCO2. BE reflects the amount of excess or insufficient level of bicarbonate in the system and BB refers to the total concentration of buffering anions in the blood, mainly HCO<sub>3</sub> along with proteins, phosphates, and haemoglobin. Altered blood gas and electrolyte changes were also associated with the severity of the disease and loss of kidney function. The assessment of blood gas, acid-base balance, electrolyte levels and timely presentation of clinical cases played a crucial role in managing canine chronic kidney disease.

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