Neutrophil gelatinase-associated lipocalin in dogs with sepsis undergoing emergency laparotomy: a prospective case-control study.

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Running head: NGAL in dogs with sepsis

Keywords: NGAL; AKI, acute kidney injury; canine

Abbreviations:
ACE: Angiotensin-converting enzyme
AKI: acute kidney injury
AKIN: Acute kidney injury network
APPLE: The acute patient physiologic and laboratory evaluation score

MODS: Multiple organ dysfunction syndrome

NGAL: Neutrophil gelatinase-associated lipocalin

NSAID: Non steroidal anti-inflammatory drug

RIFLE: Risk, injury, failure, loss of kidney function and end-stage kidney disease.

SIRS: Systemic inflammatory response syndrome

UNCR: urinary NGAL normalized to creatinine

VAKI: Veterinary acute kidney injury

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Part of this study was presented as an abstract at the European Congress of Veterinary Emergency and Critical Care in Prague, 2014.

Abstract

Background
Neutrophil gelatinase-associated lipocalin (NGAL) is an early indicator of acute kidney injury (AKI) in dogs and its use has not been evaluated in dogs with sepsis.

**Animals:**
Fifteen dogs with sepsis requiring laparotomy (study dogs) and 10 dogs undergoing surgery for intervertebral disc disease (control dogs).

**Objective:**
To determine whether NGAL increases in dogs with sepsis—undergoing emergency laparotomy and whether it is correlated with development of AKI and survival.

**Methods:**
Longitudinal study conducted at a referral teaching hospital. Serum (sNGAL), urinary NGAL normalized to creatinine (UNCR) and serum creatinine concentration were measured at 4 time points (admission, after anesthesia, and 24 and 48 hours post-surgery). Development of AKI (increase in serum creatinine concentration of 0.3 mg/dL) and in-hospital mortality were recorded.

Linear mixed-model analysis was employed to assess differences between groups over time. Mann-Whitney U test was performed for comparison of continuous variables.
between groups and Chi square or Fisher’s exact tests were used to assess correlation between discrete data.

**Results**

Serum NGAL and UNCR were significantly higher in study dogs across all time points (p = 0.007 and p < 0.001 respectively) compared with controls. UNCR in the study group was not significantly different between survivors (n=12) and non-survivors (n=3). Dogs that received hydroxyethyl starch had significantly higher UNCR across all time points (p = 0.04) than those that did not.

**Discussion – Conclusion**

sNGAL and UNCR are increased in dogs with sepsis requiring emergency laparotomy. Additional studies are needed to evaluate its role as a marker of AKI in this population.

**Introduction**

Acute kidney injury is defined as sudden injury to the renal parenchyma and reduction in renal function, regardless of underlying cause\(^1\). In critical illness and in sepsis, AKI has been identified as one of the main contributors to morbidity and mortality in human patients.\(^2,3\)
The incidence of AKI in hospitalized septic dogs managed in an intensive care setting has been reported as 12% and the in-hospital mortality rate for dogs with AKI varies from 54% to 86%.\textsuperscript{4,5}

No consensus has yet been reached in veterinary medicine however with regard to criteria that should be used to define kidney injury. There are several emerging classification systems available, including Veterinary Acute Kidney Injury staging system (VAKI) and Risk, Injury, Failure, Loss of kidney function and End-stage kidney disease (RIFLE)-like criteria.\textsuperscript{4,6}

The main pitfalls of the current systems are the use of insensitive and poorly specific markers, such as serum creatinine concentration or urine output. Serum creatinine concentration is not linearly related to GFR and can be influenced by factors other than renal function such as muscular mass and hydration state.\textsuperscript{7} Glomerular filtration rate (GFR) is the gold standard for diagnosis of renal disease but this method, apart from not being readily clinically available, only detects reduction in renal function and not cellular injury, thus, leading to delayed recognition of disease.\textsuperscript{1,8}

Since the 1990’s, there has been a perceived need to identify early markers of AKI in order to identify at-risk patients. To this end, many biomarkers, such as N-acetyl-\(\beta\)-d-
glucosaminidase (NAG), kidney injury molecule-1 (KIM-1) and neutrophil gelatinase-associated lipocalin (NGAL) have been studied in human and canine patients. Neutrophil gelatinase-associated lipocalin is a 25 kDa protein, belonging to the lipocalin family. It is covalently bound to gelatinase and secreted by neutrophils and normally is expressed at low concentrations in the kidney, lung, trachea and gastrointestinal tract. It is freely filtered through the glomerulus and re-absorbed in the proximal tubule, such that healthy individuals usually have very low urinary excretion of NGAL. Its production is highly upregulated when epithelial cell damage occurs. In clinical studies in people and dogs, its concentration in urine increases 24 to 72 hours before serum creatinine concentration, which enables earlier detection of AKI. Neutrophil gelatinase-associated lipocalin can be measured in plasma, serum or in urine and it usually is expressed as an absolute concentration. In some studies, urinary NGAL has been normalized to urinary creatinine concentration in an attempt to decrease variation associated with changes in urine volume. Plasma and urine NGAL is increased in people with sepsis and studies have shown an inconsistent ability for plasma or serum
NGAL to detect AKI in humans with severe sepsis or septic shock.\textsuperscript{16-17}

The ability of urinary NGAL to predict AKI recently has been evaluated in veterinary medicine. A number of clinical studies, published in the last few years, have shown a significantly higher urinary NGAL concentration in dogs with AKI compared to healthy dogs.\textsuperscript{12-14}

The aim of the present study was to evaluate the usefulness of both serum NGAL and urine NGAL normalized to urinary creatinine (UNCR) for detecting AKI in dogs with sepsis undergoing emergency laparotomy compared to dogs with intervertebral disc disease undergoing surgery. It was hypothesized that an increase in urinary NGAL concentration would be associated with development of AKI, longer time to discharge, increased morbidity (assessed by the Acute Patient Physiologic and Laboratory evaluation score [APPLE] full scoring system and Multiple organ dysfunction syndrome [MODS] criteria) and higher mortality.

**Material and Methods**

The study was a prospective investigation performed on client-owned dogs. The protocol was approved by the Ethics and Welfare Committee of the Royal Veterinary College, London.
(URN 2012 1155) and informed owner consent was obtained before enrolling each dog in the study.

The study group included dogs with sepsis that were to undergo emergency laparotomy. Dogs initially were recruited if they fulfilled Systemic inflammatory response syndrome (SIRS) criteria (Table 1).\textsuperscript{18} Dogs were included in the analysis if they fulfilled criteria for sepsis (i.e. presence of SIRS and confirmed or suspected infection).\textsuperscript{19} Infection was defined as the presence of intracellular bacteria on cytological examination or microbiological evidence of infection in the abdominal fluid or aspirates from an abscess. Dogs with azotemia (defined as plasma or serum creatinine concentration > 1.57 mg/dl [138 µmol/l]) on presentation, or a history of chronic kidney disease, were excluded from the study.

A control group of systemically healthy dogs undergoing emergency surgical intervention for intervertebral disk disease (IVDD) also was recruited.

\textit{Admission data}

Clinical data was collected on admission and included signalment, disease process, heart rate, respiratory rate, rectal temperature, systolic blood pressure, whole blood lactate concentration and serum creatinine concentration.
Hospitalization data

Before or during the course of hospitalization, administration of vasopressors, hydroxyethyl starches, blood products, human serum albumin and potentially nephrotoxic drugs (e.g. non-steroidal anti-inflammatory drugs [NSAID], angiotensin-converting enzyme [ACE]-inhibitors) were recorded.

Occurrence of AKI, azotemia, hypotension and survival were recorded as categorical data. Acute kidney injury was defined as an increase in serum creatinine concentration of 0.3 mg/dl (26.4 µmol/l) within 48 hours, in keeping with the Acute kidney injury network (AKIN) criteria. Serum creatinine concentration was measured on admission, post-anesthesia and 24 and 48 hours post-surgery. Hypotension was defined as a systolic Doppler pressure < 90 mmHg or a mean invasive arterial pressure < 60 mmHg. The APPLE full score on admission was calculated for each dog when data were available. Multi-organ dysfunction syndrome was defined as ≥2 organ systems having signs of dysfunctions.

Sampling and sample processing

Blood and urine samples were taken from dogs in both groups for NGAL and creatinine measurements. Samples were taken at
4 time points: (I) on admission, (II) after anesthesia (within 2 hours post-anesthesia), and (III) 24 and (IV) 48 hours post-operatively. Because ours was a clinical study performed in the United Kingdom on non-experimental animals, only residual volumes of blood taken from samples for clinical need could be used.

Urine samples were collected by voiding or from an indwelling urinary catheter, and an aliquot of each fresh urine sample was submitted to the laboratory for urinalysis (refractometric urine specific gravity, dipstick semi-quantitative chemistry and sediment examination) and urine-protein-to-creatinine ratio (UP:C) measurement. Pyuria was defined as $\geq 5$ white blood cells per high power field. Aliquots of the urine and serum samples were centrifuged at 400 $g$ for 5 minutes and the supernatant frozen at -70°C within 1 hour (urine) and 8 hours (serum) for later batch analysis.

Serum creatinine concentrations were determined from stored frozen serum samples using a biochemical analyzer. Serum NGAL was measured on the same day using the same thawed samples. All samples thawed for NGAL analysis were re-frozen within 1 hour to allow for re-analysis if required.

Urine and serum NGAL concentrations were determined using a sandwich ELISA following the manufacturer's instructions.
Urine and serum samples initially were diluted 1:200 using the diluent provided in the kit and the assay was performed according to the manufacturer’s instructions. When necessary, dilution of the samples was adjusted (1:20 to 1:10000) to ensure that the NGAL concentration fell on the linear part of the standard curve. Samples requiring further dilution were thawed and frozen a maximum of 3 times. This thaw-refreeze process has been shown not to affect NGAL stability.\textsuperscript{15} Urinary NGAL concentrations were normalized by the urinary creatinine concentration (mg/dl) to express results as UNCR. The final normalized uNGAL is expressed as ng/mg.

A dual set of pooled quality control samples (high and low concentrations) also were included within each plate for assessment of inter-assay and intra-assay variability.

\textit{Statistics}

Statistical analyses were performed in SPSS\textsuperscript{e} and figures were drawn using Graphpad Prism.\textsuperscript{f} Continuous data was presented as median (range) in the original scale because some of the data have been log transformed in the statistical analysis and the sample size was small.

Repeated measurement of UNCR and sNGAL were analyzed in a linear mixed-effects model (LMM) taking time, group (sepsis vs
control dogs), colloids, AKI, MODS, hypotension and blood product requirement as potential fixed effects and dog as random effect. Potential 2-way interactions between categorical fixed factors and time were assessed. A continuous variable was log transformed if its residuals from LMM analysis were skewed to the right.

Although many continuous data were normally distributed, admission data between clinical course and treatment groups and survivors were compared using the Mann Whitney-U test because of the small sample size. Similarly, Spearman’s correlation coefficient was used to measure linear correlation between 2 sets of continuous data to avoid the potential influence of outliers. The association between categorical variables was analyzed using Chi-squared or Fisher’s exact test if the contingency table contained a value <5. Statistical significance was set at p < 0.05.

Results

Recruitment and admission characteristics

Twenty-one dogs were enrolled in the study group and 10 dogs were enrolled in the control group between June 2012 and August 2013. One dog in the study group subsequently was excluded because no urine had been stored for pre-surgery
NGAL analysis. Twenty dogs initially were included in the study. Five of these were removed because they did not fulfill criteria for sepsis. Four of these dogs had gastric dilatation and volvulus and 1 had severe gastroenteritis. The final study group therefore was represented by 15 dogs. Of these dogs, 13 had septic peritonitis and 2 had intra-abdominal abscessation. Admission variables and days of hospitalization were compared between the study and the control groups and reported in Tables 2 and 3.

*Clinical course and treatment*

All dogs, in both the study and control groups, had blood pressure recorded over the course of the entire anesthetic period, and the study population had periodic blood pressure measurements made during hospitalization (Tables 2 and 3). Eleven of 15 dogs in the study group were hypotensive (4 pre-operatively and 10 intra-operatively). Two of 3 non-survivors were hypotensive in the immediate post-operative period and required vasopressor therapy. Blood pressure was not measured on admission for any of the dogs in the control group because they were considered to be cardiovascularly stable based on clinical examination.
None of the dogs in the control group developed AKI whereas 2/15 (13%) dogs in the study group met AKI criteria but did not become azotemic (serum creatinine concentration increased by 0.57 mg/dl [50 µmol/l] and 0.65 mg/dl [57 µmol/l] in 24 hours). These 2 dogs later were euthanized because of clinical deterioration. No difference was identified between the incidence of AKI in the study and control groups (p= 0.5; Table 3).

In the study population, 12 dogs (80%) survived to discharge, 1 dog died and 2 were euthanized after clinical deterioration. Nine dogs (90%) in the control group were discharged, with 1 being euthanized because of persistent paraplegia (Table 3).

**Serum NGAL results**

Intra-assay and inter-assay variability for sNGAL was 2.0% and 10.4% for the high concentration quality control and 7.7 and 19.3% for the low concentration quality control, respectively. No significant difference was identified for time-group interaction and among times, but sNGAL was significantly higher in the study population compared to the control population (p= 0.007; Figure 1 and Table 4). A time-colloid administration interaction was present (p=0.01). The cause of significance seemed to derive mainly from difference between
time points within the non-colloid group, and there was no
difference between colloid group at all time points (Table 4).
There was no association between AKI, MODS development,
hypotension and survival and sNGAL (Table 4).

**UNCR results**

There was no time-group interaction and no difference among
times for UNCR, but the study group had significantly higher
UNCR than did the control dogs at all measured time points
(p<0.001). No association was detected between UNCR and
development of AKI (p = 0.25) or MODS (p= 0.49; Figure 2 and
Table 4). Four of the 30 dogs (2 in each group) had pyuria.
There was no association between UNCR and the presence of
pyuria at any of the time points (p=0.55). None of the dogs in
the control group was diagnosed with a urinary tract infection
(UTI). One dog in the study group was diagnosed with a UTI,
and this dog also had prostatic abscessation and septic
peritonitis.

No time-colloids administration interaction was identified and
no difference among time points was detected. Dogs that
received hydroxyethyl starch (12/30) however had
significantly higher UNCR compared to dogs that did not across
all time points (p = 0.04; Figure 3 and Table 4), with the 2 dogs that developed AKI receiving colloids, either preoperatively or intra-operatively.

All dogs in the control group received NSAIDs after anesthesia, whereas none of the dogs in the study group received NSAIDs after admission. No time-NSAID administration interaction and no difference among times was detected, but dogs in the study group that did not receive NSAIDs before admission (n=11) had significantly higher UNCR (p = 0.04; Table 4).

There was a significant association between UNCR and days spent in the intensive care unit (ICU; Table 5) but no correlation was found between UNCR and duration of hospitalization (admission, p = 0.5; post-anesthesia, p = 0.3; 24 hours, p = 0.34; 48 hours, p = 0.09).

No significant difference in UNCR was detected between dogs that survived to discharge and those that did not (p = 0.08).

Correlation

There was moderate correlation between UNCR and sNGAL on admission (r_s 0.76) and post-anesthesia (r_s 0.82; Table 4). No correlation was found between UNCR and urinary pH and urine specific gravity (USG).
There was a correlation between UNCR and UP:C (admission, \( r_s \) = 0.61; post-anesthesia, \( r_s \) = 0.92; 24 hours, \( r_s \) = 0.87; 48 hours, \( r_s \) = 0.75) but no correlation was found between UNCR and APPLE full score at any of the time points (Table 4).

**Discussion**

Our study focused on detecting the effect of sepsis on sNGAL and UNCR and to assess their changes over the course of hospitalization. The control population was chosen to compare the effect of general anesthesia and surgery on NGAL in comparison to septic dogs undergoing emergency laparotomy. This group also was chosen because it theoretically allowed timed and non-invasive urine sampling because of routine placement of urinary catheters or bladder expression.

Sepsis has been shown to be associated with development of AKI secondary to ischemia-reperfusion injury, damage induced by pro-inflammatory cytokines and oxidative stress and tubular dysfunction leading to decreased glomerular filtration and tubular reabsorption dysregulation. Neutrophil gelatinase associated lipocalin is secreted by neutrophils and it is therefore logical to expect an increase in its concentration in inflammatory states. In particular, during sepsis, NGAL
expression increases, not only in the kidney, but also in leukocytes and liver. Therefore, higher NGAL both in blood and in urine are expected, regardless of AKI. A study in human patients showed that sNGAL was increased in patients with sepsis, severe sepsis and septic shock, but was not able to detect AKI in this population, especially in the sicker categories. The results of the present study confirm that sNGAL also in increased in dogs with sepsis compared to a control population. However, sNGAL was not associated with mortality or correlated with the severity of the disease, as assessed by the APPLE scoring system or MODS development, suggesting that this marker may not be a useful clinical severity predictor.

The UNCR differed between the study and control groups across all time points evaluated (p < 0.001). The median UNCR concentration in the study group post-anesthesia was as high as 114 times (admission) and 185 times higher than in the control group, highlighting the marked difference between the 2 groups. However, similar to sNGAL, UNCR was not associated with mortality, suggesting that this marker may not be useful to assess the severity of disease.

A study in people showed that septic AKI patients had higher urinary NGAL than non-septic AKI patients. Both sNGAL and
UNCR were not associated with AKI development over the course of hospitalization in this study. This finding may be a consequence of the inability of NGAL to distinguish between the presence of AKI and systemic inflammation in septic dogs, and as a result may suggest that NGAL is a poor marker of AKI in this population of dogs. The lack of significance to detect AKI, however, could have been a result of the small population size and thus indicate a type II error.

Neutrophil gelatinase associated lipocalin increases 3 hours after ischemic renal injury in people undergoing cardio-pulmonary bypass and peaks at 36 hours in critically ill patients developing AKI during hospitalization.\textsuperscript{24} However NGAL has been shown to increase up to 72 hours earlier than serum creatinine concentration in the event of AKI. Therefore, this study may not have detected an increase in serum creatinine concentration if it occurred after the 48 hours sample, potentially leading to AKI recognition being missed in some patients.\textsuperscript{10,23} This time window was selected based on evidence in human patients showing that adverse outcome is associated with increased serum creatinine concentration within 48 hours, and the immediate post-operative period was deemed most appropriate to detect eventual development of AKI.\textsuperscript{13,25}
The increase in UNCR in dogs with sepsis in our study perhaps represents sub-clinical AKI, not detected by the VAKI system, or may have been related to an increase in serum creatinine concentrations that occurred after 48 hours. The increase in serum creatinine concentration used to define AKI in our study was considered to be a sensitive way of identifying these dogs. However, serum creatinine concentration may not be altered in the event of tubular injury, indicated by an increase in NGAL production, because the GFR of the affected nephrons may not be altered or overall renal compensation may occur in other nephrons. Conversely, tubular injury, detected by an increase in NGAL concentration, may lead to a decrease in glomerular function as a consequence of tubular flow obstruction and tubulo-glomerular interaction. In this situation, NGAL may not be correlated with GFR, and its increase does not consistently reflect a change in serum creatinine concentration.

It is unlikely that the difference in UNCR between the 2 groups was a consequence of altered passage of NGAL across the glomerular filtration barrier. In fact, although there was an association between UP:C and UNCR, sNGAL and UNCR were only moderately correlated on admission and post-anesthesia. It appears more likely that in severe inflammation renal derangements lead to decreased re-absorptive capacity in the
proximal tubules or up-regulation of NGAL production at the
level of the thick ascending loop, distal tubule and collecting
ducts.\textsuperscript{10}

One study showed that, in dogs undergoing surgery, urinary
NGAL was significantly higher in dogs with AKI 12 hours post-
operatively, whereas plasma NGAL was not able to detect AKI.\textsuperscript{13}

Another study in dogs with a variety of diseases, including
heatstroke and snake envenomation, showed that UNCR on
admission was significantly higher in dogs with AKI (grade I to
V) and detected AKI earlier than the increase in plasma
creatinine concentration.\textsuperscript{12} However, 9 of 15 dogs (60\%) in our
study had UNCR results on admission higher than the cut-off of
238 ng/mg established by a previous study for distinguishing
non-azotemic AKI grade I patients from dogs with related renal
or urinary conditions.\textsuperscript{12} However, the pathophysiology of AKI in
sepsis is substantially different from other causes of AKI, such
as ischemia or toxic damage, and NGAL expression is
upregulated in inflammatory conditions, such as infection.\textsuperscript{21,22}

Of the 9 dogs with UNCR >238 ng/mg, only 2 developed AKI
and none of them developed azotemia. This cut-off may
therefore not be useful in dogs with sepsis and additional
studies are needed to establish a cut-off for this population.

Both UNCR and sNGAL were not significantly associated with
time to discharge but were associated with ICU stay. Timing of discharge and ICU hospitalization are clinician-dependent and financially-based decisions and these factors might have influenced this result. In addition, our control population was represented by patients with IVDD, which have predictably long hospitalization times.

Patients that did not receive NSAIDs before admission had higher UNCR compared to patients that did across all time points. This difference may have been simply a consequence of more severe disease in dogs not receiving NSAIDs, because there currently is no evidence that NSAIDs may induce urinary NGAL down-regulation.

An interesting finding was the significant difference in UNCR between dogs that received colloids and dogs that did not. The higher UNCR in dogs that received colloids was consistent across all 4 time points, and UNCR was increased before surgery although only 4 of the 9 dogs were given colloids on admission, whereas 4 received them intra-operatively and 1 post-operatively. This difference may simply reflect a difference in severity of illness, because colloids typically are used more often in dogs with more serious signs of hypoperfusion and potentially more severe inflammation. This
may have induced an up-regulation of NGAL secretion and an increase in UNCR independent of colloid administration.\(^\text{11}\) In people, hydroxyethyl starches are associated with increased risk of kidney injury, and recent studies have highlighted an association between the use of hydroxyethyl starches (including pentastarch and tetrastarch) and kidney injury or renal replacement therapy requirement.\(^\text{27,28}\) Usually, an increase in uNGAL is expected to occur at least 3 hours after the renal injury.\(^\text{11}\) If these artificial colloids induced AKI in the dogs of our study, a difference at the pre-anesthesia sample would not be expected, because most patients did not receive hydroxyethyl starches until intra-operatively or post-operatively. The results of our study should therefore be interpreted with caution.

One limitation of this study is that there were 21/100 urine samples and 40/100 serum samples missing. Analysis of all of the samples may have added further information to our analysis and allowed better interpretation of the data. The absence of results, in addition to the small number of cases enrolled, therefore may have caused the analysis to be underpowered.

A second limitation involves the choice of the control group. The median admission data for this group was not significantly
different from that of the study group, leading to a potential confounding overlap within the population (Tables 1 and 2).

However, the underlying diseases affecting the 2 groups were substantially different, being localized in the control dogs and systemic and pro-inflammatory in the study patients. Additional studies may benefit from the use of more stringent SIRS criteria, allowing clear clinical distinction between the control and the study group on admission.

In addition, our study monitored serum creatinine concentration for a 48-hours period. Because NGAL may increase up to 72 hours earlier than serum creatinine concentration in the event of renal injury, some patients with AKI may have been missed.11,24.

A further limitation includes the clinician-based decision for administration of hydroxyethyl starches, blood products or institution of vasopressor therapy, rendering objective interpretation of these data or assessment of the severity of illness difficult. It also is currently unknown whether drugs, such as human serum albumin, may affect serum and urinary NGAL release and tubular re-absorption, interfering with the results of the study.

Additional prospective studies with larger populations of dogs with sepsis and use of additional urinary markers are
warranted to assess the association between NGAL and inflammation, sepsis and AKI.

Footnotes:

a. Dog NGAL ELISA kit, BioPorto Diagnostics, Gentofte, Dk.
b. Tecn Sunrise Elisa-reader; Tecan Group, Mannedorf, Ch
c. IL600, Instrumentation Laboratory Cheshire, UK
e. SPSS version 20, Chicago, Ill, USA
f. Graphpad Prism, version 6, La Jolla, Ca, USA
g. Voluven® (6% Hydroxyethyl Starch 130/0.4 in 0.9% Sodium Chloride), Fresenius Kabi Norge A.S., Halden Norway

Bibliography:


Legends:

**Table 1:** Systemic Inflammatory Response Syndrome (SIRS) criteria used for inclusion of dogs within the study group. SIRS was defined if patients fulfilled 2 or more criteria.18

**Table 2:** Measurements taken at admission and hospitalization time between study and control patients. Measurements are expressed as median (range) and statistical significance was set at P<0.05. NA= not applicable; R.I.: Reference Interval.

**Table 3:** Pre-admission treatment, clinical course and treatment received in the study and in the control dogs. Significant differences are highlighted (significance set at a p value ≤ 0.05).

APPLE score was calculated.20

MODS (Multi-Organ Dysfunction Syndrome) was defined as dysfunction of at least two organs.5

AKI (Acute Kidney Injury) was defined as an increase of 0.3 mg/dl (26.5 µmol/l) from baseline.4
Table 4: Differences in the normalized sNGAL and UNCR for each single category and differences between time and time-category interaction. Significant difference was set at p < 0.05. AKI: acute kidney injury; NSAID: non-steroidal anti-inflammatory drug; MODS: Multi-organ dysfunction syndrome.

Table 5: Spearman's correlation ($r_s$) between UNCR and UP:C, serum creatinine, sNGAL, day in I.C.U and APPLE score. Statistical significance was set at p < 0.05.

Figure 1: The medians and interquartile range of serum NGAL at different times between the control and the study group are represented by the bar and the whiskers with all the patients displayed in a scattered plot manner. The two AKI patients are represented by x and ◆ on the graph. sNGAL concentrations were significantly higher across all time points in the study group.

Figure 2: The medians and interquartile ranges of urinary NGAL to creatinine ratio (UNCR) at different times between the control and the study group are expressed by the bar and the
whiskers respectively with the single patients displayed in a scatter plot manner. The two patients with AKI have been identified with x and ◆. The study group had significantly higher UNCR across all the time points (p < 0.001).

**Figure 3:** The medians and interquartile ranges of urinary NGAL to creatinine ratio (UNCR) at different times between patients that received colloids and patients that did not are expressed by the bar and the whiskers respectively with the single patients displayed in a scatter plot manner. Statistical significance was set at P<0.05. Dogs that received colloids had significantly higher UNCR concentration across all time points.

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<th>Variable</th>
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<td>Heart rate</td>
<td>&gt;120/min</td>
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<tr>
<td>Respiratory rate</td>
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<td>Body temperature</td>
<td>&lt;38.1°C or &gt;39.2°C</td>
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<td>White blood cell count</td>
<td>&lt;6x10³ or &gt;16x10³ /µl</td>
</tr>
<tr>
<td>Increased band neutrophils with normal neutrophil count</td>
<td>3%</td>
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**Table 1:** Systemic Inflammatory Response Syndrome (SIRS) criteria used for inclusion of dogs within the study group; SIRS was defined if patients fulfilled 2 or more criteria.¹⁸
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<td>120 (80-172)</td>
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<td>Respiratory rate (rpm)</td>
<td>36 (20-100)</td>
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<td>Temperature (°C)</td>
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<td>38.8 (36.9-39.4)</td>
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<td>White Blood cells (10⁹)</td>
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<td>-</td>
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<td>Lactate (mg/dl) (R.I. 0-22.5)</td>
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<td>16.2 (16.2-18) (n=3)</td>
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<td>Doppler systolic blood pressure (mmHg)</td>
<td>130 (60-170)</td>
<td>-</td>
<td>-</td>
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</table>

**Table 2**: Measurements taken at admission between study and control patients. Measurements are expressed as median (range) and statistical significance was set at P<0.05. NA= not applicable; R.I.: Reference Interval.
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<td>Previous surgery (5 days)</td>
<td>5/15 (33%)</td>
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<td>13/15 (86%)</td>
<td>1/10 (10%)</td>
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<td>Non steroidal anti-inflammatory medications</td>
<td>4/15 (27%)</td>
<td>6/10 (60%)</td>
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</tr>
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<td><strong>Clinical Course</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Hypotension</td>
<td>11/15 (73%)</td>
<td>0/10 (0%)</td>
<td>&lt; 0.01</td>
</tr>
<tr>
<td>APPLE score</td>
<td>28 (8-38) (14/15)</td>
<td>.</td>
<td>.</td>
</tr>
<tr>
<td>MODS</td>
<td>4/15 (27%)</td>
<td>0/10 (0%)</td>
<td>0.12</td>
</tr>
<tr>
<td>AKI</td>
<td>2/15 (13%)</td>
<td>0/10 (0%)</td>
<td>0.5</td>
</tr>
<tr>
<td>Survival to discharge</td>
<td>12/15 (80%)</td>
<td>9/10 (90%)</td>
<td>0.62</td>
</tr>
<tr>
<td><strong>Treatment</strong></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Blood products</td>
<td>5/15 (33%)</td>
<td>0/10 (0%)</td>
<td>0.06</td>
</tr>
<tr>
<td>Human serum albumin</td>
<td>3/15 (20%)</td>
<td>0/10 (0%)</td>
<td>0.25</td>
</tr>
<tr>
<td>Hydroxyethyl starch (Voluven ®)</td>
<td>9/15 (60%)</td>
<td>0/10 (0%)</td>
<td>&lt;0.01</td>
</tr>
<tr>
<td>Category</td>
<td>Time</td>
<td>Time-category interaction</td>
<td></td>
</tr>
<tr>
<td>------------</td>
<td>--------</td>
<td>---------------------------</td>
<td></td>
</tr>
<tr>
<td>sNGAL</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sepsis</td>
<td>p = 0.007</td>
<td>p = 0.13</td>
<td>p = 0.95</td>
</tr>
<tr>
<td>AKI</td>
<td>p = 0.28</td>
<td>p = 0.24</td>
<td>p = 0.83</td>
</tr>
<tr>
<td>Hypotension</td>
<td>p = 0.49</td>
<td>p = 0.23</td>
<td>p = 0.33</td>
</tr>
<tr>
<td>Survival</td>
<td>p = 0.09</td>
<td>p = 0.25</td>
<td>p = 0.34</td>
</tr>
<tr>
<td>NSAIDs</td>
<td>p = 0.12</td>
<td>p = 0.23</td>
<td>p = 0.47</td>
</tr>
<tr>
<td>Colloids</td>
<td>p = 0.72</td>
<td>p = 0.16</td>
<td>p = 0.01</td>
</tr>
<tr>
<td>MODS</td>
<td>p = 0.49</td>
<td>p = 0.24</td>
<td>p = 0.57</td>
</tr>
<tr>
<td>Blood products</td>
<td>p = 0.96</td>
<td>p = 0.24</td>
<td>p = 0.43</td>
</tr>
<tr>
<td>UNCR</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Sepsis</td>
<td>p &lt; 0.001</td>
<td>p = 0.29</td>
<td>p = 0.12</td>
</tr>
<tr>
<td>AKI</td>
<td>p = 0.25</td>
<td>p = 0.1</td>
<td>p = 0.5</td>
</tr>
<tr>
<td>Hypotension</td>
<td>p = 0.55</td>
<td>p = 0.08</td>
<td>p = 0.75</td>
</tr>
<tr>
<td>Survival</td>
<td>p = 0.08</td>
<td>p = 0.11</td>
<td>p = 0.19</td>
</tr>
<tr>
<td>NSAIDs</td>
<td>p = 0.04</td>
<td>p = 0.08</td>
<td>p = 0.21</td>
</tr>
<tr>
<td>Colloids</td>
<td>p = 0.04</td>
<td>p = 0.12</td>
<td>p = 0.07</td>
</tr>
<tr>
<td>MODS</td>
<td>p = 0.49</td>
<td>p = 0.08</td>
<td>p = 0.44</td>
</tr>
<tr>
<td>Blood products</td>
<td>p = 0.21</td>
<td>p = 0.1</td>
<td>p = 0.07</td>
</tr>
</tbody>
</table>

**Table 3:** Pre-admission treatment, clinical course and treatment received in the study and in the control dogs. Significant differences are highlighted (significance set at a p value ≤ 0.05).

APPLE score was calculated.20

MODS (Multi-Organ Dysfunction Syndrome) was defined as dysfunction of at least two organs.5

AKI (Acute Kidney Injury) was defined as an increase of 0.3 mg/dl (26.5 µmol/l) from baseline.4

**Table 4:** Differences in sNGAL and UNCR for each single category and differences between time and time-category interaction. Significant difference was set at p< 0.05. AKI: acute
kidney injury; NSAID: non steroidal anti-inflammatory drug; MODS: Multi-organ
dysfunction syndrome.
Table 5:

Spearman’s correlation ($r_s$) between UNCR and UP:C, serum creatinine, sNGAL, day in I.C.U and APPLE score. Statistical significance was set at $p < 0.05$. 

<table>
<thead>
<tr>
<th>UNCR admission</th>
<th>UP:C admission</th>
<th>Serum Creatinine</th>
<th>sNGAL admission</th>
<th>DAYS I.C.U.</th>
<th>APPLE score</th>
</tr>
</thead>
<tbody>
<tr>
<td>$p &lt; 0.001; r_s 0.61$</td>
<td>$p = 0.17$</td>
<td>$p &lt; 0.01; r_s 0.76$</td>
<td>$p &lt; 0.001; r_s 0.7$</td>
<td>$p = 0.44$</td>
<td></td>
</tr>
<tr>
<td>UNCR post</td>
<td>UP:C post</td>
<td>Serum Creatinine</td>
<td>sNGAL post</td>
<td>DAYS I.C.U.</td>
<td>APPLE score</td>
</tr>
<tr>
<td>$p &lt; 0.001; r_s -0.92$</td>
<td>$p = 0.03; r_s 0.54$</td>
<td>$p &lt; 0.01; r_s 0.77$</td>
<td>$p = 0.71$</td>
<td></td>
<td></td>
</tr>
<tr>
<td>UNCR 24</td>
<td>UP:C 24</td>
<td>Serum Creatinine</td>
<td>sNGAL 24</td>
<td>DAYS I.C.U.</td>
<td>APPLE score</td>
</tr>
<tr>
<td>$p &lt; 0.001; r_s 0.87$</td>
<td>$p = 0.11$</td>
<td>$p = 0.07$</td>
<td>$p = 0.79$</td>
<td>$p = 0.08$</td>
<td></td>
</tr>
<tr>
<td>UNCR 48</td>
<td>UP:C 48</td>
<td>Serum Creatinine</td>
<td>sNGAL 48</td>
<td>DAYS I.C.U.</td>
<td>APPLE score</td>
</tr>
<tr>
<td>$p = 0.001; r_s 0.75$</td>
<td>$p = 0.15$</td>
<td>$p = 0.33$</td>
<td>$p &lt; 0.001; r_s 0.77$</td>
<td>$p = 0.2$</td>
<td></td>
</tr>
</tbody>
</table>
**Figure 1:** The medians and interquartile ranges of serum NGAL at different times between the control and the study group are represented by the bar and the whiskers with all the patients displayed in a scattered plot manner. The two AKI patients are represented by x and ♦ on the graph. Serum NGAL concentrations were significantly higher across all time points in the study group.
**Figure 2**: The medians and interquartile ranges of urinary NGAL to creatinine ratio (UNCR) at different times between the control and the study group are expressed by the bar and the whiskers respectively with the single patients displayed in a scatter plot manner. The two patients with AKI have been identified with x and ♦. The study group had significantly higher UNCR across all the time points (p < 0.001). Statistical significance was set at p<0.05.
Figure 3: The medians and interquartile ranges of urinary NGAL to creatinine ratio (UNCR) at different times between patients that received colloids and patients that did not are expressed by the bar and the whiskers respectively with the single patients displayed in a scatter plot manner. Statistical significance was set at P<0.05. Dogs that received colloids had significantly higher UNCR concentration across all time points.