Pathology and epidemiology of oxalate nephrosis in cheetahs.

Emily P. Mitchell\textsuperscript{1,2}, Molly E. Church\textsuperscript{3}, Sarah M. Nemser\textsuperscript{4}, Betsy Jean Yakes\textsuperscript{5}, Eric R. Evans\textsuperscript{4}, Renate Reimschuessel\textsuperscript{4}, K. Lemberger\textsuperscript{6}, Peter N. Thompson\textsuperscript{7}, Karen A. Terio\textsuperscript{8}

\textsuperscript{1}Department of Research and Specialised Services, National Zoological Gardens of South Africa, Pretoria, South Africa
\textsuperscript{2}Department of Paraclinical Sciences, Faculty of Veterinary Science, University of Pretoria, Pretoria, South Africa
\textsuperscript{3}Department of Pathology, Microbiology and Immunology, School of Veterinary Medicine, University of California, Davis, California, USA
\textsuperscript{4}United States Food and Drug Administration, Center for Veterinary Medicine, Office of Research, Veterinary Laboratory Investigation and Response Network, Laurel, Maryland, USA
\textsuperscript{5}United States Food and Drug Administration, Center for Food Safety and Applied Nutrition, Office of Regulatory Science, College Park, Maryland, USA
\textsuperscript{6}Vetdiagnostics, Lyon, France
\textsuperscript{7}Department of Production Animal Studies, Faculty of Veterinary Science, University of Pretoria, Pretoria, South Africa
\textsuperscript{8}Zoological Pathology Program, University of Illinois, Brookfield, Illinois, USA

\textbf{Corresponding Author:}

Emily. P. Mitchell (neé Lane), Department of Research and Scientific Services, National Zoological Gardens of South Africa, P O Box 754, Pretoria, South Africa.

Email: emily@nzg.ac.za
Abstract

To investigate cases of acute oxalate nephrosis without evidence of ethylene glycol exposure, archived data and tissues from cheetahs (*Acinonyx jubatus*) from North America (n=297), southern Africa (n=257), and France (n=40) were evaluated. Renal and gastro-intestinal tract lesions were characterized in a subset of animals with (n=100) and without oxalate crystals (n=165) at death. Crystals were confirmed as calcium oxalate by Raman spectroscopy in 45 of 47 cheetahs tested. Crystals were present in cheetahs from 3.7 months to 15.9 years old. Cheetahs younger than 1.5 years were less likely to have oxalates than older cheetahs (p=0.034) but young cheetahs with oxalates had more oxalate crystals than older cheetahs (p<0.001). Cheetahs with oxalate crystals were more likely to have renal amyloidosis, interstitial nephritis or colitis, and less likely to have glomerular loop thickening or gastritis, than those without oxalates. Crystal number was positively associated with renal tubular necrosis (p≤0.001), regeneration (p=0.015), and casts (p≤0.001) but inversely associated with glomerulosclerosis, renal amyloidosis and interstitial nephritis. Crystal number was unrelated to the presence or absence of colitis and was lower in southern African than American and European animals (p=0.01). This study found no evidence that co-existing chronic renal disease (amyloidosis, interstitial nephritis or glomerulosclerosis), veno-occlusive disease, gastritis or enterocolitis contributed significantly to oxalate nephrosis. Oxalate-related renal disease should be considered as a potential cause of acute renal failure especially in young captive cheetahs. The role of location, diet, stress and genetic predisposition in the pathogenesis of oxalate nephrosis in cheetahs warrants further study.
Key words: Acinonyx jubatus, crystals, cheetahs, intestinal disease, nephrosis, calcium oxalate, pathology, renal disease

Since the early 1970’s worldwide sporadic cases of oxalate nephrosis have been documented in captive cheetahs and other large felids. Oxalate nephrosis has also been recorded in other non-domestic mammals including mink and koalas. While small numbers of oxalate crystals are relatively common in cheetahs dying of glomerulosclerosis and/or renal amyloidosis, large numbers of oxalate crystals and associated tubular pathology have been described in cheetah kidneys, without other renal disease being present (KAT, KL, EPM and Linda Munson, personal communication). Approximately 10% of captive cheetahs in the United States and South Africa were estimated to have oxalate crystals in renal tubules at necropsy (KAT and EPL, unpublished data). This warranted further investigation of possible risk factors for renal oxalate formation given the vulnerable conservation status of this species.

Endogenously produced oxalate is a metabolic by-product of amino acid (hydroxyproline, glycine and serine) and vitamin C catabolism. Oxalate nephrosis results from intra-tubular calcium oxalate deposition when calcium oxalate burdens exceed excretory capacity either due to genetic defects in glyoxylate metabolism that result in increased hepatic endogenous production of oxalate, or secondary or acquired disease. Most human forms of primary hyperoxaluria occur due to mutation of the pyridoxine (vitamin B6) dependent enzyme alanine:glyoxylate aminotransferase (AGT) or the enzyme glyoxalate/hydroxypyruvate reductase (GRHPR). Familial tendencies, a point
mutation in the GRHPR gene, and associated reduced enzyme function have also been identified in some domestic cats with oxalate nephrosis.\textsuperscript{6,15,26,49}

Secondary or acquired oxalate nephrosis can be due to severe liver disease\textsuperscript{46} or pyridoxine deficiency,\textsuperscript{5,7,23,65} which result in compromised oxalate metabolism; or increased absorption of oxalates due to ingestion of oxalates (or their precursors) including plants,\textsuperscript{2,3,11,39,58} ethylene glycol and other glycols,\textsuperscript{1,10,11,30,50} xylitol,\textsuperscript{13} ascorbic acid,\textsuperscript{4,47,48,55} and collagen or feathers.\textsuperscript{18,85} An altered intestinal microbiome,\textsuperscript{42,69} excessive bile or long-chain fatty acids\textsuperscript{10,56} or low intestinal calcium content due to diet,\textsuperscript{32} enterocolitis\textsuperscript{5,10} and intestinal surgical resection\textsuperscript{9,31} can also increase oxalate absorption and result in oxalate crystal deposition in the kidney. Ethylene glycol toxicity was considered, although never proven,\textsuperscript{72,76}, in cheetah cases with large numbers of crystals, but was thought unlikely in cases from the United Arab Emirates\textsuperscript{36} and southern Africa where ethylene glycol (antifreeze) is less widely utilized. Toxicological evaluation of feed and water supplies in several cases from the United States, southern Africa and France have failed to find evidence of contaminating ethylene glycol (DGA Meltzer, M Kinsel, and S Terrell, personal communications).\textsuperscript{41} Similarly, ingestion of oxalate-rich plants or sweeteners, pancreatic insufficiency, ascorbic acid toxicity, high fat or protein diets and pyridoxidine deficiency were considered unlikely primary causes in cheetahs given the widely differing environments and management practices in the USA, southern Africa and France.

However, captive cheetahs suffer from renal diseases such as glomerulosclerosis and amyloidosis\textsuperscript{8,51,59} which could result in increased oxalate deposition due to reduced glomerular filtration and urinary pH, pyridoxine depletion and hypercalcaemia due to secondary hyperparathyroidism.\textsuperscript{5,10,11,53,57} Cheetahs also
suffer from gastrointestinal tract and liver disease,\textsuperscript{51,52} which could be risk factors for oxalate formation. This retrospective pathological study was conducted to describe the lesions in cheetahs with and without oxalates in their kidneys, and to investigate the relationship between oxalate crystal deposition and concurrent renal, hepatic or gastrointestinal tract disease.

**Materials and Methods**

**Study animals**

Cases were selected from pathology databases of cheetahs kept as part of the North American Association of Zoos and Aquariums' Species Survival Plan (n=297); and from cheetah pathology databases in southern Africa (n=257) and France (n=40). Tissues from cheetahs over one month old that died between 1990 and 2013 were used as the study population. Histological sections of kidneys from all cheetahs were screened using polarized light for the presence or absence of crystals. A subset of cheetahs greater than one month old that died between 1990 and 2013 (n=265) was selected for detailed histologic examination based on the quality and availability of relevant histological material. Cases were defined as cheetahs with renal oxalates at death (n=100). Controls were defined as cheetahs with no renal oxalates at death (n=165) and similar demographic characteristics to cases including age, sex, the housing facility at death, and year of death; however, matching was approximate and not possible in all instances. Geographic distribution of the final dataset of cases and controls was North America (n=94; 40 cases, 54 controls); southern Africa (n=167; 56 cases, 111 controls) and France (four cases, no controls). All of the North American and French cheetahs and all but two of the southern African cheetahs were captive at the time of death.
Pathological evaluation

Formalin-fixed tissues were processed routinely and stained with hematoxylin and eosin, von Kossa and Masson’s trichrome stains. Crystal number and the presence and severity of renal lesions, veno-occlusive disease (VOD), gastritis, enteritis and colitis were evaluated by two pathologists (KAT and EPM) using a detailed scoring scheme (Supplemental Table S1) and previously published criteria. A subset of southern African cases was examined by both KAT and EPM to confirm evaluator concordance.

Raman Spectroscopic Characterization

 Archived, formalin-fixed paraffin embedded kidney sections from cheetahs (n=47; 28 North American, 15 southern African, 4 French) were used for Raman spectroscopic characterization. Unstained, deparaffinized sections (6-8 μm) on quartz slides (25×6×1 mm³) (SPI, West Chester, PA, USA) were examined by light microscopy for crystal presence and location. A case of confirmed canine ethylene glycol toxicity was used as a control. Raman spectroscopy measurements on the kidney sections were performed on a Nicolet Almega XR Raman spectrometer equipped with an Olympus BX51 confocal microscope (Thermo Electron, Madison, WI, USA). The camera temperature was set at -50 °C through the instrument control software OMNIC 8 for Dispersive Raman. The complimentary Atlas program was employed for imaging the samples and focusing on the birefringent crystals in the tissue. Spectra were obtained with a 532 nm solid state, diode pumped laser (24 mW) at 100% power, 50 μm pinhole aperture, 2400 lines/mm resolution grating, spectral range of 2000 to 300 cm⁻¹, and a 100× Olympus objective. Individual spectra were
acquired with a 20 sec interrogation time and three exposures after a 30 sec photobleach. All spectra were baseline corrected using the OMNIC software. Library comparison for spectra was performed using the OMNIC software and spectral libraries purchased from Thermo Electron. Specifically, the HR Raman Inorganics Library (containing a calcium oxalate spectrum) as well as additional spectra obtained from a known ethylene glycol poisoning in a dog (07N-0544-Ti, U.S. FDA) were used for confirmation that a spectrum obtained from a crystal was a spectral match with calcium oxalate. Additional spectral processing, including peak identity evaluation, was performed with Spectrus Processor 2012 (ACD/Labs, Toronto, Ontario, Canada).

Statistical analysis

Because matching of cases (n=100) and controls (n=165) was only approximate in this retrospective study, unmatched analysis was performed. Age of animals at death was categorized as <1.5, 1.5-<8, 8-<11 and ≥11 years old. Univariate analyses were conducted to compare lesion prevalence and severity in cases and controls (Supplemental Table S2), as well as to compare lesion prevalence and severity with crystal number (Supplemental Table S3). Renal edema, hemorrhage, arteritis, fibrinoid necrosis, thrombosis, cortical and medullary intra-tubular mineralization, the presence of other crystals in tubules, intratubular and intracellular pigment in tubules, tubulitis and VOD were excluded as these lesions were uncommon to rare. The cellular characteristics of interstitial inflammation were excluded as this was invariably lymphoplasmacytic. The association between the presence of oxalate crystals and the presence and severity of each remaining histological lesion (case/control status) was then assessed using mixed-effects logistic regression.
Table 1. Lesions significantly associated with the presence of renal oxalate crystals in cheetahs

<table>
<thead>
<tr>
<th>Lesion</th>
<th>Odds ratio</th>
<th>95%CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Interstitial nephritis</td>
<td>5.63</td>
<td>2.10 -15.10</td>
</tr>
<tr>
<td>Renal amyloidosis</td>
<td>2.74</td>
<td>1.06 - 7.07</td>
</tr>
<tr>
<td>Colitis</td>
<td>2.40</td>
<td>0.99 - 5.83</td>
</tr>
<tr>
<td>Glomerular loop thickening</td>
<td>0.31</td>
<td>0.14 - 0.72</td>
</tr>
<tr>
<td>Gastritis</td>
<td>0.22</td>
<td>0.07 - 0.73</td>
</tr>
</tbody>
</table>

CI, confidence interval

*aAdjusted for age, examiner and region in a mixed-effects logistic regression model*
(Table 1). Firstly, all potential predictors associated with the outcome with p<0.2 in the initial analysis were included and sequentially eliminated until all remaining predictors were significant (p<0.05). Every candidate predictor was then re-tested one-by-one in the model and retained if significant. The scores for renal amyloidosis, interstitial inflammation and gastritis were collapsed into dichotomous (0/1) scores indicating the presence or absence of each lesion, since the original ordinal scores did not contribute any additional information. Odds ratios were generated describing the strength of association between the presence and absence of crystals and each lesion.

Secondly, where oxalate crystals were present (n = 100), the association of each lesion score with crystal number (average number of cortical tubules containing crystals in three 100x fields) was assessed using mixed-effects negative binomial regression, adjusting for age, examiner and institution (random effect). Factors potentially associated with the severity of oxalate nephrosis were assessed using a mixed-effects negative binomial regression model, with the same candidate variables and modelling approach as above. All statistical analyses were done using Stata 14 (StataCorp, College Station, TX, USA). Significance was assessed at p<0.05. The scores for renal amyloidosis were collapsed into dichotomous (0/1) scores indicating the presence or absence of amyloidosis in the final multiple negative binomial regression model of factors associated with the numbers of crystals present (Table 2). Negative binomial regression in this context yields a count ratio (CR), interpreted as the ratio of the crystal number in a category with the lesion to the crystal number in the reference category (without the lesion); therefore, a CR < 1 indicates a negative or inverse association and a CR > 1 indicates a positive association with crystal number.
Table 2. Age, region and lesions associated with the number of crystals in cheetah kidneys\textsuperscript{a}

<table>
<thead>
<tr>
<th>Variable</th>
<th>Count ratio\textsuperscript{b}</th>
<th>95%CI</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Glomerulosclerosis</td>
<td>0.84</td>
<td>0.75 - 0.95</td>
<td>0.006</td>
</tr>
<tr>
<td>Amyloidosis score</td>
<td>0.60</td>
<td>0.39 - 0.94</td>
<td>0.03</td>
</tr>
<tr>
<td>Interstitial nephritis</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>absent</td>
<td>1\textsuperscript{c}</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>mild</td>
<td>0.70</td>
<td>0.40 - 1.25</td>
<td>0.23</td>
</tr>
<tr>
<td>moderate</td>
<td>1.13</td>
<td>0.60 - 2.16</td>
<td>0.70</td>
</tr>
<tr>
<td>severe</td>
<td>0.07</td>
<td>0.02 - 0.31</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Age</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>&lt;1.5 years</td>
<td>1\textsuperscript{c}</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>1.5 – 8 years</td>
<td>0.50</td>
<td>0.21 - 1.18</td>
<td>0.11</td>
</tr>
<tr>
<td>8 – 11 years</td>
<td>0.48</td>
<td>0.20 - 1.15</td>
<td>0.10</td>
</tr>
<tr>
<td>&gt;11 years</td>
<td>0.33</td>
<td>0.14 - 0.80</td>
<td>0.01</td>
</tr>
<tr>
<td>Region</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Africa</td>
<td>1\textsuperscript{c}</td>
<td>–</td>
<td>–</td>
</tr>
<tr>
<td>USA</td>
<td>1.82</td>
<td>1.14 - 2.88</td>
<td>0.01</td>
</tr>
<tr>
<td>Europe</td>
<td>2.53</td>
<td>1.06 - 6.06</td>
<td>0.04</td>
</tr>
</tbody>
</table>

CI, Confidence interval

\textsuperscript{a}Adjusted for age, examiner and region in a mixed-effects negative binomial regression model
The count ratio is the ratio of the crystal count in animals in that category to the crystal count in animals in the reference category; or, the fold change in crystal count for each unit increase in the glomerulosclerosis or amyloidosis score.

Reference category
Results

Histopathology

Histologically, kidneys from cheetah cases contained small to very large numbers of colorless refractive crystals (average 0.3-81 crystals in three 100x fields) in cortical and medullary tubules forming rosettes, globules and acicular fragments (Fig. 1) that were birefringent with polarized light (Fig. 2). Crystals were not uniformly distributed in the renal cortex, with clusters apparently occurring in single or groups of tubules; crystals were not associated with tracts of inflammation. Affected tubules contained small amounts of sloughed necrotic cellular debris mixed with variable amounts of pale amorphous eosinophilic material and, in many cases, were lined by a discontinuous layer of epithelial cells with variable degrees of necrosis characterized by shrunken cells with hypereosinophilic cytoplasm and dark basophilic pyknotic nuclei (Fig. 1). Adjacent tubules were variably dilated and lined by regenerative low cuboidal epithelial cells with pale basophilic cytoplasm and large crowded oval nuclei (Fig. 3). Variable degrees of mild interstitial fibrosis and small numbers of intra-tubular cellular casts were present. Crystals and fine mineral deposits variably present on tubular basement membranes in the cortex and medulla stained variably positive with von Kossa which was interpreted as the presence of calcium phosphate and/or carbonate salts, since melanin pigment was not visible in crystals or on tubular basement membranes on hematoxylin and eosin stains (Fig. 4). Renal lesions were present histologically in 46 of 165 (28%) cheetahs without oxalate crystals and in 87 of 100 (87%) cheetahs with oxalate crystals.

Additional renal lesions included small numbers of deeply basophilic mineralized crystals in the lumina of both cortical and medullary tubules, rare pale eosinophilic or

Figure 1. Oxalate crystals are associated with epithelial necrosis and tubular dilatation; the glomerulus is normal. Hematoxylin and eosin (HE).

Figure 2. Polarization highlights moderate numbers of oxalate crystals in tubular lumina (an average of 37 crystals per three 100 fields). HE.

Figure 3. Cortical tubular epithelial necrosis, sloughed epithelial cells in the tubular lumina, and regeneration of tubular epithelium. HE.

Figure 4. Oxalate crystals and cortical tubular basement membranes are mineralized. von Kossa.
yellow pigment in renal tubular epithelial cells or in tubular lumina; and variable microvesiculation consistent with lipidosis and or hydropic degeneration in renal tubular epithelial cell cytoplasm. Tubulitis, renal edema or hemorrhage, arteritis, vascular fibrinoid necrosis were rarely noted.

**Raman spectroscopy**

On Raman spectroscopic examination, crystals were easily observed as bright spheres in the dark tissue background. Crystals in the control case of canine ethylene glycol poisoning and in 27 (96%) North American, 14 (93%) southern African, and four (100%) French cases were confirmed as calcium oxalate with no laser photodegradation of the crystals (Fig. 5). Laser photodamage prevented crystal identification in two cheetahs (one North American and one southern African). In addition to calcium oxalate crystals, calcium carbonate, calcium phosphate or calcium sulphate crystals were present in the kidney tubules of three North American cheetahs, and one additional North American cheetah sample had a Raman multi-component spectrum that contained calcium oxalate bands. All of these cheetahs were >10 years of age and had other underlying renal disease including glomerulosclerosis and amyloidosis.

**Statistical analysis**

The presence and numbers of crystals were not significantly different between male and female cheetahs ($p=0.454$). The average age of cheetahs with renal oxalate crystals at death was similar in the North American, French and southern African populations (3.7 months - 15.9 years old; median = 8.7 years (interquartile range [IQR]: 6.4-11.3 years). Age distribution was similar in cheetahs without oxalates
Figure 5. Representative Raman spectrum of a crystal from an 8-year-old captive female cheetah from South Africa. Characteristic bands for calcium oxalate monohydrate are assigned to the C-O asymmetric stretch (1628 cm\(^{-1}\)), C-O symmetric stretch (1490 and 1463 cm\(^{-1}\)), C-C stretch (896 cm\(^{-1}\)), and O-C-O in-plane bending (502 cm\(^{-1}\)).
Figure 6. Association between cheetah age and average number of oxalate crystals in renal tubular lumina.
(median 8.6 years; IQR: 3.6-11.3 years). Cheetahs older than 1.5 years were much more likely to have oxalates than younger cheetahs (p=0.034, Supplemental Table S2) but cheetahs <1.5 years old had significantly more crystals than older cheetahs (p<0.001, Supplemental Table S3). Although three young cheetahs from one North American institution had very high numbers of crystals (average of 69-81 (per three 100x fields)) no grouping of crystal number based on age was present (Fig. 6). On the multivariate mixed effects negative binomial regression analysis cheetahs >11 years old had significantly fewer crystals than those < 1.5 years old (p=0.01, Table 2).

When comparing cheetahs with renal oxalate crystals (cases) with those without such crystals (controls), tubular changes were significantly associated with the presence of crystals (Supplemental Table S2). Cases had significantly more renal tubular necrosis, regeneration, dilatation, protein and cellular casts than controls (p≤0.001 for each parameter). Compared to controls, cases also had significantly more lymphoplasmacytic interstitial nephritis (p≤0.001), amyloidosis (p≤0.001), chronic infarcts (p≤0.001), efferent arteriole arteriolosclerosis (p≤0.001), arcuate artery arteriosclerosis (p=0.011), cortical (p=0.02) and medullary (p=0.005) tubular basement membrane mineralization, cortical fibrosis (0.049), hyperplastic collecting tubule epithelium (p=0.063), enteritis (p=0.001) and colitis (p=0.017). In contrast, gastritis was significantly more common in controls than cases (p=0.012). Sex (p=0.899), glomerulosclerosis (p=0.952), glomerular loop thickening (p=0.273), glomerular hypercellularity (0.743), and medullary fibrosis (p=0.898) had no effect on the likelihood of having renal oxalate crystals. In the 250 cheetahs for which liver was available hepatic VOD was present in 22 of 88 (25%) of cases and 32 of 162 (20%) of controls. In addition, 6% of cheetahs without oxalates but only 3% of
cheetahs with oxalates had severe VOD. In the multivariable mixed effects logistic regression analysis comparing cases and controls, the only factors associated with an increased likelihood of having renal oxalates were interstitial nephritis, and to a lesser extent, renal medullary amyloidosis and colitis (Table 1). Case cheetahs had reduced odds of having glomerular loop thickening and gastritis compared to controls.

When crystal number was evaluated in cheetahs with oxalate crystals in the kidney (Supplemental Table S3), the number of crystals present was positively associated with tubular necrosis (p≤0.001), regeneration (p=0.015), cellular casts (p≤0.001), glomerular hypercellularity (p=0.007) and arteriosclerosis of the efferent arteriole (p≤0.001). Crystal number was inversely associated with protein casts (p=0.043), medullary tubular basement membrane mineralization (p=0.002), collecting duct epithelial hyperplasia (p=0.023), glomerulosclerosis (p=0.009), glomerular loop thickening (p≤0.001), amyloidosis (p≤0.001), chronic infarcts (p=0.001), renal cortical (p≤0.001) and medullary (p=0.002) fibrosis, enteritis (p=0.001) and colitis (p=0.019). In 31 cheetahs with large numbers of oxalates (average of 10-81 per three 100x fields), there was no evidence of any underlying renal disease. Crystal number was unrelated to sex (p=0.455), tubular dilatation (p=0.409), cortical tubular basement membrane mineralization (p=0.382), arteriosclerosis of the arcuate artery (p=0.118) or gastritis (p=0.826). The relationship between crystal number and interstitial nephritis was significant but non-linear (p≤0.001) since cheetahs with grade two nephritis had more crystals than those with nephritis grades zero, one and three. In the multivariable mixed effects negative binomial regression analysis, geographical region was the only factor positively associated with crystal number (Table 2). On average, cheetahs from the
USA and France had significantly more oxalate crystals than those from Africa (p=0.01). Nine of the 10 cheetahs with highest average number of cortical tubules containing oxalate crystals (26-81) were from North America (5 different institutions), and the remaining one was from a South African institution (24). Among the 10 southern African cheetahs with the highest number of crystals (14-24), cheetahs from one institution were over-represented (50%). The four cheetahs from three French institutions had an average number of cortical tubules containing crystals of 15 (10-20). One free-ranging Glomerulosclerosis and amyloidosis were negatively associated with crystal number in the multivariate analysis (Table 2).

Discussion

This study found that the presence of intra-tubular renal oxalate crystals was associated with renal tubular necrosis, regeneration and the presence of tubular casts in cheetahs in North America, southern Africa and France; and that increased numbers of crystals was associated with increased severity of these tubular changes. This effect of oxalate crystals on tubular epithelium is well documented in humans and animals.11,28,43,44,49,66,78,83

Raman spectroscopy confirmed that the majority of crystals were calcium oxalate. Melamine and other renal crystals were ruled out because the Raman spectral fingerprints did not match these compounds.37,61,68 The two cases in which oxalate crystals were not found were cheetahs with only a few crystals in the H&E stained sections, and it is possible that sections prepared for spectroscopy contained no crystals or that they were damaged by the laser. Calcium carbonate, calcium phosphate or calcium sulphate crystals, present in the kidney tubules of three North
American cheetahs, were not distinguishable from calcium oxalate on routinely stained sections but were readily identifiable by their Raman spectra. Therefore, while calcium oxalate monohydrate was the predominant crystal noted other crystal types can occur in cheetahs. Compound oxalate crystals have also been described in domestic dogs.  

Oxalate crystals and associated tubular damage occurred in 31 cheetahs without other renal diseases, all of which were reported to have clinical evidence of renal failure (inappetance, weakness, vomiting, azotemia, dehydration, polyuria and depression) and showed varying degrees of parathyroid gland hyperplasia and metastatic mineralization but no other renal disease. Similar findings have been described previously in cheetahs and other animals with oxalate nephrosis. Oxalate nephrosis, therefore, occurs independently of other renal diseases in some captive cheetahs and should be considered in the differential diagnosis of renal disease. Although rare scattered oxalate crystals may occur secondarily to many types of chronic renal disease, cheetahs with chronic renal disease also had large numbers of crystals suggesting that oxalate nephrosis may co-exist with other renal diseases at least in some cheetahs. Tubular lesions, glomerular hypercellularity and efferent arteriolosclerosis could contribute to oxalate crystal formation in chronic renal disease as these were the only renal lesions positively associated with crystal number in these cheetahs. 

While a few oxalate crystals may occur secondarily to concurrent chronic renal disease in some cases, factors other than concurrent renal disease likely affect renal oxalate deposition in cheetahs. 

No clear evidence was found in our study that oxalate nephrosis occurs secondary to gastro-intestinal or hepatic disease in cheetahs. Gastritis was more
common in controls than cases, and was unrelated to crystal number. On the
univariate analysis, more cheetahs with renal oxalate crystals had enteritis and colitis
than controls, and on the multivariate analysis colitis was associated with an
increased likelihood of having renal oxalates. However, both lesions were inversely
associated with crystal number and therefore likely do not play a key role in the
development of oxalate nephrosis. In humans, fat malabsorption due to enterocolitis
or pancreatic insufficiency may result in reduced intestinal calcium which increases
intestinal oxalate absorption because calcium binds intestinal oxalates\textsuperscript{5,10,46,56,81} However, no relationship between enterocolitis and renal oxalate crystal number was
found and pancreatic insufficiency is rare in the parent population of cheetahs.\textsuperscript{51,52}
Veno-occlusive disease was uncommon in animals with and without renal oxalate
crystals. Increased intestinal absorption of oxalates may also result from loss of
intestinal lactic acid bacteria, including \textit{Oxalobacter formigenes}, that degrade oxalate
into carbon dioxide and formate.\textsuperscript{42,46,70,82} Research is needed to determine whether
or not cheetahs that receive broad-spectrum antibiotic therapy for \textit{Helicobacter}-associated gastritis or other infections have depleted \textit{O. formigenes} populations,
which may predispose them to oxalate nephrosis.\textsuperscript{35}

Significant geographical and institutional clustering of cheetahs) with abundant
renal oxalate crystals suggests that genetic, dietary or management factors
may influence the prevalence of oxalate nephrosis in captive cheetahs. So far, no
proof of ethylene glycol toxicity has been found in cheetahs, but exposure to
ethylene glycol, xylitol or other oxalate precursors should be thoroughly investigated
in cases of oxalate nephrosis. Identification of dietary factors was beyond the scope
of this retrospective necropsy study. However evaluation of the fluid intake and
protein, fat, collagen, feathers, calcium, magnesium, beet pulp, pyridoxine, ascorbic
acid and arachidonic acid levels in captive cheetah diets is needed as these factors can influence oxalate excretion. Dietary calcium:oxalate ratios affect O. formigenes colonization of the intestine in rats so this ratio may be important in cheetah diets. Obesity and stress may contribute to oxalate nephrosis in humans, and their roles in oxalate nephrosis may warrant further study since obesity may be present in underactive captive cheetahs and stress has been documented in captive animals.

Primary hyperoxaluria (PH) in humans is an autosomal recessive condition associated with mutations in three genes involved in oxalate metabolism: AGT gene in PH type 1; GRHPR gene in PH type 2; and 4-hydroxy-2-oxoglutarate aldolase (HOGA1) in PH type 3. Similarly, both PH type 1 and PH type 2 have been described in domestic cats. Clear evidence that oxalate nephrosis in cheetahs is a primary genetic disease was not found in our study, however we did not test genetic relatedness. In humans and cats with inherited oxalate nephrosis the disease is seen in juveniles. In this study, although cheetahs <1.5 years old had significantly more crystals than older cheetahs, which might indicate an inherited predisposition, relatively large numbers of crystals were seen in cheetahs in all ages. Lesions which are characteristic of primary disease in humans and cats, including granulomatous nephritis associated with interstitial oxalates, widespread tissue oxalate deposition and neurological disease were not seen in the cheetahs in this study. Manifestation of genetic disease is complex with environmental influences, multigenetic inheritance, epigenetic effects, incomplete penetrance and variable expression resulting in differing disease severity and phenotypes. Therefore, whether variations in crystal number are due to differing manifestations of the same disease or differences in pathogenesis among cheetahs...
in this study is uncertain. Since cheetahs from the USA, southern African and French cheetah populations share common founders a detailed genetic analysis, including sequencing of key genes involved in oxalate metabolism, and resultant enzyme activity, is needed to determine whether or not cheetahs suffer from a genetically limited capacity for glyoxylate metabolism.48

Early diagnosis of oxalate nephrosis is important since, this disease is potentially treated by fluid therapy to reduce oxaluria44,85, pyridoxine to stabilize and enhance the activity of AGT,5,7,23,65 urine alkalinisation,66,85 and n-3 fatty acid supplementation to decrease urinary oxalate excretion and free-radical injury.5,24,38,46,57,62,66,70

Urinalysis data was limited to a few cases in this retrospective study, but could be a valuable diagnostic tool since small numbers of oxalate crystals may be present in the urine of affected cheetahs.30 However, confirming the presence of oxalate nephrosis and differentiating it from concurrent glomerulosclerosis or renal medullary amyloidosis, in azotemic cheetahs is not simple if urinary oxalates are absent. Renal ultrasonography is non-specific in domestic dogs and cats with oxalate nephrosis.1 Computed tomography scans are used to identify renal crystals in humans,5 but are not a practical diagnostic modality in cheetahs. At one Southern African institution, urinary oxalate was elevated in an acutely azotemic cheetah later diagnosed with oxalate nephrosis without other renal lesions (8000 mg oxalate/gram creatinine, compared 128 mg /g in a healthy enclosure mate F. Reyers, Golden VetPath, unpublished data). Since crystal number was higher in young cheetahs and chronic renal disease in cheetahs is age-related,25,51 oxalate nephrosis should be a primary consideration in younger cheetahs with renal failure.

In conclusion, this study found that oxalate nephrosis unrelated to ethylene glycol toxicity occurs in cheetahs. We found no convincing evidence that oxalate nephrosis
in captive cheetahs is secondary to renal, hepatic or gastro-intestinal tract disease although it may occur concurrently with such diseases. Oxalate nephrosis should be considered as a differential diagnosis in captive cheetahs with renal failure, especially in young animals that are less likely to be suffering from renal amyloidosis or glomerulosclerosis. Oxalate nephrosis is of uncertain etiology in captive cheetahs but a multifactorial pathogenesis, including a primary genetic predisposition, diet, altered gut microbiome and or stress, is suspected.

The data analyzed in this study are available as Supplemental Materials.

Acknowledgements
This study was made possible by the veterinarians and cheetah holding and breeding institutions that submitted clinical data and carcasses from captive and free-ranging cheetahs for examination from the USA: The Association of Zoos and Aquariums’ Cheetah Species Survival Plan including Anonymous (5 facilities), Albuquerque Biopark Zoo, Binder Park Zoo, Brevard Zoo, Caldwell Zoo, Chehaw Wild Animal Park, Cincinnati Zoo and Botanical Garden, Cleveland Metroparks Zoo, Columbus Zoo and Aquarium, Dickerson Park Zoo, Fort Wayne Children’s Zoo, Fort Worth Zoo, Fossil Rim Wildlife Center, Honolulu Zoo, Jackson Zoo, Jacksonville Zoo and Gardens, Kansas City Zoo, The Living Desert, Maryland Zoo in Baltimore, Mesker Park Zoo and Botanic Gardens, Montgomery Zoo, Nashville Zoo at Grassmere, Oklahoma City Zoo and Botanical Garden, Phoenix Zoo, San Antonio Zoo and Aquarium, San Diego Zoo Global, Tulsa Zoo and Living Museum, Utah’s Hogle Zoo, White Oak, Wildlife Safari, Zoo Knoxville, and Zoo New England); from southern Africa: AfriCat, Cango Wildlife Ranch, Cheetah Conservation Botswana, Dr.
A. Tordiffe, Dr. D. Zimmerman, Dr. K. Good, Dr. P. Buss, Dr. P. Caldwell, Dr. P. Swartz, Farm Inn, Hoedspruit Endangered Species Centre, Johannesburg Zoo, Letsatsi La Africa, Lory Park Zoo, Matobo Veterinary Centre, National Zoological Gardens of South Africa, Rhino and Lion Park, Seaview Predator Park, The Ann van Dyk Cheetah Centre, Tshwane Nature Conservation; from France: the Safari de Peaugres, Dr. C. Vitaud, Dr. D. Sarran; Parc des Félins, Mr. G. Breton, Dr. F. Ollivet-Courtois. Data and cases provided by Drs. Linda Munson and Nadia Robert are also acknowledged. Pathology laboratory staff at the NZG and Faculty of Veterinary Science, University of Pretoria, at Anipath, France, and the University of Illinois Zoological Pathology Program and Veterinary Diagnostic Laboratory provided excellent technical assistance. The authors thank Dr. Chuck Mohr for his guidance in developing the renal scoring system. Professors John Lawrence and Leon Prozesky, Faculty of Veterinary Science, University of Pretoria provided valuable editorial input. The National Research Foundation, through a core grant to the National Zoological Gardens of South Africa, provided the funding to conduct pathological examinations in southern African cheetahs. The views expressed in this publication are those of the authors and do not necessarily reflect the official policy of the Department of Health and Human Services, the U.S. Food and Drug Administration, or the U.S. Government.
References


17. Dijcker JC, Hagen-Plantinga EA, Hendriks WH. Changes in dietary macronutrient profile do not appear to affect endogenous urinary oxalate


33. Jansen JH, Arnesen K. Oxalate nephropathy in a Tibetan spaniel litter - a


41. Lemberger K, Sarran D, Robert Terio, K N. Renal oxalate nephrosis in several


59. Papendick RE, Munson L, O’Brien TD, Johnson KH. Systemic AA amyloidosis


68. Selvaraju R, Raja A, Thiruppathi G. FT-Raman spectral analysis of human


76. Spelman LH, Cambre RC, Pessier AP, et al. Renal oxalosis in a cheetah


