Characterization of the intrarenal renin-angiotensin system in cats with naturally occurring chronic kidney disease

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Abstract

Background: The role of the renin-angiotensin-aldosterone system in cats with chronic kidney disease (CKD) is incompletely understood.

Objective: To characterize components of the intrarenal renin-angiotensin system (RAS) in cats with CKD.

Animals: Eleven cats with naturally occurring CKD (CKD group) and 8 healthy control cats.

Methods: Renal tissue samples were evaluated by reverse-transcription polymerase chain reaction for renin, angiotensinogen, angiotensin-converting enzyme (ACE), and angiotensin II type 1 receptor transcript levels, and by liquid chromatography-mass spectrometry for quantification of angiotensin I, II, III, and IV concentrations. Linear mixed models were used to compare gene transcript levels and concentrations of angiotensin peptides between groups.

Results: Cats of the CKD group were significantly older (P < .001) and more likely to be neutered (P = .007) than healthy control cats. Kidneys from cats with CKD had significantly higher transcript levels of angiotensinogen (P < .001) and lower transcript levels of ACE (P < .001) than those from control cats. Renal angiotensin I concentrations were increased in CKD compared with control kidneys (P = .001). No other significant differences in renal transcript levels or angiotensin peptide concentrations were noted between groups.

Conclusion and Clinical Importance: The intrarenal RAS might be activated in cats with CKD. Small sample size and differences in age, neuter status, and dietary sodium intake between groups might have limited the ability to identify a significant difference in concentration of renal angiotensin II.

KEYWORDS
angiotensin-converting enzyme, angiotensinogen, equilibrium analysis, tissular RAAS

Abbreviations: ACE, angiotensin-converting enzyme; ACE, gene for angiotensin-converting enzyme; AGT, angiotensinogen; AGT, gene for angiotensinogen; Ang, angiotensin; AGTR1, gene for angiotensin II type 1 receptor; BP, blood pressure; CKD, chronic kidney disease; GAPDH, gene for glyceraldehyde 3-phosphate dehydrogenase; LC-MS/MS, liquid chromatography-mass spectrometry/mass spectrometry; PRA, plasma renin activity; RAS, renin-angiotensin system; REN, gene for renin; RPS7, ribosomal protein S7; SBP, systolic blood pressure.

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1 | INTRODUCTION

The renin-angiotensin-aldosterone system (RAAS) is a major regulator of systemic arterial blood pressure (BP) and fluid and electrolyte balance. Its main effector peptide, angiotensin (Ang) II, promotes renal and gastrointestinal sodium reabsorption, aldosterone secretion, vascular smooth muscle contraction, and diuresis, pressor and tachycardic responses, while inhibiting renin release—actions that are achieved predominantly through interaction with the Ang II type 1 receptor. In health, these responses conserve sodium and maintain normal circulating blood volume and BP. Interaction of Ang II with the Ang II type 1 receptor also induces the expression, transcription or both of growth factors and proliferative cytokines, including transforming growth factor-β1, and perpetuates the production of nephrotoxic reactive oxygen species, leading to inflammation, fibroblast formation, and collagen deposition. As a result, in human beings, chronic activation of the RAAS is recognized as a critical factor in the progression of chronic kidney disease (CKD), and its inhibition is an integral part of renal disease treatment.

Studies characterizing the circulating RAAS in cats with CKD have yielded variable results. As a population, cats with naturally occurring kidney disease and systemic arterial hypertension have lower mean plasma renin activity (PRA) and higher mean plasma aldosterone concentration than healthy controls, suggesting that RAAS activation may not play a major role in the pathogenesis of CKD-associated hypertension in this species. However, in a subset of these patients, evidence of RAAS activation was suggested by PRA and aldosterone concentrations that fell inappropriately within or above associated reference ranges. Similarly, in experimentally induced CKD in cats, both PRA and circulating aldosterone concentrations were increased in some, but not all, described models.

Recently, focus on the role of the RAAS in the pathophysiology of organ injury has been extended to that of the local, tissue renin-angiotensin system (RAS). Rodent models suggest that the kidney is capable of producing all RAS components, allowing local formation of Ang II by pathways that are subject to feedback mechanisms that operate independently of the traditional, systemic RAAS. For example, data from rodent models of systemic hypertension suggest the intrarenal RAS can be active in the absence of evidence of systemic RAS activation, with hypertensive individuals experiencing appropriately suppressed PRA and juxtaglomerular renin immunoreactivity along with increased distal nephron renin expression and local angiotensinogen (AGT) production, the latter combination contributing to increased intratubular Ang I and Ang II concentrations.

Little work has been done to characterize the intrarenal RAS in companion animal species. Although a study of cats with CKD reported tubular and interstitial Ang II immunoreactivity that correlated with glomerulosclerosis and cellular infiltration, but not tubulointerstitial fibrosis, most investigations to date have evaluated components of the systemic RAAS, leaving the importance of the local tissue system largely undescribed.

Our objective was to characterize the renal expression of RAAS components in healthy cats and in cats with naturally occurring CKD. We hypothesized that compared to kidneys from healthy control cats, those from cats with CKD would demonstrate increased transcript levels of genes encoding RAS proteins and increased concentrations of angiotensin peptides.

2 | MATERIALS AND METHODS

2.1 | Study design

Our study was cross-sectional and performed on banked renal tissue samples collected from client-owned cats with naturally occurring CKD (CKD group), and healthy control cats (control group) as part of a separate study designed to investigate pathways of renal fibrosis.

All activities related to the original study were approved by the University of Georgia Institutional Animal Care and Use Committee (Animal Use Protocol A2017 05-008-Y3-A1).

2.2 | Animals and enrollment criteria

Enrollment criteria used in the study from which renal tissues originated, including methods used to establish a diagnosis of CKD, have been published previously. Briefly, renal tissues from the CKD group were obtained from client-owned cats presented to primary care and referral veterinary centers for euthanasia. Available clinicopathologic data reflecting renal function and systemic arterial BP were recorded upon review of each cat’s medical record. Cats were excluded if they were known to have received a RAAS antagonist (ie, an angiotensin-converting enzyme [ACE] inhibitor, angiotensin receptor blocker, or mineralocorticoid receptor antagonist) or a short-acting corticosteroid in the 14 days preceding euthanasia, or if they received a depot corticosteroid injection in the 6 months before euthanasia. For the present study, samples also were excluded if cats had received other medications with known impact on the RAAS (eg, beta-adrenergic blockers, calcium-channel blockers, furosemide) in the 14 days preceding euthanasia.

Renal tissues for the healthy control group were obtained from adult, purpose-bred, mixed breed, male neutered cats participating in unrelated terminal studies having no impact on renal structure and function, and from adult cats that were euthanized as part of population control measures at a local animal control facility. Blood and urine samples were collected immediately before or after euthanasia with pentobarbital. Complete blood count, serum biochemistry profiles, urinalysis and urine protein-to-creatinine ratios (Clinical Pathology Laboratory of the University of Georgia), and symmetric dimethylarginine concentration measurement (IDEXX Laboratories) were performed for all control cats. Control cats were deemed healthy based on normal findings of physical examination and necropsy and were deemed to have normal renal function and structure on the basis of renal histology, serum biochemistry, and urinalyses. Cats obtained from the animal control facility population were screened for retroviral infections using feline immunodeficiency virus antibody and feline leukemia virus antibody.
2.3 | Renal tissue sample collection

For all cats, renal tissues were collected within 1 hour of euthanasia. Kidneys were removed through a midline laparotomy and sectioned longitudinally. One-half of each kidney was minced and placed in RNA stabilization solution (RNAlater, QIAGEN, Valencia, California). Thus, renal cortex and medulla, in proportions equal to those present in vivo, were present in each specimen. After overnight incubation at 4°C, tissues were removed from RNA stabilization solution, homogenized using a mortar and pestle, divided into 30 mg aliquots, and stored at −80°C until further analysis.

2.4 | Renal RAS gene transcription

Reverse-transcription and quantitative PCR were performed to quantify transcript levels of AGT, renin (REN), ACE, and Ang II type 1 receptor (AGTR1) in banked renal tissue homogenates from each kidney from all cats in the CKD and control groups. Feline gene specific primers were selected from previously reported studies23,24 when available, or designed using oligonucleotide prediction software (Table 1).25

For each sample, total RNA was extracted from 30 mg of tissue homogenate using a commercially available RNA extraction kit (RNeasy Plus Mini Kit, QIAGEN, Valencia, California). Integrity of the isolated RNA was confirmed by quantification using a spectrophotometer (NanoDrop Spectrophotometer, Thermo Fisher Scientific, Waltham, Massachusetts), followed by visualization of 18S and 28S ribosomal bands on 1.2% agarose gels. A total of 1 μg of RNA extracted from each sample was treated with deoxyribonuclease (ezDNase, Invitrogen, Carlsbad, California), and reverse-transcribed using a cDNA reaction master mix (SuperScript IV VILO Master Mix, Invitrogen, Carlsbad, California). Quantification was performed in an automated cycler (CFX96, Bio-Rad Laboratories, Hercules, California) using 20 μL reactions containing 10 μL of SYBR Green Supermix (SsoAdvanced SYBR Green Supermix, Bio-Rad Laboratories, Hercules, California), 5 PMol of each primer (Table 1), and 9 μL of complementary DNA sample at a 1:40 dilution. Thermal cycling conditions consisted of an activation step at 95°C for 30 seconds, followed by 36 amplification cycles (95°C for 15 seconds for denaturation and 60°C for 30 seconds for annealing and extension), and a melt curve step (60-95°C, increasing at increments of 0.5°C every 5 seconds). All reactions were performed in triplicate and average values were used for further analyses. A no reverse transcriptase control was included for each sample and 3 no template controls were used in each plate. In accordance with the sample maximization method,26 for each gene, all samples were analyzed in two 96-well plates. A standard sample was analyzed in triplicate with glyceraldehyde 3-phosphate dehydrogenase (GAPDH) primers in each plate and used as an inter-run control for the entire gene study.

Transcript levels of AGT, REN, ACE, and AGTR1 were normalized to those of 2 reference genes, GAPDH and ribosomal protein S7 (RPS7), using the GENorm method.27 Reference genes were selected based on the efficiency of their primers, and their stability value (M value <0.5) and coefficient of variation (<25%) when tested in a complete set of the experiment's samples. After normalization, transcript levels were scaled to those of the lowest sample for each target gene.

 Primer efficiencies were calculated using the Pfaffl method based on standard curves performed with serial dilutions of cDNA (factor of 2 or 4; 3 replicates) starting at a 1:20 dilution. Quantitative PCR products were confirmed on 1.2% agarose gels. Amplicons were cloned into a vector system (PGEM-T easy vector, Promega, Madison, WI) and sequenced bi-directionally by chain termination (Molecular Cloning Laboratories, San Francisco, California). Resulting sequences were confirmed using the National Center for Biotechnology Information Basic Local Alignment Search Tool against the feline genome.28

### Table 1 | Primer sequences for quantitative PCR

| Gene   | Ensemble ID/ NCBI Access. No. | Primer sequence (5’ → 3’) | Fragment size (bp) | Source       |
|--------|-------------------------------|---------------------------|--------------------|--------------|
| GAPDH  | ENSEFCAG 00000006874          | for: GCTGCCCAAGACATCCATCC rev: GTGATCCAGGACGGACAC | 134                | Riedel et al.23 |
| RPS7   | NM_001009832                  | for: GTCCCGAAGCCGGCAGCTTT T rev: CACATCTGGCCTGGGAA A | 74                 | Kessler et al.24 |
| AGT    | XM_011287265.1                | for: TCT TGA CGT GGC TGA AGA ATC rev: AGC AGG TCC TGA AGG TCA TA | 94                 | IDT PrimerQuest25 |
| REN    | ENSEFCAG 00000002902          | for: CGA CCC CCA GTA TTA CCA AG rev: CCT CCTCAC AGA CCA AGG TG | 116                | Riedel et al.23 |
| ACE    | XM_003997083.1                | for: GGT GAC ACC CCT GTA CCA AG rev: CCT GGT TTC AGT CTC CA | 133                | Riedel et al.23 |
| AGTR1  | ENSEFCAT00000000716           | for: AAC ATC GCT TCA GCC AGT GT rev: TTT GCC ATC AGC ATT GTA CG | 134                | Riedel et al.23 |

Abbreviations: ACE, gene for angiotensin-converting enzyme; AGT, gene for angiotensinogen; AGTR1, gene for angiotensin II type 1 receptor; GAPDH, gene for glyceraldehyde 3-phosphate dehydrogenase; IDT, integrated DNA technologies; REN, gene for renin; RPS7, gene for ribosomal protein S7.
2.5 | Renal angiotensin peptide equilibrium analysis

Equilibrium concentrations of Ang I (ie, Ang 1-10), Ang II (ie, Ang 1-8), Ang III (ie, Ang 2-8), and Ang IV (ie, Ang 3-8) were quantified in RNA-stabilization-solution-preserved renal tissue homogenates by a commercial laboratory (Attoquant Diagnostics, Vienna, Austria) using previously described methods.29 Each angiotensin peptide was measured in samples from both kidneys in CKD group cats, and from the right kidney only in control group cats. Briefly, frozen renal tissue samples (80-100 mg) were homogenized using a mortar and pestle under liquid nitrogen. The frozen tissue powder was dissolved at 100 mg/mL in 6M aqueous guanidinium chloride (Guanidinium chloride, Sigma-Aldrich, Steinheim, Germany) supplemented with 1% (v/v) trifluoroacetic acid (Trifluoroacetic acid, Sigma-Aldrich, Steinheim, Germany) by cooled sonication using a 2-mm microtip (2-mm microtip, Sonics and Materials, Newton, New Jersey). Stable isotope-labeled internal standards for individual angiotensin metabolites were added to tissue homogenates at 200 pg/mL and stored at −80°C until analysis. Liquid chromatography-mass spectrometry/mass spectrometry (LC-MS/MS) analysis for tissue angiotensin quantification was performed using a reversed-phase analytical column operating in line with a mass spectrometer (Xevo TQ-S triple quadruple mass spectrometer, Waters Corporation, Milford, Massachusetts).

The LC-MS/MS analytical assay for biological samples was validated according to the European Medicine Agency’s Guideline on bioanalytical method validation adopted on 21 July 2011 (https://www.ema.europa.eu/en/documents/scientific-guideline/guideline-bioanalytical-method-validation_en.pdf). Between- and within-run precision and accuracy of the assay were <15% at the lower limit of quantification and <5% in the mid-range of quantification for all analytes of interest.

2.6 | Statistical analyses

Distributions of values for clinical variables by study group were examined for normality by visual assessment of histograms and normal quantile plot, and the Shapiro-Wilk test. Normally distributed clinical data are presented as mean ± SD and compared between groups using the 2-sample, unequal variance t-test. Non-normally distributed data are presented as median (range) and compared between groups using the Mann-Whitney U test. For the purposes of statistical analyses, urine specific gravity values reported as >1.060 and serum urea nitrogen concentrations reported as >100 mg/dL or >180 mg/dL were assigned values of 1.061, 101, and 181, respectively. Categorical data were compared using the Fisher’s exact test.

Differences between groups with respect to gene transcript levels and angiotensin peptide concentrations were assessed using linear mixed models with cat included as a random effect to account for repeated measurements (ie, analyses of each kidney from the same individual), and group as a fixed nominal effect. Models used restricted maximum likelihood estimation, and degrees of freedom were computed using the Kenward-Roger procedure. Models with different residual covariance structures were compared using Akaike’s information criterion to determine which model provided the best fit to the data. Normality of residuals was assessed using normal quantile plots and the Shapiro-Wilk test. Positively skewed variables were log-transformed before analysis to improve the approximation to a normal distribution. Geometric means and 95% confidence intervals (CI) were calculated by back transforming the means and confidence limits that were estimated on the log-transformed scale. Fisher’s exact test was used to compare groups with respect the proportion of cats that had angiotensin IV concentrations below the assay’s lower limit of quantification (ie, <4 fmol/g). All tests assumed a 2-sided alternative hypothesis and P-values <.05 were considered significant. Analyses were performed using commercially available software (Stata version 16.0, StataCorp LLC, College Station, Texas; GraphPad Prism for Mac, version 8, GraphPad Software Inc, La Jolla, California).

3 | RESULTS

3.1 | Animals and samples

Tissue samples from 11 cats with CKD and 8 healthy cats were included. Samples from 1 cat with CKD, which was enrolled in the original study,22 were excluded from the present study because the cat had been receiving atenolol chronically. A summary of the demographic and clinical data for cats from which tissues for the present study originated is presented in Table 2. Cats in the CKD group were significantly older (P < .001) than control cats. Two CKD cats, aged 3.1 and 4.4 years, were considered young adults according to the American Association of Feline Practitioners/American Animal Hospital Association life stage guidelines,30 with all others being >9.1 years old. Relative to control cats, cats in the CKD group were also more likely to be neutered (P = .007). Samples from cats with both azotemic and non-azotemic CKD were evaluated. Complete staging of CKD according to the recommendations of the International Renal Interest Society (IRIS)31 was not possible in all cats. Urine protein-to-creatinine ratios were known for 3 CKD cats, 2 of which were proteinuric. Systolic blood pressure (SBP) measurements were noted in the medical records of 2 cats with CKD, both of which were normotensive. The most recent SBP measurements for each of these cats were 104 and 118 mm Hg, determined using Doppler sphygmanometry 0 and 4 days before euthanasia, respectively. Both SBP and urine protein-to-creatinine ratio were known for only 1 cat with CKD, with disease that was IRIS stage 4 with proteinuric and normotensive sub-staging.

For cats with CKD, histologic diagnoses made by 1 of 2 board-certified veterinary pathologists included typical ischemic CKD (ie, chronic tubular atrophy and tubulorhexis with interstitial inflammation, lipid, and fibrosis; n = 6), ischemic CKD with oxalosis (n = 2), amyloidosis (n = 1), membranoproliferative glomerulonephritis (n = 1), and possible renal maldevelopment with secondary ischemic changes (n = 1).
### Table 2: Clinical and laboratory data from cats with chronic kidney disease (n = 11) and from healthy control cats (n = 8)

|                                | Chronic kidney disease group | Healthy control group | P-value |
|--------------------------------|-----------------------------|-----------------------|---------|
| Number                         | 11                          | 8                     |         |
| Age (years)                    | 17.3 (3.1-19.4)             | 1.7 (1–7)             | <.001   |
| Sex                            |                             |                       | .007    |
| Female spayed                  | 6                           | 0                     |         |
| Female intact                  | 0                           | 3                     |         |
| Male neutered                  | 5                           | 4                     |         |
| Male intact                    | 0                           | 1                     |         |
| Breed                          |                             |                       | .23     |
| Domestic short haired          | 7                           | 8                     |         |
| Domestic medium haired         | 3                           | 0                     |         |
| Abyssinian                     | 1                           | 0                     |         |
| Body weight (kg)               | 3.65 ± 1.45                 | 4.22 ± 0.95           | .35     |
| Serum creatinine concentration (mg/dL) | 5.4 ± 3.2 | 0.9 ± 0.5 | <.001 |
| Serum urea nitrogen concentration (mg/dL) | 89.0 ± 42.6 | 24.9 ± 5.1 | <.001 |
| Serum symmetric dimethylarginine concentration (μg/dL) | 14.5 (9-20) | 10.5 (6-14) | .60 |
| Urine specific gravity         | 1.016 ± 0.003               | 1.051 ± 0.007         | <.001   |
| Urine protein-to-creatinine ratio | 0.70 (0.39-23.41)  | 0.19 (0.07-0.41) | .02 |
| Time between collection of samples for laboratory data and renal tissues (d) | 17 (0-525) | 0 (0-0) | .001 |

**Note:** Numerical data are presented as mean ± SD or median (range), where appropriate. Number of cats for which data are available is provided if different from the number of individuals in each group.

### 3.2 | Dietary sodium intake

Information regarding dietary sodium intake was available for 6 of 11 cats in the CKD group and 4 of 8 cats in the healthy control group. All cats in the control group for which diet information was known had been fed a maintenance dry cat food (Purina Cat Chow) containing 165 mg sodium/100 kcal. Of the 6 cats in the CKD group for which diet information was known, 5 cats were being fed a prescription renal diet (Hill’s k/d [56 mg sodium/100 kcal] in n = 4 and Royal Canin Feline Renal Support D [90 mg sodium/100 kcal] in n = 1). One CKD cat was being fed a combination of dry and canned forms of a maintenance diet (Purina Pro Plan Focus urinary tract formula [60 and 90 mg sodium/100 kcal for dry and canned forms, respectively]).

### 3.3 | Intrarenal RAS gene transcription

The geometric mean transcript level of AGT was 15.4 (95% CI, 6.3, 37.8) times greater in renal tissue samples from the CKD group compared to those from the control group, whereas the geometric mean transcript level of ACE was 5.3 (95% CI, 2.6, 11.0) times greater in renal tissue samples from the control group compared to those from the CKD group (P < .001; Figure 1). No significant differences were found in renal transcript levels of REN (P = .72) or AGTR1 (P = .51) between groups.

### 3.4 | Renal angiotensin peptides

The geometric mean concentration of Ang I (Ang 1-10) was significantly higher in kidneys from the CKD group (1052 fmol/g; 95% CI, 621, 1783) compared to those from the control group (167 fmol/g; 95% CI, 69, 406; P = .001; Figure 2). Although the geometric mean concentration of Ang II (Ang 1-8) was numerically higher in CKD kidneys (618 fmol/g; 95% CI, 341, 1122) than in control kidneys (319 fmol/g; 95% CI, 155, 656), this difference was not statistically significant (P = .12). No significant difference was found in geometric mean concentration of Ang III (Ang 2-8) between tissues of the CKD group (69 fmol/g; 95% CI, 39, 121) and those of the control group (112 fmol/g; 95% CI, 56, 226; P = .23). Renal concentrations of Ang IV (Ang 3-8) were <4 fmol/g for 10 (91%) of the 11 cats with CKD and 5 (63%) of the 8 healthy control cats; these data are not presented graphically. The proportion of cats with Ang IV concentrations below the lower limit of quantification did not differ significantly...
FIGURE 1  Dot plots of normalized transcript levels of (A) angiotensinogen (AGT), (B) renin (REN), (C) angiotensin-converting enzyme (ACE), and (D) angiotensin II type 1 receptor (AGTR1) in renal tissue homogenates from cats with naturally occurring chronic kidney disease (CKD; n = 11) and healthy control cats (n = 8). Values from the left kidney (open diamond) and right kidney (open circle) of the same individual are connected by solid lines. Levels of each target gene were normalized to those of the reference genes GAPDH and RPS7. For each gene, transcript levels are scaled to those of the lowest sample; note, therefore, that scales differ among genes. ***P < .001

FIGURE 2  Dot plots of renal concentrations of (A) angiotensin (Ang) I (Ang 1–10), (B) angiotensin II (Ang 1–8), and (C) angiotensin III (Ang 2–8) in renal tissue homogenates from cats with chronic kidney disease (CKD; n = 11) and control cats (n = 8). For the CKD group cats, values from the left (open diamond) and right kidney (open circle) of the same individual are connected by solid lines. **P < .01
In our study, kidneys from aged cats with naturally occurring CKD had higher AGT and lower ACE transcript levels, and higher mean Ang I concentrations, than those from healthy, adult cats. Although mean concentration of Ang II was numerically higher in the kidneys of CKD cats compared to those of healthy controls, this difference was not statistically significant. These findings suggest, as in other species, that there may be activation of the intrarenal RAS in cats with naturally occurring CKD.

Inappropriate and deleterious RAAS activation is well documented in CKD in mammals and is a known factor for disease progression. In cats, the systemic RAAS has been shown to be inappropriately activated in some, but not all, CKD models and in a subset of individuals with naturally occurring disease. Recently, local, tissue RASs have gained attention as crucial mediators of end-organ damage. Importantly, these local systems appear to be subject to regulatory mechanisms that are independent of the classical, systemic RASs. Renal concentrations of Ang II may exceed those from healthy kidneys, that the intrarenal RAS might be activated in cats with CKD.

Angiotensinogen is the only known precursor of the angiotensin peptides and its regulation may be of key importance to the activity of the intrarenal RAS. Circulating AGT is produced constitutively by the liver and is relatively abundant, with renin activity representing the rate-limiting, major regulatory step of the systemic RAS. Conversely, within the kidney, generation of Ang II is controlled by numerous independent mechanisms, of which intrarenal generation of AGT appears to be a critical factor. One important distinction between the systemic and the intrarenal RAS is that stimulation of renal Ang II type 1 receptor by Ang II exerts a positive feedforward loop, in part by increasing transcription of AGT. Our data suggest, based on the relative abundance of AGT mRNA in diseased as compared to healthy kidneys, that the intrarenal RAS might be activated in cats with CKD.

Our study identified lower ACE mRNA levels in tissues from affected cats as compared to those from healthy individuals. Additionally, although Ang I levels were significantly higher in cats with CKD, Ang II levels were not significantly different between groups. Although this finding could be a reflection of lower relative expression of ACE, work in other species suggests that ACE-independent pathways of Ang II production predominate in renal tissues. One such pathway relies on the activity of the enzyme chymase, which was not evaluated in our study.

Although the profibrotic and proinflammatory effects of Ang II are primarily mediated by the Ang II type 1 receptor, binding of the Ang II type 2 receptor leads to numerous counterbalancing effects (e.g., systemic and renal vasodilatation, decreased renal sodium reabsorption, nitric oxide release, and inhibition of proliferation, growth and fibrosis). Angiotensin II also is degraded by various enzymes to several smaller, biologically active, angiotensin peptides, including Ang III (Ang 2-8) and Ang IV (Ang 3-8), both studied here, as well as others (e.g., Ang 1-9, Ang 1-7, Ang 1-5) not evaluated in our study. Like Ang II, Ang III can stimulate both Ang II type 1 and type 2 receptors, but has higher selectivity for the Ang II type 2 receptor than does Ang II. Angiotensin IV may be generated from Ang II or III and interacts with the Ang II type 4 receptor, which has functions associated with cognition (e.g., memory acquisition and recall), regulation of blood flow, inhibition of renal tubular sodium reabsorption, and cardiac hypertrophy. We were unable to identify differences in transcript abundance of Ang II type I receptor or concentrations of Ang III and Ang IV between tissues from cats with CKD and those from healthy cats.

Cats in the CKD group were significantly older and more likely to be sexually altered than those in the control group. In human beings, there exist both sex and age differences in the expression levels of RAS components, with young children having higher supine PRA and plasma aldosterone concentrations than adolescents. Similarly, a veterinary study documented an association between aging and a decrease in PRA in healthy cats. In human beings, differences in circulating, urinary or both renin and aldosterone concentrations among men, pre- and post-menopausal women, and women receiving estrogen replacement, have been documented in multiple studies, although evidence occasionally is conflicting with regard to the directionality of the observed disparities. In cats, neutering has been associated with lower PRA, but no change in plasma aldosterone concentration. Significant differences in age and neuter status between the groups in our study represent potential competing factors that may have influenced our results.

Because the RAAS plays a key role in the regulation of fluid and electrolyte balance, its activity is necessarily influenced by dietary sodium intake. In health, an inverse relationship exists between sodium intake and circulating RAAS activity, but this relationship can be disrupted in certain individuals and specific disease states. In healthy dogs, progressive increases in sodium intake have been associated with decreases in PRA, and plasma Ang II and aldosterone concentrations. In another study, supplementation of dietary sodium chloride led to decreased aldosterone concentrations in cats with induced renal insufficiency, but not in healthy control cats. Interestingly, although feeding diets rich in sodium tends to suppress circulating markers of RAAS, high sodium intake appears to induce intrarenal RAS activation in some rodent models. Diet information was available for approximately half of the cats in our study. Of these, all CKD cats were fed a relatively sodium-restricted diet, whereas control
cats were fed a high-sodium diet. How dietary sodium intake differentially impacts the systemic and intrarenal RAS in cats is presently unknown. However, it is plausible that differences in sodium intake between groups might have impacted our results.

Our study had some limitations. First, a relatively small number of banked samples was available for analysis. This, along with a high degree of inter-individual variability, likely limited our power to detect differences between groups. In addition, expressions of renin, AGT, ACE, and the Ang II type 1 receptor were evaluated at the transcriptional level only—concentrations of the corresponding proteins may differ from those of the mRNA transcripts. For financial reasons, equilibrium analyses of angiotensin peptides from only 1 kidney in the control group was possible, which might have further increased the likelihood of type II error. Furthermore, concurrent evaluation of the systemic RAAS was not performed, leaving an important gap in the assessment of this system. Other limitations include lack of complete information regarding systemic arterial BP and dietary sodium intake, which may have impacted our observations if systematic differences existed between groups. Future studies may include the evaluation of a larger number of individuals for which SBP and dietary data are known. For financial reasons, we did not assess components of the so-called alternative angiotensin peptide pathways (eg, ACE 2, Ang 1-9, Ang 1-7, Ang 1-5), or ACE-independent pathways of Ang II production, which could have benefited the interpretation of our data. Finally, because homogenized tissues were used, assessment of regional differences in the expression of the evaluated intrarenal RAS components was not possible.

Presently, assessment of the intrarenal RAS in cats is expensive and requires renal tissue sampling, limiting its usefulness in clinical cases. Studies in humans suggest that urinary AGT is a suitable biomarker of intrarenal Ang II and is associated with increased risk of progressive renal dysfunction in patients with CKD.35,36 Evaluation of urinary AGT as an antemortem, non-invasive marker of intrarenal RAS activation in cats warrants further investigation.

In conclusion, our data suggest that the intrarenal RAS might be stimulated in cats with naturally occurring CKD. Future efforts to identify biomarkers reflecting the status of the intrarenal RAS are crucial for in vivo research in cats.

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CONFLICT OF INTEREST DECLARATION
Dr. Bianca Lourenço was the recipient of a Boehringer Ingelheim Postdoctoral Scholarship. No other authors have a conflict of interest.

OFF-LABEL ANTIMICROBIAL DECLARATION
Authors declare no off-label use of antimicrobials.

INSTITUTIONAL ANIMAL CARE AND USE COMMITTEE (IACUC) OR OTHER APPROVAL DECLARATION
This study was performed using banked samples. All activities related to the original study for which samples were obtained were approved by the University of Georgia IACUC (Animal Use Protocol A2017 05-008-Y3-A1).

HUMAN ETHICS APPROVAL DECLARATION
Authors declare human ethics approval was not needed for this study.

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