Molly, a 2-year-old, 40-lb (18-kg) spayed springer spaniel living in the United Kingdom, was presented for a 4-day history of progressive, multifocal, ulcerative skin lesions and a 24-hour history of lethargy, anorexia, and vomiting. The skin lesions initially affected only the right pelvic limb; however, the day before presentation, additional ulcers developed over both inguinal areas, the lips, and the nasal planum. Molly was up-to-date on vaccinations and received monthly imidacloprid and moxidectin topical parasite prevention.

**Physical Examination**
Mucous membranes were slightly dry, and clinical dehydration was assessed at approximately 6%. Examination showed interdigital ulceration and soft tissue swelling of digits IV and V on the right pelvic limb (Figure 1), superficial erosion of the skin in the inguinal area, and multiple erosions and ulcers over the rostral aspect of the upper and lower lips and nasal planum (Figure 2, next page). No other abnormalities were noted.

**Diagnostics**
Systolic blood pressure measured by Doppler ultrasonography was persistently increased (195 mm Hg; range, 100-150 mm Hg). CBC showed thrombocytopenia (platelet count, $50 \times 10^3/\mu L$; range, $150-500 \times 10^3/\mu L$), and serum chemistry profile revealed hypoalbuminemia, hyperbilirubinemia, and azotemia. International Renal Interest Society (IRIS) grade III acute kidney injury (AKI) was suspected (Table, next page). Basal cortisol concentration (4.4 µg/dL; normal, >2 µg/dL) was within normal limits, making glucocorticoid-
deficient hypoadrenocorticism unlikely. PCR for *Leptospira* spp in a single blood sample was negative. Antinuclear antibody test results were not suggestive of systemic lupus erythematosus.

Urine specific gravity was 1.025; urinalysis was otherwise unremarkable. Urine protein: creatinine ratio was elevated at 2.5 (normal, <0.4). Urine culture was negative. A urinary catheter was placed to monitor urinary output and adjust fluid therapy accordingly as part of standard management of AKI.

No significant findings were identified on abdominal ultrasonography or radiography of the thorax, abdomen, and limbs. Skin biopsy samples were not obtained because of concerns regarding the effects of sedation on renal perfusion.

**Presumptive Diagnosis**
Skin lesions and clinical abnormalities were suggestive of cutaneous and renal glomerular vasculopathy (CRGV). Differential diagnoses included leptospirosis, pyelonephritis, intoxication (eg, grape, ethylene glycol), immune-complex glomerulonephritis, and previous or current renal hypoperfusion (eg, prerenal azotemia), although these conditions are not typically associated with ulcerative skin lesions. A concurrent or associated dermatologic condition (eg, immune-mediated vasculitis, chemical burn) could not be excluded but was considered less likely.

**Treatment**
Standard AKI and supportive treatments were initiated (see *Treatment at a Glance*), including amlodipine (0.1 mg/kg PO q24h initially, then increased to 0.1 mg/kg PO q12h in the absence of clinical response), maropitant (1 mg/kg SC q24h), omeprazole (1 mg/kg PO q12h), and methadone (0.2 mg/kg IV q4h). Antibiotic therapy (ampicillin [15 mg/kg IV q8h]) was initiated to manage secondary infection of skin lesions, pending PCR for *Leptospira* spp and urinalysis results.

Crystalloid fluid therapy (ie, lactated Ringer’s solu-
tion) was initiated at 12.75 mL/kg/hr to provide maintenance fluid therapy and correct 6% dehydration. Urine output measurement after 6 hours of fluid therapy demonstrated urine production of 0.4 mL/kg/hr (normal, >1 mL/kg/hr). Fluid therapy rate was adjusted to match the dog’s urine output and avoid volume overload. A furosemide bolus (1 mg/kg IV) was administered, and furosemide (0.5 mg/kg/hr CRI) was started; urine output remained <0.5 mL/kg/hr over the next 6 hours.

Outcome
Eighteen hours after admission, serum chemistry profile showed worsening azotemia with a creatinine concentration of 7.7 mg/dL (IRIS Grade IV oliganuric AKI). The patient was obtunded. Several options were discussed with the owner, including use of mannitol and/or fenoldopam to promote diuresis, referral for therapeutic plasma exchange and/or continuous renal replacement therapy, and euthanasia. Therapeutic plasma exchange has been used in humans with conditions similar to CRGV. After consideration, Molly was euthanized. Postmortem examination disclosed renal and cutaneous lesions consistent with thrombotic microangiopathy, including fibrinoid necrosis of the glomerular arterioles with frequent vessels occluded by thrombi and concurrent evidence of tubular necrosis and cutaneous coagulative necrosis with rare intravascular thrombi. The presence of these lesions with Molly’s clinical signs and progression confirmed CRGV.

The Take-Home
CRGV is an uncommon disease of unknown cause and is also referred to as Alabama rot in reference to a North American disease of unknown origin that is characterized by similar histopathologic lesions. CRGV has been reported in greyhounds in North America and a Great Dane in Germany. Until 2012, few (<5) cases had been reported in Europe. Since then, more than 150 cases in the United Kingdom, including one in Northern Ireland, have been confirmed by postmortem examination in nongreyhound breeds; 30 of these cases were reported in 2018, making CRGV a rare condition. Although a definitive cause has not yet been identified, research investigating environmental and genetic causes (mainly complement system dysfunction) is ongoing.

Ulcerated, vasculitis-like skin lesions of the extremities, ventrum, face, and tongue are typically reported first. Some dogs develop skin lesions and azotemia; others develop skin lesions only and do not become azotemic, although the percentage of dogs that develop skin lesions only is

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TREATMENT AT A GLANCE
- Fluid therapy (typically over 4-6 hours to allow accurate assessment of urine output) should aim first to correct suspected hypovolemia (rare) and/or dehydration. Once the fluid deficit has been corrected, urine output should be monitored, and the fluid rate should be adjusted to match urine production and avoid volume overload.
- Systemic hypertension should be managed with amlodipine (or hydralazine if immediate control is necessary). ACE inhibitors may reduce glomerular filtration rate and are contraindicated in these cases.
- CRGV skin lesions tend to heal spontaneously over days to weeks with appropriate management, including broad-spectrum antibiotic therapy, if indicated, and appropriate wound management.

AKI = acute kidney injury
CRGV = cutaneous and renal glomerular vasculopathy
IRIS = International Renal Interest Society
unknown due to the lack of gold standard testing antemortem and the lack of specificity of the skin lesions. The average time from onset of skin lesions to development of azotemia is 3 days (range, 1-10 days). Once AKI is present, prognosis is poor, with a mortality rate of 85%. Dogs that do not develop azotemia have an excellent prognosis, provided that wounds are adequately managed with standard local care and systemic antibiotic therapy. Thus far, there have been no reports of CRGV recurrence in a dog that survived the disease or of CRGV transmission between dogs. CRGV does not appear to be zoonotic, although standard precautions should be followed (eg, wearing gloves and a protective apron) until more is known about the disease etiology.

References