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First report of Lyme borreliosis leading to cardiac bradydysrhythmia in two cats

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Abstract

Case series summary Two cats were presented for investigation of bradyarrhythmia detected by their referring veterinarians during routine examination. Both cats had extensive investigations, including haematology, serum biochemistry with electrolytes and thyroxine concentrations, systolic blood pressure measurement, echocardiography, electrocardiography and infectious disease testing. Infectious disease testing included serology for Toxoplasma gondii, Ehrlichia canis, Anaplasma phagocytophilum and Borrelia burgdorferi, and PCR for B burgdorferi antigen in both cats. Case 1 was also assessed by PCR for Bartonella henselae antigen and case 2 was assessed for Dirofilaria immitis by serology. All infectious disease tests, other than for B burgdorferi, were negative. Case 1 was diagnosed with Lyme carditis based on marked bradydysrhythmia, positive B burgdorferi serology, a structurally normal heart and clinical resolution with appropriate treatment with a 4-year follow-up. Case 2 was diagnosed with Lyme carditis based on marked bradydysrhythmia and positive B burgdorferi PCR; however, this cat had structural heart disease that did not resolve with treatment.

Relevance and novel information This small case series describes two B burgdorferi positive cats presenting with newly diagnosed cardiac abnormalities consistent with those found in humans and dogs with Lyme carditis. Both cats were asymptomatic as perceived by their owners; the arrhythmia was detected by their veterinarians.

Keywords: Borrelia burgdorferi; Lyme disease; Lyme carditis; bradyarrhythmia

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Introduction

Lyme borreliosis is caused by a group of Gram negative spirochete bacteria in the Borrelia burgdorferi sensu lato complex. Pathogens in this group are an important cause of morbidity and mortality in humans, with a wide geographical distribution.1–4 Dogs and cats are frequently seropositive, but clinical disease is rare and most commonly reported in dogs.5

Transmission of B burgdorferi requires Ixodes species ticks.6,7 Ixodes ticks live for 2 years, with a three-stage life cycle, where they feed on a variety of different-sized animals, giving them ample opportunity to be infected with and transmit Borrelia organisms.8

In the UK, ticks on cats are usually Ixodes species, predominantly I ricinus (57%) and I hexagonus (41%), with
1.8% of all ticks found to harbour B burgdorferi, although in certain areas the prevalence of infection in ticks is as high as 67%.7,9 These findings are echoed worldwide, with geographical variation in seroprevalence.10–12

Clinical disease in cats with Lyme borreliosis is poorly characterised. Experimentally infected cats have most commonly been asymptomatic.13–15 To our knowledge, there has previously only been one case series of clinical Lyme borreliosis in cats in the USA, and one report of suspected clinical Lyme borreliosis in two cats in the UK.16,17 The former publication reported positive serology for B burgdorferi in cats with clinical signs, including lethargy, lameness, anorexia and hindlimb ataxia, with response to treatment with doxycycline.16 The report from the UK describes recurrent pyrexia in two cats that were identified as PCR-positive for the organism, although treatment and outcome are not reported.17

Lyme carditis is an uncommon clinical manifestation of Lyme borreliosis in people, occurring in approximately 1–10% of cases, depending on geographical location.18,19 The hallmark of Lyme carditis in people is bradydysrhythmia (most commonly second- or third-degree atrioventricular block) and, less commonly, perimyocarditis.18 In dogs, reports of suspected Lyme carditis are rare and the most common presentations are sudden death due to myocarditis, and by dilated cardiomyopathy or cardiomyopathy secondary to infectious disease in humans (Lyme carditis in particular) and cats, we pursued further infectious disease testing.23–28 In-house blood testing for feline immunodeficiency virus and FeLV (SNAP FIV/FeLV Combo Test; IDEXX) was negative. Serology for Toxoplasma gondii (IgG and IgM antibodies; Biobest Laboratories) was negative. Serology for B burgdorferi was positive at 5.5 (RI <4.5 [IgG ELISA; LVPD, University of Liverpool]), with a test that has been validated for use in cats.29 Whole-blood PCR

**Case series description**

**Case 1**

A 7-year-old male neutered Maine Coon cat was presented to the Royal (Dick) School of Veterinary Studies (RDSVS), for assessment of a newly detected arrhythmia. The cat had outdoor access and was fully vaccinated (against feline calicivirus [FCV], feline herpes virus [FeHV], feline panleucopenia virus [FPV] and feline leukaemia virus [FeLV]). The cat was fed a commercial dry food. It had been more than 1 year since any parasite treatment. The arrhythmia was identified during a routine physical examination. There was no significant previous medical history, other than chronic osteoarthritis (OA) of the hips. The owners reported a circular erythematous lesion resembling a tick bite target lesion of about 1 cm diameter on the cat’s ventral abdomen 10 months previously, following a camping trip with the owners in the Scottish Highlands.

Physical examination revealed the cat to be bright, alert and responsive (body weight 6.97 kg; body condition score [BCS] 6/9). Respiratory rate and character were normal (32 breaths per min), oral mucus membranes were pink and moist, and capillary refill time (CRT) was <2 s. Heart rate was variable, ranging between bradycardia at 60 beats per min (bpm) and 140 bpm; no murmurs were evident. Heart rhythm was irregular, with multiple suspected ectopic beats audible; pulse strength was strong, but pulse deficits were present. Rectal temperature was normal. Both hips had reduced mobility, with some discomfort (consistent with chronic OA of the hip), but no other joints were painful or swollen, and the remainder of the physical examination was unremarkable.

Electrocardiography (ECG) was performed using a standard six-lead technique, revealing periods of sinus rhythm at 160 bpm, interspersed with periods of brady-dysrhythmia with ventricular bigeminy and ventricular ectopic beats. There were multiple rhythm abnormalities — including triplets (relatively malignant as R on T in the central beat), singular ectopy, ventricular ectopy from different foci, periods of sinus, and periods of bigeminy and trigeminy.

Blood pressure was normal (130 mmHg, Doppler method). Atropine (0.04 mg/kg IV) resulted in a sustained sinus rhythm for 30 mins, followed by a sustained idioventricular rhythm.

Echocardiography (ECHO) revealed no myocardial changes and no gross structural disease. There was a mild decrease in systolic function based on a decreased fractional shortening (27%, reference >30%) and borderline left ventricular internal diameter in systole (14.1 mm, reference interval [RI] 6.1–14.1 mm).

A 24 h ECG (Holter monitor) was fitted and revealed marked dysrhythmia, including a third-degree atrioventricular (AV) block, plus multifocal ventricular ectopy occurring both singly and in triplets, with periods of bigeminy and trigeminy (Figures 1 and 2). Sustained sinus rhythm occurred only occasionally with a heart rate of 140–160 bpm.

Routine haematology and serum biochemistry, including total thyroxine concentration and electrolytes, revealed no abnormalities. Serum cardiac troponin I (Beaufort) was within the reference interval (RI) (0.02; RI <0.03), as was taurine concentration (IDEXX). Urine analysis, including culture and urine protein:creatinine ratio, was unremarkable.

Our differentials for the cardiac arrhythmias were primary cardiomypathy or cardiomypathy secondary to systemic disease. Owing to sporadic reports of cardiomypathy owing to infectious disease in humans (Lyme carditis in particular) and cats, we pursued further infectious disease testing.23–28 In-house blood testing for feline immunodeficiency virus and FeLV (SNAP FIV/FeLV Combo Test; IDEXX) was negative. Serology for Toxoplasma gondii (IgG and IgM antibodies; Biobest Laboratories) was negative. Serology for B burgdorferi was positive at 5.5 (RI <4.5 [IgG ELISA; LVPD, University of Liverpool]), with a test that has been validated for use in cats.29 Whole-blood PCR
tests for *Borrelia*, *Bartonella* and *Ehrlichia/Anaplasma* species (Acarus) were negative for all three groups of pathogens. Seropositivity in isolation is not diagnostic for clinical Lyme disease,12 and, as such, the presumptive diagnosis of Lyme carditis was made based on positive serology of *B burgdorferi* in conjunction with compatible clinical signs. The cat was treated with doxycycline (Ronaxan [Boehringer Ingelheim Animal Health UK], 10 mg/kg q24h PO) for 30 days. A home visit a month later by one of the authors (DGM) revealed a heart rate of 120 bpm, no obvious dysrhythmia and regular matched pulses. The cat remained clinically asymptomatic, and was re-examined 1 year later; ECHO and ECG revealed a structurally normal heart with a heart rate of 140–160 bpm, and no dysrhythmia. At 17 months after initial presentation, *B burgdorferi* serology was repeated and found to be unchanged from previously (5; RI <4.5). Regular re-examination until euthanasia owing to a thymoma 2.5 years later consistently found a heart rate of 160–200 bpm, with a regular rhythm and good strength matching pulses. Post-mortem examination revealed a grossly and histologically normal heart, with no signs of remaining Lyme disease-associated interstitial myocarditis. This case demonstrates clinical resolution of bradydysrhythmia believed to have been caused by *B burgdorferi* infection, following treatment of doxycycline; the bradydysrhythmia did not return in the 4 years after treatment.

**Case 2**

A 15-year-old male neutered Siamese/Burmese cross cat was presented to the RDSVS, for assessment of brady-cardia and erratic heart rhythm auscultated during a routine evaluation. The cat lived indoors, but had outdoor access when the owners visited their holiday home on the west coast of Scotland, last visiting 4 months previously. The cat was up to date with vaccinations (FCV, FeHV, FPV, FeLV) and treated for endoparasites. No treatments for ectoparasites were given. The cat lived with its biological sibling, a male on whom a tick had been found during their last visit to the same location; the area is endemic for ticks and deer were regularly seen in the garden.

On presentation, the cat was reported to be well, apart from mild weight loss. Physical examination revealed a quiet, alert and responsive cat (weight 3.67 kg, BCS 3/9). Oral mucous membranes were pink, with a normal CRT. Heart rate was 88 bpm, with a regular rhythm, no murmur and matching femoral pulses of good strength. Respiratory rate was 36 breaths per min; effort, pattern and auscultation were normal. The rest of the clinical examination was normal; there were no signs of joint swelling or discomfort, and rectal temperature was normal. Systolic blood pressure was normal (140 mmHg; Doppler method).

ECG revealed third-degree AV block with a ventricular escape rate of 88–90 bpm and an atrial rate of 120 bpm. Ventricular premature complexes of three different morphologies were interspersed within the ventricular escape rhythm, with a maximum rate of 180 bpm. No sinus rhythm was evident.
ECHO revealed moderate/severe right ventricular dilation, left ventricular dilation with normal contractility, normal wall thickness, and prominent left and right atria consistent with an unclassified cardiomyopathy. Mild aortic, pulmonic, tricuspid and mitral insufficiencies were evident and were considered likely to be secondary to chamber dilation.

Further investigations were undertaken to determine a potential cause of the cardiac dysfunction. Haematology revealed mild lymphopenia, while serum biochemistry revealed mild increases in alanine aminotransferase (152 U/l; RI 6–83 U/l), urea (15 mmol/l; RI 2.8–9.8 mmol/l) and creatinine (187 µmol/l; RI 40–177 µmol/l) concentrations; total thyroxine concentration was unremarkable. Urine protein was not evaluated. Serum cardiac troponin I concentration was slightly increased (0.08 ng/ml; RI < 0.03 ng/ml). The reasons for pursuing infectious disease testing in case 2 were the same as in case 1. Serology for Anaplasma phagocytophilum, B burgdorferi, Ehrlichia canis and Dirofilaria immitis (IDEXX SNAP 4Dx) and Toxoplasma gondii (IgG and IgM antibodies; Biobest Laboratories) was negative, as was Ehrlichia/Anaplasma species DNA by PCR on whole blood (Acarus). Borrelia species PCR on blood was positive (Acarus).

A diagnosis was made of B burgdorferi infection suspected of causing Lyme carditis, and the cat was treated with doxycycline (Ronaxan [Boehringer Ingelheim Animal Health UK], 10 mg/kg q24h PO) for 3 weeks.

The cat was re-assessed after completing the antibiotic course. The cat was clinically well with a home respiratory rate of 20 breaths per min. In the clinic, the heart rate was 88 bpm, regular, with matched strong pulses. The rest of the physical examination was unremarkable. Systolic blood pressure was normal (130 mmHg). There were no significant changes from previously on ECG and ECHO.

Other possible causes of the arrhythmia were assessed by abdominal ultrasound and thoracic radiography, neither of which revealed any abnormalities. Serology for B burgdorferi (4; RI < 4.5; LVPD, University of Liverpool), and Borrelia species PCR on blood (Acarus) were both now negative.

The cat was assessed after 3 months and 10 months, and was still asymptomatic. The heart rate, ECG and ECHO findings were largely unchanged from the previous examination. Haematology and serum biochemistry, revealed chronic kidney disease (CKD) IRIS stage 3 (creatinine 255 µmol/l [RI 40–177 µmol/l]; urea 17.1 mmol/l [RI 2.8–9.8 mmol/l]); total thyroxine concentration was unremarkable. Urine protein quantification was not evaluated.

Fourteen months later, aged 17 years, the cat presented with tachypnoea and was diagnosed with congestive heart failure. ECHO revealed severe dilatation of the right-sided chambers, especially the right atrium. ECG revealed a third-degree AV block. The ECG and ECHO findings suggested advanced cardiomyopathy, with a phenotype suggestive of arrhythmogenic right ventricular cardiomyopathy.

Despite treatment, congestive heart failure progressed and the cat was euthanased.

Discussion

We present the first two feline cases of Lyme carditis, which include full case details, treatment and outcomes; case 1 is also the first case of feline Lyme carditis with clinical resolution on treatment.

Prior to diagnosis, the cats had been in the Scottish Highlands, a high-risk area for B burgdorferi in people. Neither of the cats had other clinical abnormalities that could be attributed to B burgdorferi, although case 2 was not evaluated for proteinuria, which would have been pertinent in relation to a potential concurrent Lyme nephritis (and as a part of a standard investigation of CKD). Both cats had third-degree AV block, which is the most common finding in humans with Lyme carditis.

Case 1 had a history of erythema migrans, positive B burgdorferi serology and a bradydysrhythmia that resolved upon treatment. While it could be argued that there may have been another pathogen present that responded to treatment with doxycycline, this seems unlikely considering the cat was clinically well and tested negative for Bartonella, Anaplasma, Ehrlichia and Toxoplasma species. Lyme carditis has been reported in dogs; however, there is no report of complete clinical resolution after treatment in dogs.

In case 2, it is possible that while the cat was PCR positive for B burgdorferi, the cardiomyopathy and arrhythmia may not have been caused by this infection. However, the cat did have a third-degree AV block, which is the most common abnormality in human Lyme carditis, with dilatation of all cardiac chambers, which is the second most common abnormality in canine Lyme carditis. In addition, as with all reported cases of canine Lyme carditis, this cat’s cardiac pathology did not resolve with appropriate treatment. This cat had not seroconverted a month after diagnoses; however, it can take more than 7 weeks for cats to develop seropositivity, and in some cats antibodies are only detectable for 1 week. This is in contrast to dogs, where seroconversion typically occurs 4–6 weeks postinfection and lasts for years. PCR positivity is rare with B burgdorferi as bacterial burden is low and the pathogen sequesters to tissue, which is also the reason why sensitivity is higher in body tissues than in body fluids.

Of note, both cases had a normal and a slightly elevated troponin I. It may be expected that this would be more increased in myocarditis; however, human cases of Lyme carditis have also been reported where the troponin was within normal limits or only mildly elevated.
There were a number of limitations to our study, particularly pertaining to case 2, including the lack of urine protein quantification, lack of a post-mortem examination and lack of response to treatment, although the latter is common in dogs with Lyme carditis.

It is likely that further cases of Lyme carditis may emerge in cats, as most practitioners currently do not test for *B. burgdorferi* in cats with cardiac abnormalities, especially as the cats may be otherwise asymptomatic and many cats do not see a veterinarian regularly. In humans and dogs, Lyme carditis is reported to cause sudden death, which may also be the case in cats and could explain under recognition.\textsuperscript{19,22}

**Conclusions**

Lyme carditis can affect cats and based on this small case series, a positive response to treatment with clinical resolution is achievable. Further investigations are needed to fully elucidate the nature of Lyme borreliosis in cats, but given the effect of climate change with globally increasing prevalence of ticks and tick-borne infections, we suspect further cases will emerge.\textsuperscript{36,37}

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**Ethical approval**

This work involved the use of non-experimental animal(s) only (owned or unowned), and followed established internationally recognised high standards (‘best practice’) of individual veterinary clinical patient care. Ethical approval from a committee was not necessarily required.

**Informed consent**

Informed consent (either verbal or written) was obtained from the owner or legal custodian of all animal(s) described in this work for the procedure(s) undertaken. For any animals or humans individually identifiable within this publication, informed consent for their use in the publication (verbal or written) was obtained from the people involved.

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