Fibrocartilaginous embolic encephalopathy of the cerebellum and brainstem in a cat

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Abstract
A 12-year old male castrated Siamese cat was evaluated for a one-day history of mild obtundation and decerebellate rigidity. Neurological examination findings were consistent with multifocal disease including lesions within the right cerebellum and right brainstem. Investigations included hematology, biochemistry, urinalysis, and urine culture. A definitive diagnosis was not achieved clinically and differential diagnoses included vascular disease, neoplasia, and infectious disease. About nine hours after admission, the cat had an episode where it became rigid and then limp. The cat became agonal and went into cardiopulmonary arrest and attempts of resuscitation were unsuccessful. Post-mortem examination revealed evidence of multiple fibrocartilaginous emboli (FCE) within the cerebellum and within meningeal vessels on the ventral aspect of the brainstem. This is the first reported case of fibrocartilaginous embolism causing an encephalopathy in a cat. While this is a rare disease, it is an important differential diagnosis that should be considered in cases of acute encephalopathies in cats.

Keywords: Cerebellum, Embolus, Encephalopathy, Feline, Fibrocartilaginous.

Introduction
Fibrocartilaginous embolism (FCE) is typically an acute, non-painful, non-progressive myelopathy that results from an ischemic or hemorrhagic infarction of the spinal cord (Gandini and Cizinauskas, 2003). Histological examination of the material found within the vasculature of the spinal cord is typically identical to the nucleus pulposus (Gandini and Cizinauskas, 2003). The pathophysiology of FCE is much debated and several hypotheses have been proposed, but the exact mechanism is poorly understood. The most common theory is that the material comes from an intervertebral disc.

FCE within the spinal cord is commonly reported in the dog and occasionally in the cat (Cauzinille, 2000; Gandini and Cizinauskas, 2003; De Risio and Platt, 2010). It has also been reported in several other species including humans, horses, turkeys, and pigs (Taylor et al., 1977; Johnson et al., 1988; Fuentealba et al., 1991; Stedman et al., 1998; AbdelRazek et al., 2016). There have been eleven reported cases of FCE causing myelopathies in cats, some confirmed on histopathology and some suspected based on clinical and diagnostic findings. These cases reportedly presented as an acute, lateralizing, non-painful myelopathy and lesions were reported within various regions of the spinal cord (Zaki et al., 1976; Turner et al., 1995; Scott and O’Leary, 1996; Abramson et al., 2002; Coradini et al., 2005; Mikszewski et al., 2006; Barker et al., 2014). FCE causing an encephalopathy has been rarely reported in both human and veterinary medicine and has never been documented in a cat. To date, this is the first case of an FCE causing an encephalopathy in a cat reported.

Case Details
A 12-year-old, 3.8 kg, neutered male Siamese cat was referred to the University of Wisconsin-Madison, Veterinary Medical Teaching Hospital for acute onset of mild obtundation and cerebellar signs. The owner had found the cat in lateral recumbency and vocalizing earlier that day. Prior to presentation the cat had been treated by the primary veterinarian with Lactated-Ringers Solution (LRS) (10ml/kg IV), dexamethasone (0.5mg/kg IV), and clindamycin (1.6mg/kg PO) without any improvement. The cat had a prior history of inflammatory bowel disease and had recently been started on oral prednisone every other day (dose unknown).

On presentation the cat was mildly obtunded, hypothermic (99.6°F), bradycardic (100 bpm), and had tacky mucous membranes. He was non-ambulatory, laterally recumbent, vocalizing, and displaying decerebellate rigidity (extended thoracic limbs, flexed pelvic limbs at the level of the hips) and opisthotonus. Neurologic examination revealed mildly obtundated mentation, reduced menace response in the right eye, reduced palpebral reflex in the right eye, and absent physiologic nystagmus.

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The remainder of cranial nerves were normal. Postural reactions were intermittently absent in the right thoracic and right pelvic limbs. Muscle tone and spinal reflexes were normal and there was no pain elicited on spinal palpation. Other physical examination findings were unremarkable.

Based on findings on neurologic examination, multifocal central nervous system disease including the right cerebellum and right brainstem was suspected. Differential diagnoses for the lesion included vascular disease (hemorrhage, infarct), neoplasia (lymphoma, meningioma, glioma, secondary metastasis), and infectious diseases (fungal, protozoal, viral, bacterial). A complete blood count (CBC) and biochemical profile were performed. Clinicopathological abnormalities included lymphopenia of 824 cells/µL (reference range, 1500 to 7000 cells/µL), hyperproteinemia of 7.9 g/dL (reference range, 6 to 7.8 g/dL), hyperglycemia of 198 mg/dL (reference range 56 to 153 mg/dL), elevated CK of 7718 mg/dL (reference range, 55 to 688 mg/dL), elevated AST of 99 U/L (reference range, 14 to 54 U/L), and elevated total bilirubin of 0.3 mg/dL (reference range, 0 to 0.2 mg/dL). These findings were most consistent with stress and prolonged muscle rigidity. On urinalysis triple phosphate crystals, moderate rods, and many cocci were present. A urine culture was negative.

The cat was placed on intravenous fluid therapy (Lactated Ringers Solution + 20mEq KCl/L at 8ml/hr) and was given a dose of buprenorphine (0.01mg/kg IV once) due to concerns of the cat being painful. About nine hours after admission, the cat had an episode where it became rigid and then limp. The cat became agonal and went into cardiopulmonary arrest. Resuscitation was attempted but was unsuccessful. A postmortem examination was performed. Gross examination revealed chronic, diffuse, mild fibrous visceral adhesions of the gastrointestinal tract and mesentery; mild diffuse pulmonary edema; chronic, multifocal siderofibrotic plaques of the spleen; and a focal nodule within the wall of the mid-urethra. Initial sectioning of the brainstem and cerebellum revealed no gross lesions. Histopathologic examination of the cerebellum revealed diffuse, multifocal, extensive hemorrhages within all layers. Multifocally, small to medium sized blood vessels were occluded by amorphous, glassy, blue-gray emboli that occasionally appeared cellular, reminiscent of chondrocytes (Fig. 1). The material stained positive with Alcian blue, indicative of a cartilaginous matrix (Fig. 2).

Several large meningeal blood vessels, including one on the ventral aspect of the brainstem contained similar emboli. The remainder of the central nervous system was unremarkable. Histopathologic examination of the remainder of the body revealed hepatic, pulmonary, and renal hemorrhages.

The cause of these hemorrhages was unknown though extensive endothelial damage may have triggered disseminated intravascular coagulation. The final histopathological diagnosis was multiple fibrocartilaginous emboli (FCE) with hemorrhage.

**Discussion**

Further diagnostics were not performed in this case due to the rapid decline and ultimate death of the patient. However, diagnostics to consider include MRI and cerebrospinal fluid analysis. These diagnostics are not likely to be sensitive or specific to the diagnosis of FCE and findings are likely to be consistent with other vascular-type events. FCE causing an encephalopathy has been rarely reported in both human and veterinary medicine and has never been documented in a cat. Axlund et al. (2004) reported a case of FCE of the brainstem and midcervical spinal cord in a 4-year old Maltese dog. The dog presented for an acute onset of ataxia and left thoracic limb paralysis. Neurologic examination findings were suggestive of multifocal disease affecting both the left lateral cervical intumescence and caudal brainstem. The dog showed no clinical improvement following treatment with corticosteroids and neurologically declined.
The dog was euthanized and a postmortem examination revealed fibrocartilaginous material within the veins of the brainstem and midcervical spinal cord (Axlund et al., 2004). In addition, a case of FCE causing a multifocal ischemic encephalopathy in a 1-week old, female Suffolk cross Texel lamb has been reported. The lamb presented for generalized tremors but was otherwise neurologically normal. Post-mortem examination revealed infarction to the medulla, cerebellum, and multiple areas of the spinal cord. Histological analysis revealed evidence of material identical to nucleus pulposus within the vessels (Jeffrey and Wells, 1986).

To date, a total of sixty-seven cases of FCE in humans have been reported; forty-one tissue-confirmed and twenty-six clinically suspected. Of the forty-one tissue confirmed cases, seven showed evidence of extension into the medulla oblongata and one showed evidence of extension into the cerebral arteries (Kepes and Reynard, 1973; Toro-Gonzalez et al., 1993; AbdelRazek et al., 2016). This is the first case of an FCE causing an encephalopathy in a cat reported to the author’s knowledge. Vascular events causing encephalopathies are commonly associated with hypertension secondary to comorbidities in cats and the prognosis and treatment depend on the underlying cause (Boudreau, 2018). A recent report of feline ischemic myelopathy and encephalopathy in five cats showed evidence of hyaline degenerative arteriopathy within the ventral spinal and/or basilar arteries. Co-morbidities reported included hypertrophic cardiomyopathy, hyperthyroidism, and chronic renal disease (Rylander et al., 2014). Another case series of eight cats with acute non-progressive neurologic signs showed evidence of ischemic myelopathies and encephalopathies. Co-morbidities were reported in all but one of these cases and included hypertrophic cardiomyopathy, hyperthyroidism, and chronic renal disease (Simpson et al., 2014).

Treatment recommendations for FCE induced encephalopathies would likely align with treatment for any vascular-induced encephalopathy and is mostly supportive care such as fluid therapy and physiotherapy. Prognosis is difficult to state with only one case reported. While this is an extremely rare disease, it is an important differential diagnosis that should be included in cases of acute encephalopathies in cats.

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Conflict of interest
The authors declare that there is no conflict of interest.

References


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