Feline Hypertrophic Osteopathy Associated with Congenital Megaesophagus: Two Case Reports and Literature Review

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ABSTRACT: This report describes two cases of feline hypertrophic osteopathy (HO) associated with congenital megaesophagus (ME). The diagnosis was based upon case history, physical examination, radiography and laboratory investigations. The clinical signs of both cats included; regurgitation, weakness and reluctance to walk since birth. Physical examination revealed painful and thickened long bones, swollen joints, difficult flexion of all four limbs and normal urination, defecation, appetite as well as heart and respiratory rates. The abnormal laboratory findings were thrombocytosis, leukocytosis with lymphocytosis and monocytosis. Survey and contrast thoracic radiography showed generalized esophageal dilatation. Radiography of appendicular skeleton revealed symmetrical and massive subperiosteal new bone formation of the affected long bones and pelvis. In conclusion, radiography of extremities is highly recommended for cats admitted with congenital ME and lameness to investigate the possibility of HO. Moreover, the HO should be listed during the differential diagnosis of thickened limbs in cats.

Keywords: Domestic shorthair cat, Hypertrophic osteopathy, Lameness, Megaesophagus, Periosteal proliferation.

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INTRODUCTION

Congenital megaesophagus (ME) or esophageal hypomotility is a rare disease in cats affecting mostly Siamese cats (Hoenig et al., 1990). Acquired ME can be seen in all cat breeds. It is characterized by a decreased or absent esophageal motility. The most common clinical signs include; poor body condition, weight loss, regurgitation, changes in eating habits and difficulty in swallowing. Respiratory distress, aspiration pneumonia, excessive salivation, halitosis, coughing, nasal discharge and fever are the most common complications of ME (Forbes and Leishman, 1985). Megaesophagus may be congenital, due to ill-developed signaling pathway of the esophagus or, acquired due to numerous causes (Hoenig et al., 1990).

Hypertrophic osteopathy (HO) is a diffuse periosteal proliferative disease particularly affecting the appendicular long bones (Hime et al., 1972). Most of the reported cases of HO are secondary to pulmonary or abdominal neoplasms or inflammatory disease of unknown pathogenesis (Rohr 2003; Johnson and Lenz, 2011).

According to the available literature, several cases of feline HO have been reported over the years (1971-2019) as a secondary to several disorders disease such as pulmonary, digestive and renal lesions as well as kidney, adrenal gland and injection site neoplasms as shown in table (1). Moreover, four reports of idiopathic HO have been recorded (Table 1). The prognosis of HO in cats is poor as most of the recorded cases were died shortly afterwards (Richards, 1977; Gram et al., 1990).

Secondary HO is presented with lameness, long bone pain and symptoms from the primary thoracic or abdominal lesions. The diagnosis of such cases is based upon radiography that detects the primary lesions and hypertrophic changes of long bones (Nafe et al., 1981; Gram et al., 1990; Rohr, 2003). Computed tomography (Johnson and Lenz, 2011); Magnetic Resonance Imaging (Salguero et al., 2015) and histopathology (Grierson et al., 2003; Salguero et al., 2015) have also been used in the diagnosis of HO in cats.

Nutritional and supportive treatment such as suitable fluid therapy and electrolyte correction according to the abnormalities in laboratory test results were recommended for treatment of feline HO (Huang et al., 2010).

This study aims to present two cases of feline congenital megaesophagus associated with HO.

Table 1. The primary lesions and prognosis of the recorded feline hypertrophic osteopathy

<table>
<thead>
<tr>
<th>Primary lesion</th>
<th>No of cases</th>
<th>Outcome</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Pulmonary lesion</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Carr (1971)</td>
</tr>
<tr>
<td>Pulmonary neoplasia</td>
<td>One</td>
<td>Died shortly afterwards</td>
<td>Richards (1977)</td>
</tr>
<tr>
<td>Pulmonary neoplasia</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Roberg (1977)</td>
</tr>
<tr>
<td>Renal papillary adenoma</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Nafe et al. (1981)</td>
</tr>
<tr>
<td>Pulmonary carcinoma</td>
<td>One</td>
<td>Died shortly afterwards</td>
<td>Gram et al. (1990)</td>
</tr>
<tr>
<td>Adrenocortical carcinoma</td>
<td>One</td>
<td>Recovered after 4 months of surgery</td>
<td>Becker et al. (1999)</td>
</tr>
<tr>
<td>Pulmonary sarcoma</td>
<td>One</td>
<td>Recovered after 6 months of surgery</td>
<td>Grierson et al. (2003)</td>
</tr>
<tr>
<td>Pulmonary lesion</td>
<td>One</td>
<td>Died shortly afterwards</td>
<td>Rohr (2003)</td>
</tr>
<tr>
<td>Megaesophagus</td>
<td>One</td>
<td>Recovered after medical therapy</td>
<td>Mills (2010)</td>
</tr>
<tr>
<td>Renal adenoma</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Johnson and Lenz (2011)</td>
</tr>
<tr>
<td>Injection-site sarcoma</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Salgüero et al. (2015)</td>
</tr>
<tr>
<td>Cardiointerventricular septal defect</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>De Sousa et al. (2019)</td>
</tr>
<tr>
<td>Idiopathic</td>
<td>Seven</td>
<td>6 cats recovered after medical therapy and one died shortly afterwards</td>
<td>Huang et al. (2010)</td>
</tr>
<tr>
<td>Idiopathic</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>De Melo et al. (2006)</td>
</tr>
<tr>
<td>Idiopathic</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Ocarino et al. (2006)</td>
</tr>
<tr>
<td>Idiopathic</td>
<td>One</td>
<td>Euthanized at the time of diagnosis</td>
<td>Guizelini et al. (2019)</td>
</tr>
</tbody>
</table>

In addition, HO has been previously reported in canine, equine, bovine, wild animals and human secondary thoracic neoplasms, sarcoma of the urinary bladder, bacterial endocarditis and infection by Spirocerca lupi or Dirofilaria immitis (Brodey 1971; Ndikusere and Hill, 1989; Masegi et al., 1994; Mair et al., 1996, Hassan et al., 2019).
CASE HISTORY OF THE FIRST CAT

A 3-month-old female intact domestic shorthair kitten was admitted to the surgery clinic at Faculty of Veterinary Medicine, Cairo University, Egypt with a history of intermittent regurgitation since birth, stiffness and weakness (Fig 1). The case history revealed that the kitten was an orphan stray animal and fed on cat milk until weaning at the age of 2 months. The appetite remained normal despite the intermittent regurgitation and defecation and urination pattern was unremarkable.

![Figure 1. (a) A 3-month-old female domestic shorthair kitten with megaesophagus and hypertrophic osteopathy showing poor body condition and slight thickening of both forearms. (b) The same cat at the age of 4 years showing marked thickening of the limbs.](image-url)

The kitten had also dental malocclusion, changes in eating habits, stiffness and poor condition at the age of 3 months. A thorough physical examination was unremarkable. Symptomatic and supportive therapeutic measures for presumed gastritis were initially performed in the form of intravenous injections of Dextrose 5% solution as boluses (40 mL daily) and 0.3 mg/kg of metoclopramide (Primperan®, Sanofi Aventis) as an antiemetic drug twice daily for 3 days. Three months after initial presentation, the kitten with poor body condition was readmitted for intermittent regurgitation. Plain lateral thoracic radiography and positive contrast esophagography barium sulfate paste (E-Z- Paste® Barium sulphate esophageal cream (60% w/w), Bracco Diagnostics Inc., USA) dosed as 5–10 mL oral boluses together with the previous case history confirmed severe generalized congenital megaeosophagus (Fig 2). The owner was advised to administer a liquid diet from an elevated position and hold the kitten in upright position after each meal. Over the following months, the general clinical status improved yet occasional regurgitation still occurred.

At the age of one year, the cat was re-admitted for investigation of weakness, thickening of all four limbs, reluctance to walk, stiffness in gait and occasional regurgitation. Palpation of the long bones elicited slight pain and showed hard thickening of the humerus, radius, ulna, femur, tibia and fibula. Flexion of all limbs was difficult, particularly the forelimbs.
Routine complete blood count (CBC) revealed thrombocytosis (915x10³/µL, reference interval [RI]: 300-800 x10³/µL), leukocytosis (25.3x10³/µL, RI: 5.5-19.51x10³/µL) with lymphocytosis (9.5x10³/µL, RI:1.5-7x10³/µL) and monocytcosis(1.3x10³/µL, RI: 0.0-0.9x10³/µL). Radiographic examinations of the appendicular skeleton showed extensive subperiosteal new bone proliferation suggesting HO. Radiographic changes included intense and symmetric formation of new subperiosteal bone through all diaphyses and metaphyses of the affected bones (Fig 3). The new bone formation had uniform opacity and a ‘palisading’ pattern in some areas. The most affected bones were radius, ulna and femur in both limbs. Interestingly, the new bone formation was more clearly seen in fore limbs than hind limbs. Oral treatment with meloxicam 0.1 mg q24h (Metacam; BoehringerIngelheim, Germany) was given for pain control.

The animal’s weight was 2 kg, 2.6 kg, 2.7 kg, 2.2 kg and 2.5 kg at the age of 6, 12, 18, 30 and 48 months, respectively. After puberty, the cat exhibited abnormal and scarce estrus cycle that lasted for 2-3 days without any obvious signs. CBC was carried out...
again at the age of 2.5 and 3 years and revealed the same previous abnormal findings.

Serum biochemistry tests including albumin, creatinine, urea, alkaline phosphatase, alanine aminotransferase, aspartate aminotransferase and γ-glutamyltransferase tests revealed high urea level, 49 mg/dL (reference range 10-30 mg/dL) as the only abnormality. Serum creatinine was 0.9 mg/dL (reference range <1.6 mg/dL). Urinalysis was unremarkable with inactive sediment. Until this writing (Four years after the initial presentation), in periodic follow up examinations, the cat remains in stable condition, presenting occasional regurgitation episodes.

CASE HISTORY OF THE SECOND CAT
A one-year-old female intact domestic shorthair cat was admitted with a history of intermittent regurgitation, thickened limbs, weakness and reluctance to walk. Physical examination was unremarkable. Palpation of the limbs showed hard thickening of all four limbs with slight pain. Plain lateral thoracic radiography and positive contrast esophagography (E-Z-Paste® Barium sulphate esophageal cream (60% w/w), Bracco Diagnostics Inc., USA) dosed as 5–10 mL oral boluses revealed dilated esophagus. Radiographic examination of all limbs showed periosteal proliferation in humeri, femurs and tibias (Fig 4). It was not known how long periosteal proliferation had been present prior to initial presentation. Feline hypertrophic osteopathy associated with congenital megaesophagus was confirmed. The owner was advised to feed the cat 5-6 times daily small amounts of liquid diets (such as; beef/chicken broth, baby cereals, blend soft cat food with broth or commercial feline liquid diets) from a high position that resulted in marked improvement in the regurgitation episodes.

The owners of both cases refused any surgical treatment (Percutaneous endoscopic gastrostomy tube placement or esophagectomy) for megaesophagus and we could not follow up the second case.

Figure 4. (a) Latero-medial radiographs of a one-year-old female domestic shorthair cat showing marked periosteal proliferation in humerus, radius and ulna in the forelimb as well as femur and tibia in hindlimb (b).

DISCUSSION
Hypertrophic osteopathy also known as Marie’s disease is a rare disease of uncertain pathogenesis in cats (Langley-Hobbs, 2012). Several suggestions have been mentioned such as periosteal irritation by circulating toxins originated from the primary disease, disturbance in peripheral blood flow to the limbs, neurogenic stimuli, vagal and intercostal nerves stimulations, distal vascular production of platelet-derived growth factor and vascular endothelial growth factor (Richards, 1977; Gram et al., 1990; Salguero et al., 2015). In our recorded cases, the pathogenesis of HO development due to ME was not determined; however, one or more of the aforementioned theories could have contributed. The increase of blood flow to the limbs is one of the most important factors responsible for development of bone lesions. According to the neuron theory, ME may stimulate the vagus nerve...
and increase the peripheral blood flow with less tissue oxygen stimulating periosteal proliferation (Roberg, 1977; de Melo et al., 2006; Huang et al., 2010). In addition, we couldn’t exclude the possibility of simultaneous independent occurrence of idiopathic HO and congenital ME in the recorded cases. Both idiopathic HO and HO secondary to ME were previously reported in cats as shown in table (1). We recommend further studies to identify the exact association between HO and ME in cats.

The recorded cases in this report were young adult female cats. No age, breed or sex predisposition has been reported in literature so far (Huang et al., 2010).

The weakness, poor body condition and reluctance to walk were attributed to deficient nutrition as a result of the intermittent regurgitation. Moreover, the pain, difficult flexion, enlarged joints and hard thickening of all four limbs could be attributed to the massive subperiosteal new bone proliferation. Regurgitation is the most common clinical sign noted with ME. Frequency and timing of regurgitation after feeding may vary considerably. The hematological abnormalities presented by the cats were compatible with those reported in the literature (Ocarino et al., 2006; Salgúero et al., 2015).

Medical treatment of ME includes small, frequent meals from an upright position in order to facilitate passage of the food through the esophagus into the stomach. The decrease in the frequency of regurgitation in our recorded cases could be attributed to the improvement of esophageal function with age due to maturation of the esophageal neuromuscular system after the first year of age. Similar findings were recorded in dogs (Brodey, 1971).

Resolution of the HO after surgical removal of the primary lesion has been reported in several studies (Becker et al., 1999; Grierson et al., 2003; Mills, 2010). Although medical treatment of ME decreased regurgitation episodes in both recorded cases, it did not improve HO. This may be due to severity of both cases and the absence of complete resolution of ME in both cats.

**CONFLICT OF INTEREST**

The authors declare no conflict of interests regarding publishing of this article.

**REFERENCES**


