An unusual case of pheochromocytoma and unruptured abdominal aortic aneurism in a male Yorkshire terrier

Ongebruikelijk voorkomen van een feochromocytoom en niet-geruptureerd aneurysma van de abdominale aorta bij een mannelijke yorkshireterriër


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ABSTRACT

A six-year-old, male Yorkshire terrier was presented with acute vomiting, anorexia, depression, watery diarrhea and sudden blindness. On the basis of a transabdominal ultrasonographic examination, the presence of a prominent aortic aneurysm was established. The aneurysm of the aorta was confirmed at post-mortem examination. Unexpectedly, a pheochromocytoma of the left adrenal gland was found to be involved with the aneurysm. In this case report, the unusual occurrence of a large, unruptured abdominal aortic aneurism (AAA) concurrent with a pheochromocytoma in a male Yorkshire terrier dog is discussed.

SAMENVATTING

Een zes jaar oude, mannelijke yorkshireterriër werd aangeboden met acuut braken, anorexia, depressie, waterige diarree en plotse blindheid. Op basis van een transabdominaal uitgevoerd echografisch onderzoek werd een prominent aneurysma van de abdominale aorta vastgesteld. Bij post-mortemonderzoek werden de aanwezigheid en de locatie van het aneurysma bevestigd. Bovendien werd een feochromocytoom van de linkerbijnier vastgesteld dat betrokken was bij het aneurysma. In deze casuïstiek wordt het uitzonderlijk voorkomen van een groot, niet-geruptureerd aneurysma van de abdominale aorta (AAA) samen met een feochromocytoom van de bijnier bij een mannelijke yorkshireterriër beschreven.

INTRODUCTION

Ante-mortem diagnosis of abdominal aortic aneurysms (AAAs) in dogs has been rarely reported in the literature. Currently, little is known about the events leading to the development of AAAs in the canine species. The term “aneurysm” is derived from the Greek ανευρυσµα (aneurism), meaning “widening”, and is defined as the permanent and irreversible localized dilation of a vessel (Sakalihasan et al., 2005). Large-vessel aneurysm is a disorder of the tunica media of the arterial wall, characterized by destruction of the extracellular matrix components associated with remodelling of the vessel wall in an aneurysmal fashion (Michel, 1998). The usual pathophysiology of an aortic aneurysm involves disruption of the aortic intima and media, along with the formation of a hematoma (Waldrop et al., 2003). Hemodynamic stress facilitates the atheromatous process and impairs the mechanical properties of the abdominal aorta (Michel, 1998).
Many factors contribute to the pathogenesis of AAAs. Local inflammation, apoptosis and oxidative stress, which activate proteases and degrade the aortic wall, play a role (Rizas et al., 2009). This structural instability leads to weakening of the aortic wall and to aortic aneurysm. Additionally, hypertension has consistently been shown to be a major risk factor for AAAs in humans (Vardulaki et al., 2000) and in mice (Kanematsu et al., 2010). Pheochromocytomas can cause episodic or continuous systemic hypertension in 43% of dogs (Barthez et al., 1997; Loste et al., 2013). These tumors are uncommon in dogs (prevalence of 0.17-0.76%) (Myers, 1997).

Canine cases of AAAs have been associated with systemic fungal disease (Gershenson et al., 2011; Murata et al., 2015), spirocercosis (van der Merwe et al., 2008; Kirberger et al., 2013), and familial inheritance in Leonberger dogs (Chetboul et al., 2003). The coexistence of an AAA with a pheochromocytoma has been previously reported as uncommon in humans (Kota et al., 2013). The aim of this case report is to discuss the unusual occurrence of a concurrent large unruptured abdominal aortic aneurism (AAA) and a pheochromocytoma in a male Yorkshire terrier dog.

CASE REPORT

A six-year-old, male Yorkshire terrier presented with acute vomiting, anorexia, depression, watery diarrhea and sudden blindness. Physical examination revealed that the dog was very dehydrated and had pale mucous membranes. The rectal temperature was within normal limits, and the body condition score was 2/5. The skin of the ventral abdomen showed petechiae and multifocal ecchymosis. The hind limbs had mild subcutaneous edema. Routine (red and white) blood cell count was within normal limits for the species. Increased alkaline phosphatase (ALP) activity (583 IU/L; reference interval 20-156 IU/L) and increased urea (73.5 mmol/L; reference interval 10.7-28.6 mmol/L) were observed on the serum chemistry panel. Additionally, thrombocytopenia (0 /μL; reference interval 200,000-500,000/μL) and platelet aggregation (+++) were identified in the blood sample.

An abdominal ultrasonographic (US) examination was performed using a multiple frequency (5-8 MHz) convex transducer. Imaging results revealed moderate peritoneal effusion, bilaterally enlarged kidneys (dimensions of the left and right kidney were 5.24 cm and 5.21 cm, respectively; the normal reference interval for dogs weighing 0-4 kg is 3.2-3.3 cm) (Finn-Bodner, 1995) with mild left and right pyelectasis and urolithiasis in the bladder and the prostatic urethra. A hypoechoic, irregular nodular lesion (3.08 cm x 3.11 cm) was present in the pancreas. The pancreatic lesion was confirmed as a metastasis with necrosis at histopathology. Multiple enlarged lymph nodes with dimensions varying between 1.13 cm and 2.18 cm were visualized. The ultrasonographic features of the lymph nodes were that they were round in shape, hypoechoic relative to the surrounding fat and showed a homogeneous echotexture. Additionally, US examination revealed a pulsatile anechoic structure, which reflected a focal dilation of the abdominal aorta (4.39 cm x 3.94 cm) at the level of the branching of the left celiac artery (Figure 1A). Within the dilation, the blood flow was turbulent, with an appearance partially resembling a yin-yang sign; this pattern is observed on color Doppler as blood within one side of the aneurysm travelling towards the probe (colored red) and blood on the other side travelling away from the probe (colored blue) (Figure 1B). Additionally, remarkable invasion of the aorta by a hypochoic thrombus was visualized (Figure 1A). Standard transverse and longitudinal views of the aneurysm components by US followed previous descriptions in the literature (Bhatt and Dogra, 2008). On the basis of the US findings, the diagnosis of an aortic aneurysm was established.

The patient’s initial treatment was supportive and consisted of isotonic crystalloids (lactated Ringer’s solution; sodium chloride 0.9% 40 mL/kg IV per day)
administered in an alternately basis, empirical anti-
bacterial treatment (enrofloxacin 2.5% and amoxicillin,
5 mg/kg sid SC and 10 mg/kg bid SC, respectively),
fructose with vitamin B complex (one ampule IV per
day), ondansetron (0.1 mg/kg bid IV) and prednisone
(1 mg/kg IM per day). After two days without
signs of clinical improvement, the dog died. A full
necropsy was then authorized and performed.

Initial post-mortem gross examination identified
free abdominal fluid and extensive abnormal tissue
overlapping the aneurysm’s adjacent structures
(Figures 2 and 3A). On the cut section, the abdomi-
nal aorta showed wall dissection at the level of the
branching of the dilated celiac artery, with clot and
thrombus formation (dissecting aneurysm), as speci-
fied by the pathologist (Figures 3A, 3B, 3C and 4).
The left kidney exhibited subscapular hemorrhage
and had multifocal whitish wedge-shaped areas in the
cortical region (anemic infarction). The renal pelvis
was dilated (slight hydroureter). The left kidney
exhibited multifocal to coalesc-
ing yellow areas, firm nodules that were somewhat
delimited (2.0 to 4.0 cm in diameter) and that, when
cut, stretched to the parenchyma. The suprarenal aorta
had multifocal firm, yellow nodules (0.5 cm in diameter).
The small intestine showed multifocal petechiae on
the serous layer and focal adhesion to the pancreas.
Histopathology revealed marked necrosis in the intesti-
nal submucosa, along with fibrinoid degeneration in the
vessel wall associated with infiltration by degener-
ated neutrophils.

Furthermore, the left adrenal gland was enlarged
(5 x 3 x 2 cm), diffusely reddish, soft and friable on the
cut surface. Histopathology was consistent with pheo-
chromocytoma. The tumor cells had a round to poly-
hedral shape. The cytoplasm was eosinophilic and occa-
ionally had a finely granular appearance. The nu-
clei of these cells were oval to reniform and contained
finely granular chromatin and inconspicuous nucleoli.
Moreover, the cells showed marked anisocytosis and

![Figure 2. Initial post-mortem gross examination: free abdominal fluid (white arrow) (D. Driemeier).](image)

![Figure 3A. Tumor and aneurysm overlapped by abnormal tissue (yellow circle) (B.A. Rodrigues). B. Clot at the original site in the AAA (white arrow) (D. Driemeier). C. Fusiform aneurysm after clot removal (black circle). The suprarenal aorta can be visualized (D. Driemeier).](images)
anisokaryosis, with occasional binucleated cells and rare mitotic figures. Additionally, extensive necrosis, multifocal hemorrhage, inflammatory infiltration of lymphocytes and plasma cells and mild multifocal areas of dystrophic mineralization were observed. Metastatic lesions of the left adrenal tumor were diagnosed by histology in the liver, pancreas, kidneys, spleen, lungs and mesenteric lymph nodes. Additionally, metastatic neoplastic bone marrow invasion was confirmed. A diagnosis of malignant pheochromocytoma causing multiple metastases was established. The coexistence of pheochromocytoma and AAA was unique and was challenging to diagnose.

**DISCUSSION**

The present case reveals the coexistence of an AAA and a large tumor (pheochromocytoma) in the left adrenal gland of a six-year-old, male Yorkshire terrier. The simultaneous occurrence of an AAA with a pheochromocytoma has been previously reported to be uncommon in humans (Kota et al., 2013).

Adrenal tumors are usually slow-growing, and their size is extremely variable, ranging from a few millimetres to more than 10 cm in diameter (Reusch, 2006). In the present case report, the tumor in the left adrenal gland of the patient, which was diagnosed post-mortem by histopathology, was large enough (5 x 3 x 2 cm) to press on the surrounding abdominal structures and invade local blood vessels (Laste et al., 2013).

Clinical signs associated with pheochromocytomas are related to the secretion of catecholamines, which may cause episodic hypertension. Blood pressure measurements are valuable in cases of pheochromocytomas. An oscillating hyper- and hypotensive status endorses the pathological condition (Lindquist et al., 2015). Nevertheless, the failure to identify systemic hypertension does not exclude the presence of this adrenal tumor (Nelson, 2002). Generalized weakness, episodic collapse, tachypnea, panting, tachycardia and cardiac arrhythmias may be listed as clinical features described in dogs (Lindquist et al., 2015).

In the present case, although not proven (measurement of blood pressure would aid in verifying hypertension), the exposure to high catecholamines produced by the tumor could be considered an etiologic factor associating the pheochromocytoma with the AAA, similar to reports in the existing literature on human patients (Ehata et al., 1999). Catecholamine release is known to damage the vascular wall and favors the development of an aneurysm. The shape of the aortic aneurysm in the patient in this case report was fusiform with symmetrical bulges around the circumference of the aorta. The presence of a concurrent thrombus in the aneurysm was assumed to have been instigated by intimal aortic damage. Since histopathology of the aorta was not performed, invasion of the aortic wall by the pheochromocytoma remained unclear.

Based on the dilation observed by US, it was presumed that the aneurysm increased with time, similar to what has been previously described in humans (Battaglia et al., 2010). This dilation occurs in conjunction with thinning and weakening of the aortic wall, which can increase the risk of an arterial rupture due to the high pressures and stresses that the aorta must endure. Interestingly, despite the aneurysm's large dimensions (4.39 cm x 3.94 cm) in this case, it remained unruptured. In a Doppler duplex ultrasonographic study of ten beagles (five males; five females) weighing 17-20 kg, the normal aortic diameter at the level of the bifurcation of the external iliac arteries was reported to be approximately 0.88 ± 12 cm (Miño et al., 2004). In the present report, the aneurysm was diagnosed in a male Yorkshire terrier with a prominent abdominal aortic dilation with at least fivefold the diameter expected for a 20 kg dog; the growth exceeded the expected expansion rate. Siegel et al. (1994) supported the theory that a mural thrombus has a protective effect on an AAA by cushioning the pulsations of the flowing blood. The findings in the present case suggest that the primary factor leading to the coexistence of the AAA and the pheochromocytoma was persistent or episodic exposure to high levels of catecholamines produced by the tumor and the likely concurrent hypertension. Additionally, in this current case, hypertensive episodes might have contributed to retinal bleeding and sudden blindness in the dog.

Moreover, histopathologically diagnosed necrosis in the intestinal submucosa could be attributed to catecholamines secreted by the tumor. Hypercatecholaminemia can inhibit peristalsis and promote intense mesenteric artery vasoconstriction, with subsequent ischemic intestinal necrosis (Manger, 2006). Unfortunately, it was not possible to measure the levels of catecholamine because the patient died 48 hours after presentation of the clinical complications (generalized organ failure).
The laboratory parameters of platelet aggregation and severe thrombocytopenia soon after presentation in the clinic may support a disseminated intravascular coagulation (DIC) state. Large aortic aneurysms, as observed in the patient in this case report, may lead to the local activation of coagulation factors, and cause DIC (Levi, 2004). The syndrome itself is characterized by the systemic activation of coagulation. Microvascular thrombosis compromises organ perfusion and may contribute to organ failure.

Ante-mortem diagnosis of pheochromocytomas is challenging because the clinical signs are often intermittent (Boston, 2014), and laboratory findings are not specific. Affected animals frequently have concurrent diseases (hyperadrenocorticism, diabetes mellitus or neoplastic processes), which may further hinder a clear diagnosis (Herrera et al., 2008). The diagnostic work-up of suspected pheochromocytoma in dogs includes immunohistochemical staining for endocrine cells with chromogranin A and synaptophysin (Rosa et al., 2012). Unfortunately, this technique is only possible after removal of the affected gland. In humans, advanced imaging techniques such as magnetic resonance imaging (MRI) and computer tomography (CT) can reliably identify 95% of pheochromocytomas 1 cm or larger (Manger, 2006). Additionally, CT diagnosis of adrenal tumors in dogs has been reported (Spall et al., 2011). However, availability, the need for anesthesia, costs and the time required for the execution of these techniques limit the ability to perform these procedures. The measurement of the ratio between urinary metanephrine and creatinine levels has been reported in the literature by Héron et al. (1996) as a diagnostic tool in human patients. In a recent study by Salesov et al. (2015) urinary and plasma catecholamines and metanephrines were compared in healthy dogs (n=10), in dogs diagnosed with pheochromocytoma (n=7), hypercortisolism (n=10), and in dogs with nonadrenal diseases (n=14) with the aim to determine the test with the best diagnostic performance for dogs with pheochromocytoma. It was shown that dogs with pheochromocytoma had significantly higher urinary normetanephrine and metanephrine : creatinine ratios and significantly higher plasma-total and free normetanephrine and plasma-free metanephrine concentrations than the three other groups. The authors concluded that measurement of normetanephrine was the preferred biochemical test for pheochromocytoma, and urine was superior to plasma. The results of the study allowed the authors of the present case report to envision the use of the test as a reliable diagnostic tool to detect the disease.

In accordance with the findings of other authors (Spall et al., 2011), in addition to the adjacent vessel invasion (30-50%) (Baily and Page, 2007), the tumor can induce pressure on the surrounding abdominal structures, and metastatic spread occurs. The liver and lungs are common sites of tumor metastasis (Baily and Page, 2007; Boston, 2014), similar to the findings in the present case. Additionally, as in the presented case, other sites of metastasis are regional lymph nodes, the spleen, kidneys, pancreas and bone marrow.

In the case presented here, clinical disease chronicity contributed to the development of multiple histologically confirmed metastases distributed in various organs (liver, pancreas, lungs, lymph nodes, bone marrow and spleen), which in turn culminated in generalized organ failure. Moreover, the pheochromocytoma probably was the underlying disorder causing the AAA. The end-disease pathway resulted in the DIC state and the death of the patient.

In conclusion, transabdominal US with color Doppler served as an effective screening modality for the detection of the AAA. US accurately identified the site and extent of involvement of the aneurysm sac and the presence of a thrombus. However, non-visualization of the adrenal tumor was a major drawback of this examination. Notably, the abnormal tissue and adjacent structures overlapped in the US investigation, thereby precluding visual access to the gland. The necropsy and histopathology findings confirmed the existence of a pheochromocytoma. Aneurysms are still incidentally found during routine veterinary, ultrasonographic examination.

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REFERENCES


