Pericardial effusion and cardiac tamponade in a dog with hypothyroidism

Efusión pericárdica y taponamiento cardíaco en un perro con hipotiroidismo

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RESUMEN

La relación entre hipotiroidismo y enfermedades cardiovasculares en perros frecuentemente está asociada con la presencia de signos clínicos como bradicardia sinusal, alteraciones miocárdicas y complejos QRS de bajo voltaje y onda T invertida en el electrocardiograma. Varios estudios en perros hipotiroides no han mostrado la presencia de efusión pericárdica, sin embargo, en humanos es bien conocida la presencia de efusión pericárdica asociada a hipotiroidismo, pero el taponamiento cardíaco no es un signo frecuente. Un perro Golden Retriever fue presentado con historial clínico de letargia progresiva, intolerancia al ejercicio, intolerancia al frío y dificultad respiratoria durante 3 semanas. Al examen clínico el perro presentó disnea, tonos cardíacos tenues e hipertensión. El examen ecocardiográfico demostró la presencia de fluido en el saco pericárdico y taponamiento cardíaco. Se realizó una pericardiocentesis y el análisis citológico del fluido descartó la presencia de células neoplásicas. La información entregada por los propietarios, además de la detección de zonas alopecias en cola y abdomen sumado a la evolución clínica llevó a analizar las hormonas tiroideas, confirmando el hipotiroidismo. Aunque la etiología de la efusión pericárdica no fue confirmada es posible que el hipotiroidismo haya causado la efusión pericárdica, o bien el perro padeció una efusión pericárdica idiopática y coincidentemente hipotiroidismo. Sin embargo, de acuerdo con los signos clínicos descritos y evolución de este caso, se sugiere que aun cuando la efusión pericárdica es un signo no descrito en el hipotiroidismo canino, se debe considerar la evaluación de las hormonas tiroideas en perros con efusión pericárdica idiopática.

Palabras clave: hipotiroidismo, efusión pericárdica, taponamiento cardíaco, perros.

SUMMARY

The relationship between hypothyroidism and cardiovascular diseases in dogs is frequently associated with clinical signs like sinus bradycardia, myocardial alterations and ECG findings such as low voltage QRS and inverted T wave. Echocardiographic studies in hypothyroid dogs have not identified pericardial effusion. Hypothyroidism in human patients is a well-known cause of pericardial effusion, but cardiac tamponade is not a frequent clinical sign. A Golden Retriever was presented with a clinical history of progressive lethargy, exercise intolerance, cold intolerance and increased respiratory effort for 3 weeks. At the clinical examination the dog presented dyspnea, muffled heart sound and hypotension. Echocardiography showed fluid in the pericardial sac and cardiac tamponade and pericardiocentesis was performed. Analyses of the fluid showed a non-neoplastic etiology. The information given by the owners, the alopecia zones detected in tail and abdomen and the clinical response, led to analyse the thyroid hormones, and hypothyroidism was confirmed. Following two pericardiocenteses and levothyroxine supplementation, the effusion resolved. Although the etiology of pericardial effusion was not clear, it is possible that the pericardial effusion had been caused by hypothyroidism or the dog had idiopathic pericardial effusion and coincidental hypothyroidism. However, according to the clinical signs and the evolution of the dog, an evaluation of thyroid hormones in dogs with pericardial effusion must be considered even though pericardial effusion is a non-described sign in canine hypothyroidism.

Key words: hypothyroidism cardiac tamponade, pericardial effusion, dogs.

INTRODUCTION

Cardiac abnormalities associated to hypothyroidism in dogs are common, but their clinical significance is higher in patients with preexisting cardiac conditions. The relationship between hypothyroidism and cardiovascular diseases in dogs is frequently associated with clinical signs like sinus bradycardia, myocardial alterations and some findings in ECG such as low voltage QRS and inverted T wave. However, echocardiographic studies in hypothyroid dogs do not describe the development of pericardial effusion (PE) (Flood and Hoover 2009, Panciera 1994, Phillips and Harkin 2003, Sangster et al 2012). The presence of PE in humans with hypothyroidism represents between 3-5% and it is dependent on severity and duration of hypothyroidism (Chiang et al 2012).

The purpose of this report is to describe an atypical clinical sign of an adult Golden retriever with hypothyroidism that was diagnosed with pericardial effusion and cardiac tamponade.

MATERIAL AND METHODS

A 5 year old unspayed female Golden Retriever, weighing 37.5 kg, was referred. The dog had presented with a clinical history of weakness, extremely sensitive
to cold temperatures, exercise intolerance and increased respiratory effort for 3 weeks. At time of presentation, the dog was quiet and slowly responsive. A marked inspiratory dyspnea, dry and cyanotic mucous membranes, weak femoral pulse and jugular distension were observed. During thoracic auscultation, marked bronchial and vesicular sounds were heard but the intensity of the heartbeat was decreased. Systemic blood pressure values were 200/140 mmHg. Rectal temperature was 36.5 °C and pulse oximetry was at 76%. The abdomen was distended and tense, indicating the presence of fluid in the abdominal cavity. Also, hyperpigmentation with alopecia on the dorsal neck area, nose and tail was observed. Initial treatment included oxygen supplementation and placement of intravenous catheter to administer medication. The dog was eating and drinking and no hypovolaemic status was detected. An enlarged, globoid shaped cardiac silhouette was observed in thoracic radiographs. Moreover, a diffuse alveolar pattern was observed.

RESULTS AND DISCUSSION

Echocardiographic findings included pericardial fluid and severe collapse of the right ventricle consistent with cardiac tamponade (figure 1). According to the normal values for Golden Retriever (Morrison et al 1992) a decreased left ventricular systolic diameter (2.1 cm) and decreased left ventricular diastolic diameter (3.26 cm) was diagnosed. The shortening fraction was calculated to be approximately 35%. The left atrium:aorta ratio was 1.35. Abdominal ultrasound detected ascites. Findings were conclusive with cardiac tamponade. Pericardiocentesis was performed and 428 ml of hemorrhagic fluid was removed. An ECG was done before and after pericardiocentesis. Before the pericardiocentesis the ECG showed low voltage QRS and ST complexes and slight electrical alternans on QRS an ST. These findings resolved immediately after the pericardiocentesis (figure 2). Pericardial fluid cytology revealed 13,900 cells/uL and hematocrit of 10%. The observed cells were consistent with pleomorphic reactive mesothelial cells with prominent nucleolus and occasional mitotic figures were detected in some microscopic fields. In contrast, abundant neutrophils and thrombocytes were also present. These findings were associated to aseptic fluid characterised by reactive mesothelial cells. However the mesothelioma was not discarded due to the presence of malignancy features in some cells.

A blood test revealed regenerative normocytic normochromic anemia (PCV 27%, reference range 37-50%; MCV 70 fl, reference range 60-77 fl; MCHC 28.9 g/dl, reference range 32.0-37.0 g/dl), and thrombocytopenia (11,2 x10^3/ ml, reference range 177-398 x10^3/ml). Leukocytosis was observed (29,4x10^3/ml reference range 5,3- 19,8 x10^3/ml)

Figure 1. M-mode echocardiography from the right parasternal long axis view. Note the fluid in the pericardial sac before the pericardiocentesis. The diagnosis of cardiac tamponade was made according the collapse of the right ventricular during diastole (arrow).

Eje largo de vista para esternal derecho en modo M ecocardiografico. Se observa fluido en el saco pericárdico previo a la pericardiocentesis. El taponamiento cardiaco fue diagnosticado observando el colapso del ventrículo derecho durante el diástole (flecha).

Figure 2. Electrocardiogram showed a low voltage QRS in all leads and slight electrical alternans on QRS and ST complexes, before the first pericardiocentesis (A). Electrocardiogram after the pericardiocentesis (B), QRS complexes increased the voltages without presence of the electrical alternans. Paper speed 25mm/s; 1cm= 1mV.

El electrocardiograma muestra un bajo voltaje en el complejo QRS en todas las derivadas, además se observa leve alternancia eléctrica en los complejos QRS y ST, antes de realizar la pericardiocentesis (A).
with moderate neutrophilia (25.5 x 10^3/ml, reference range 3.1-14.4 x 10^3/ml). Biochemical analysis revealed high creatine kinase (220 U/L, reference range 25-125 U/L) and mild azotemia (urea nitrogen 55mg/dl, reference range 5-30 mg/dl, creatinine 3.1mg/dl, reference range 0.7-1.8 mg/dl).

Clinical findings, laboratory results and echocardiography confirmed the diagnosis of pericardial effusion. According to the findings, the most suggestive diagnosis was idiopathic pericardial effusion.

After 24 hours the dog showed a marked improvement. Seven days after initial treatment the ascites disappeared but still the dyspnea and arterial hypertension (190/140) persisted. A treatment with 0.5 mg/kg enalapril orally twice a day and oxygen (5L/min) was initiated. The hematological parameters worsened, a severe neutrophilia (neutrophils 49373 10^3/ml reference range 3300-10000 10^3/ml) was detected. Thoracic radiographies showed a persistence of alveolar pattern. Therefore, secondary bacterial contamination of pulmonary tissue was associated with this possibly due to pulmonary congestion or intubation procedure. Antibacterial therapy was initiated using 20mg/kg Amoxicillin/ clavulanic acid orally twice a day. Moreover, azotemia increased (Urea nitrogen 70 mg/dl, reference range 5-30 mg/dl; Creatinine 4.7mg/dl, reference range 0.7-1.8mg/dl) with a marked hyperphosphatemia (8.43 mg/dl, reference range 2.8-6.1 mg/dl) possibly associated to a decreased renal blood flow due to hypertension. Intravenous maintenance fluid was maintained and diuresis therapy with furosemide 2mg/kg orally twice a day was included. On day 15, although blood pressure and body temperature were normal and respiratory signs and renal function improved the dog remained cold intolerance. Radiographs findings showed cardiomegaly with a globoid shape but less than the first radiography approach. Echocardiography showed cardiac tamponade again. A second pericardiocentesis was performed and 196 ml of the reddish fluid was removed containing 2700 cells/µl and hematocrit at 4%. Fluid analysis contained 5.2 g/dl of total protein, pH 7.0 and glucose 150mg/dl. Cytological analysis showed monocytes (95%), neutrophils (5%) and a light red background. At the same time, blood samples were collected that showed that hematological and biochemical parameters were abnormal. Anemia still was present but the leukocytosis was resolved. Cholesterol, glucose, albumin and triglycerides were higher than reference values. Because of this thyroid hormones were measured^1. The results showed decreased values of T4t and T4f (T4t 0.5 ug/dl, reference range 1.3-3.8 ug/dl; T4f 0.2 pg/dl, reference range 0.3-1.3 pg/dl) with increased cTSH (1.0 ng/ml, reference range < 0.6 ng/ml). Based on clinical signs, clinical pathological findings and thyroid hormones, primary hypothyroidism was diagnosed. At day 20, after the first pericardiocentesis, the thoracic radiographs and echocardiography were normal. For this reason the dog was discharged. Oral treatment with Levothyroxine at dosage of 0.02 mg/kg q12h, furosemide (3mg/kg/q12h) and enalapril (0.5mg/kg/q24h) were recommended. Thyroid hormones were normal (T4t 2.1 ug/dl, T4f 0.9 pg/dl, TSH 0.5ng/dl) on day 60 and the cutaneous signs improved totally on day 120. Within 8 months of hormonal supplementation and monthly echocardiographic monitoring, the presence of pericardial effusion, heart tumor or new cardiovascular signs did not develop.

Pericardial diseases (PD) represent nearly 8% of all cardiovascular disorders. Most dogs with PD develop pericardial effusion (PE) as a common clinical sign (Shaw and Rush 2007^2). PE is frequently associated as a consequence of cardiac neoplasias or the idiopathic form representing 58% and 19%, respectively (Stafford et al 2004). PE associated to hypothyroidism is a rare condition in dogs but it has been previously described in humans (Chiang et al 2012, Cohen et al 2012, MacGregor et al 2004). Reports of PD in dogs with hypothyroidism are scarce. Only a single report describes cholesterol-based pericardial effusion in a dog with hypothyroidism, where other different causes could not be ruled out (MacGregor et al 2004).

The clinical signs described in the present report were muffled heart sounds, jugular venous distention, poor pulse quality, ascites, lethargy and exercise intolerance. These signs are frequently reported in dogs with cardiac tamponade (Shaw and Rush 2007^2, Stafford et al 2004).

Cardiac tamponade was diagnosed and considered an emergency and pericardiocentesis was immediately performed. After pericardiocentesis the dog improved its clinical and electrocardiographic parameters but the pericardial effusion recurred albeit to a lesser degree than at presentation. The hypertension did not resolve after the initial pericardiocentesis, so furosemide and enalapril were administered until the hypothyroidism had resolved with levothyroxine supplementation. Until now hypertension had not been reported in hypothyroid dogs as a clinical sign. However in humans many studies have reported a high prevalence of hypertension in hypothyroidism. In the case of cardiac tamponade with hypothyroidism, the hypertension could be explained by a possible excessive adrenergic drive caused by the tamponade (Cohen et al 2012). Also, increased peripheral vascular resistance and low cardiac output have been suggested to be the possible link between hypothyroidism and diastolic hypertension. Also, T3 deficit has been associated with peripheral vasoconstriction and the renin level is reduced in hypothyroid patients (Stabouli et al 2010).

Although the use of furosemide is not recommended as management of cardiac tamponade because it can reduce cardiac filling pressure and thus lower cardiac output (Ware 2009), we justified it use due to persisted hypertension. Moreover, the furosemide treatment was administered while the dog developed the second tamponade, and the severe hypotension described as a potential complication was not detected.

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^1 Immulite canine immunoassay, Siemens Health Care.
After the second pericardiocentesis, with Levothyroxine supplementation, no pericardial fluid was subsequently collected. However, in this report it is not possible to confirm clearly that the pericardial effusion was attributed to hypothyroidism, mainly because golden retrievers have a significantly increased risk for development pericardial effusion and hypothyroidism. Therefore, it is probably that the pericardial effusion was caused by hypothyroidism or the dog had idiopathic pericardial effusion and coincidental hypothyroidism.

Cytological studies described that the cytological features of the recurrent effusion differed during the course of the disease. Therefore, cytological findings must be associated with clinical signs present in the dog (Cagle et al 2014,Stepien et al 2000). However, the diagnostic possibilities using cytology are varied but increase when the pericardial fluid has a hematocrit <10% (Cagle et al 2014, de Lafourcade et al 2007). In humans, pericardial fluid from hypothyroid cardiac tamponade patients is typically an effusion with a color yellowish, with lymphocytes and hematocrit < 10%, although the serosanguinous fluid has been reported. The protein content oscillate between 5.0 and 6.2 mg/dl. Also, in cytological examination the presence of mononuclear cells or reactive mesothelial cells is uncommon. This allows to rule out easily some neoplastic diagnoses.

In our case the cytological findings showed reactive mesothelial cells with the first pericardial fluid removed but the hematocrit was 10%. In the second sample of fluid removed the hematocrit was 4% and mesothelial cells were not detected and the protein was 5.2 mg/dl. Therefore, the second and last effusion was a modified transudate. But, the definitive non neoplastic diagnosis was made using echocardiographic checkups during 8 months ruled out the presence of pericardial disease. Retrospective echocardiographic studies detected that 60% of the dogs with idiopathic pericardial effusion have reactive mesothelial cells without echocardiographic evidence of mesotheliomas during a period at more than 120 days (MacGregor et al 2004, Rajagopalan et al 2013, Stepien et al 2000).

A diagnosis of hypothyroidism in this dog was suspected based on typical clinical signs such as chronic lethargy described by the owners, nasal and tail alopecia, cold intolerance, also non regenerative mild anemia and hypercholesterolemia (Panciera 1994).

The diagnosis was confirmed with T4 and TSH screenings. In humans hypothyroidism is thought to result in increased capillary permeability and impaired lymphatic drainage with protein leakage into the interstitial space, resulting in PE. Furthermore, hypothyroidism causes salt and water retention (Lin et al 2000). For these reasons, hypothyroidism in human patients is a well-known cause of PE, but cardiac tamponade is not a frequent clinical sign. Echocardiography in humans diagnosed early with hypothyroidism, identifies PE in 3-5% cases. The PE is minimal, representing < 5mm of the M mode width and no more than 200 ml of fluid collected (Lin et al 2000). In dogs, the velocity at which the fluid accumulates is more important than the total volume accumulated to cause tamponade. (Cohen et al 2012, Lin et al 2000, Shaw and Rush 2007, Stafford et al 2004). Finally the hypothyroidism may have caused the effusion, but is more likely that idiopathic PE did. Unfortunately, there is no way to prove either of them. However, the thyroid hormone supplementation in this case was not associated to a third pericardial fluid collection (Chiang et al 2012, Cohen et al 2012, MacGregor et al 2004). The evaluation of thyroid hormones in dogs with pericardial effusion must be considered.

REFERENCES