A Hemangiosarcoma Case in a Dog

Gamze KARABAĞLI1*, Oktay DÜZGÜN1, Esma YILDAR1, Özge ERDOĞAN2, Aydin GÜREL2

1İ.Ü. Veteriner Fakültesi Cerrahi Anabilim Dalı, 34320, Avcılar, İSTANBUL
2İ.Ü. Veteriner Fakültesi Patoloji Anabilim Dalı, 34320, Avcılar, İSTANBUL

*Corresponding Author: Gamze KARABAĞLI İstanbul Üniversitesi Veteriner Fakültesi Cerrahi Anabilim Dalı, 34320, Avcılar, İstanbul, e-mail: gamze.vet@gmail.com, Tel: +90 212 4737070-17304

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SUMMARY
The material of this study is a 3.5 years old male Rottweiler dog presented to Istanbul University Veterinary Faculty Surgery Clinics with a complaint of swelling in the right shoulder region. A diagnosis of hematoma was made and treatment was performed, but swelling started to grow gradually and general well-being deteriorated. When the drain was controlled, drainage of fluid containing blood was observed. Anemia and leucocytosis was determined on examination of the blood. The patient was operated. The mass was observed to expand in the thoracal cavity and it was found that organs in that regions were disfunctioned. So the patient was considered to be inoparable and euthanasia was performed. Specimens taken were examined histopathologically by using streptavidin-biotin immunperoxidase (IMP) method and a diagnosis of hemangiosarcoma of soft-tissue origin was made.

Key Words: Hemangiosarcoma, dog, histopathology

ÖZET
BİR KÖPEKE HEMANJİYOSARKOM OLGUSU

Anahtar Kelimeler: Hemanjiosarkom, köpek, histopatoloji

Introduction
Hemangiosarcoma is a malignant tumor with high metastasis rates also known as angiosarcoma or hemangioendothelioma originating from vascular endothelium (Douglas, 2007; Goldschmidt et al. 2002; Hidaka, 2006; Hilbe et al. 2002). Hemangiosarcoma is seen more frequently in
dogs compared to other animal species. It is predominantly reported in dogs of 9-10 years old. However, there are also publications reporting its presence in dogs under the age of three. German Shepherd dogs, Golden Retriever, Bernese Mountain dogs and Boxer dogs are among sensitive breeds (Bar-Am, 2006; Douglas, 2007; Vail, 2000). Hemangiosarcomas in dogs are aggressive and malignant tumors with poor prognosis (Bar-Am, 2006; Clifford, 2000; U’Ren, 2007). They are responsible of 12-21% of all mesenchymal tumors and 5% of all non-cutaneous primary malignant tumors in dogs. 2.3-3.6% of cutaneous tumors and 45-51% of splenic tumors in dogs are hemangiosarcomas. Although the etiology of hemangiosarcoma is unknown, thinning of hair and minimal pigmentation is found in laboratory dogs exposed to ultraviolet light (Brown, 1985; Douglas, 2007; Erdem, 2000; Hargis, 1992; Liapis, 2004; Mellanby, 2004; Ward, 2008). In the present study, the objective is to transfer clinical state and histopathologic examination of a case of hemangiosarcoma originating from soft-tissue found in a 3.5 years old Rottweiler dog to literature.

**Case Report**

Our case is a 3.5 years old, male, Rotweiler dog presented to Istanbul University Veterinary Faculty Surgery Clinics (presentation date: 09-25-2009, Protocol number: 2009003291) with gradually increasing swelling in the right shoulder joint and neck region despite therapy performed.

On examination a fluctuant mass about the size of a soccer ball was observed including the right shoulder and neck region (Figures 1, 2). It was reported that the general welfare of the patient gradually deteriorated and it could hardly walk. It was also reported that treatment of hematoma had been performed on the mass for 4 weeks. When the drain was controlled, drainage of fluid containing blood was observed. Anemia and leucocytosis was determined on examination of the blood. A date for operation was determined for extirpation of the mass or amputation of the extremity. Following anesthesia induction with 6 mg/kg propofol (IV 200 mg/20 ml, Pofol ® Sandoz) shaving and disinfection was completed. For general anesthesia Isoflurane (100 ml, Forane®, Abbott laboratories) inhalation anesthesia was performed. Circumferential incision (melon slice) was performed on the mass and tissues were dissected. The mass was found to be connected to the thorax. It was decided that the mass could not be removed by operation. With the will of the owner euthanasia was performed. The red-brown soft mass with a size of 10 cm x 7 cm x 5 cm and grey areas on the section surface was fixed with 10% formaldehyde, treated with alcohol and xylol series and embedded into paraffin blocks. Sections of 5μ...
thickness were taken and stained with Haematoxyline-Eosine. On microscopic examination of sections it was seen that endothelium-like cells with shapes changing from oval to fusiform and with vesicular nuclei and one or more nucleoli contracted cellular component. Many fissures, irregular capillaries and larger vascular structures, numerous typical and atypical mitoses, large necrosis areas, diffuse tromboses in the vessels and inflammatory cell infiltration were found (Figure 3). A diagnosis of hemangiosarcoma originating from soft-tissue was made and immunohistochemical labeling with the tumor marker Factor VIII (F-VIII) was performed on sections using streptavidin-biotin immunoperoxidase method to confirm the diagnosis. Neoplastic cells thought to be originating from endothelial cells were determined by cytoplasmic F-VIII staining (Figure 4).

Discussion and Conclusion

Hemangiosarcoma is a malignant tumor originating from vascular endothelial tissue (Bar-Am, 2006; Hidaka, 2006). Although hemangiosarcoma is mostly seen in middle-aged animals, publications reporting this tumor in dogs younger than 3 years old are compatible with our case.

Figure 3. A. Trombose in vein (arrow mark). B. Hemangiosarcoma, Hematoxylin-Eosin.
2.3-3.6% of cutaneous tumors in dogs are hemangiosarcoma. In a study performed in 104 dogs, 65 of the cases of hemangiosarcoma were reported to have developed in the spleen (62%), 18 in soft-tissue of the thorax and extremities (17%), 3 in the heart (3%), 3 in the long bones, 2 in the lung (2%), 1 in the bladder, 1 in the aorta and 1 in the prostate (4). Our case is in a large percentage segment (Douglas, 2007).

In dogs, hemangiosarcomas are metastatic tumors and rapidly lead to death (U-Ren, 2007). Metastasis occurs by transabdominal implantation or by hematogeneous route rapidly. Hemangiosarcoma has been found in semimembranous muscle histopathologically. Metastases are most frequently seen in the liver, omentum, mesenterium and lung (Bar-Am, 2006). The fact that the mass covered nearly the whole thoracic cavity in our case suggested that metastasis occurred rapidly by hematogenous route. It is compatible with the literatures.

Clear demonstration of hemangiosarcoma includes hematologic and biochemical laboratory findings, coagulation tests, thoracoabdominal imaging, abdominosynthesis +/- or echography (Douglas, 2007). Hematologic abnormalities have reported to include normocytic-normochromic anemia, thrombocytopenia, hemolysis, hypofibrinogenemia, prolongation of coagulation time (Bar-Am, 2006). The diagnosis of our case was evaluated biochemically-physiologically and histopathologically as far as possible and found to be compatible with the literatures.

Surgery and chemotherapy has limited success for survival (Clifford, 2000). When immunotherapy and chemotherapy is combined, survival has been reported to prolong (U’Ren, 2007). Because of too many metastatic areas were present in our patient, treatment methods reported in the literature could not be performed.

Finally, hemangiosarcoma may occur in any region of the body. We believe that early diagnosis of the pathologic mass with biopsy and subsequently total extirpation of the mass would prolong survival and decrease possible function losses and pain to a minimal level.
REFERENCES


