INTRODUCTION

Hemangiosarcoma (HSA) is a malignant neoplasm derived from precursor vascular endothelial cells. HSA is an uncommon neoplasm in domestic animals, being more common in dogs than in cats. For that reason the information that we have about feline HSA is poor. (Johannes, et al., 2007)

OBJECTIVES

Characterize the biological behavior, treatment outcomes, hematological and biochemical changes, prognostic risk factors and survival times of feline HSA.

RESULTS AND DISCUSSION

- Middle-aged to older animals (ages ranged from 7 to 17 years, mean was 11,42±2,91 and median was 12).
- No breed or sex predilection were observed.
- There are different types of HSA classified according to their location as cutaneous, subcutaneous, visceral, oral cavity and conjunctival. There is no agreement on the frequency of each form.
- There is no agreement about which are the most frequent cutaneous and subcutaneous HSA locations.
- Some studies and our own results support that the most frequent locations in visceral feline HSA are spleen, liver mesentery, small and large intestine (Clup, et al., 2010; Ogilvie and Moore, 2001, Scavelli, et al., 1985).
- Although right atrium is the third most common site of occurrence of HSA in dogs, primary cardiac HSA in the cat has been poorly reported (Merlo, et al., 2002).
- Kraje et al. (1999) reported metastatic disease in animals with cutaneous and subcutaneous HSA, located in lung, lymph nodes, spleen, kidney and eyes. The locations of metastasis reported in our study were spleen, kidney and lung.
- Scavelli, et al. (1985) claims that 60% of the cats suffering from visceral HSA developed metastasis. Our results support that the most frequent metastasis sites in visceral HSA were: liver, small bowel, mesentery and colon.
- The treatment of choice is the complete surgical extirpation of the neoplasm. Benefits of intravenous chemotherapy is unclear.
- Comparisons of median survival time based on age, gender, tumor site, margins and biochemical changes were not statistically significant (Figure I and III). Comparison of median survival times based on the presence of anemia instead of normal red blood cells count was statistically significant (value p <0.05) (Figure II).

CONCLUSIONS

- Most of the claims about feline HSA are controverted because all the information that we have about it comes from retrospective cases studies and published cases. There is discussion on some items. Release a prospective study could be warranted.
- Cutaneous HSA in cats may have had more aggressive and metastatic behavior than previously thought, because some cats in this study have developed recurrence, metastasis and comparing median survival time based on tumor site showed no statistically significant difference.
- The visceral HSA diagnosis usually is delayed. It may be the reason why often those animals suffer from metastasis at the time of diagnosis.
- Anemia is a usual complication of HSA, caused by acute or chronic hemorrhage derived from primary or metastasis rupture and microangiopathic hemolytic disease.
- Most of the biochemical alterations associated to HSA are caused by the tumor location, or by the hypotension derived from hemorrhage caused by primary or metastasis rupture

References:
6. Dr. Elenia Martinez de Emeiros and Diego Esteban. Who helped me to get cases. Thanks especially to Dr. Joseph Pastor, who mentioned this work and has been an inspiration to me.

Figure I: Kaplan-Meier median survival time for cat with hemangiosarcoma.

Figure II: Kaplan-Meier median survival time data comparing cats with normal red cells count (blue) and with anemia (green).

Figure III: Kaplan-Meier median survival time data comparing cats with visceral (green), cutaneous (blue) and other location (yellow) HSA.

Diagnostic test | Results
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Hematology | Regenerative and non regenerative anemia.
Blood smear | Polychromatophilic and hyperchromatic erythrocytes, Howell-Jolly bodies, anisocytosis, macrocytosis, microcytosis, poikilocytosis, schistocytes and erythroblasts.
Neutrophilic leukocytosis, neutrophilia, leukopenia with neutropenia, leucopenia and lymphopenia.
Thrombocytosis and thrombocytopenia.
Biochemistry | Hypoproteinemia, hypalbuminemia, increased alkaline phosphatase (ALT), reduced and increased blood urea nitrogen (BUIN), reduced and increased creatinine. Increased citrate prothrombin time and activated partial thromboplastin time (ct-PT and co-APTT).
Thorax and abdominal radiography | Effect Mass Pulmonary metastasis
Ascites and pulmonary edema
Abdominal ultrasonography | Masses and metastasis
Cytology | Erythrocytes and exceptionally fusioplastic neoplastic cells.
Histology | Blood-filled vascular spaces lined by neoplastic endothelial cells.
Immunohistochemistry | Markers such as factor VIII related antigen allow us to differentiate well differentiated HSA from sarcomas and granulocyti stiocarcinomas.

Table I: Diagnostic test and results to expect in animals with HSA.