



DIFFICULTIES IN MANAGING OF CANINE CUTANEOUS HISTIOCYTOSIS

Introduction

Cutaneous histiocytosis (CH) is a rare dermatology condition in which the histiocytic cells proliferation causes nodular skin lesions⁽¹⁾. We described a clinical case that reflects the difficulties in the management of this disease.

Material and methods (clinical case description)

A 5-years-old intact male mixed breed dog was attended in our dermatology service with a 1-month history of nodular lesions. Multiple solid and non-pruritic nodules were located in face, trunk and limbs (Figures 1 and 2). No other symptoms were founded at clinical examination. Multiple biopsies of the nodular lesions were taken. The histopathologic findings were consistent with a CH. One month of treatment with tetracycline/niacinamide 500mg of each one twice daily and cyclosporine 5mg/kg daily, was not effective. Administration of prednisone 1mg/kg twice daily was able to resolve the skin lesions. After five weeks, adverse side effects occurred and prednisone was reduced at 1mg/kg daily, and azathioprine (2 mg/kg daily) was added. Lesions relapsed in one month and prednisone doses had to be increased. Three months later, dog development serious systemic symptoms and was euthanased.

Results

CH is a disorder of reactive dermal dendritic cells⁽²⁾. A dysregulation of immune response mechanism have been proposed⁽³⁾. Some immunomodulatory/ immunosuppressive drugs have been used in the management of CH. Tetracycline/niacinamide combination has proved to be successful⁽⁴⁾. However, it was not effective in our patient that had to receive immunosuppressive doses of glucocorticoids +/- azathioprine to control the clinical course of lesions. An ongoing immunosuppressive therapy has also been reported to avoid relapses in some dogs⁽⁴⁾, increasing the risk of development adverse side effects, as we saw in our case.

Conclusion

The management of canine CH can be complicated, especially in patients who need a continuously immunosuppressive therapy that can lead to a fatal outcome.

References

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Figure 1.



Figure 2

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