

# Comparative study between Uveodermatologic syndrome in dogs and Vogt-Koyanagi-Harada disease in humans



Javier Lagioiosa

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### INTRODUCTION

Uveodermatologic syndrome (UVD) is a rare disease of the dog characterized by association of clinical signs of uveitis and cutaneous signs of depigmentation (vitiligo and poliosis)

The Vogt-Koyanagi-Harada (VKH) disease is a bilateral panuveitis associated with dermatological alterations, central nervous system and auditory system involvement in humans

Unknown etiology: virus infection??

### **OBJECTIVES**

- Bibliographic review of two similar diseases in different species (dogs & humans)
- Comparative study
- Describe similarities and difference

### **PATHOGENESIS**

- Autoimmune aggression against melanocytes
- Ag of the immune response: Tyrosinase protein (ez melanin synthesis)
- Cellular immune response-> LT
- Genetically susceptible individual

VKH: Altered expression of HLA (human leukocyte antigens)

**UVD**: Altered expression of DLA (dog leukocyte antigens)

# Th1 cytokines Th2 cytokines VKH disease clinical features Uveitis Disacousia Meningism

### **UVD** syndrome

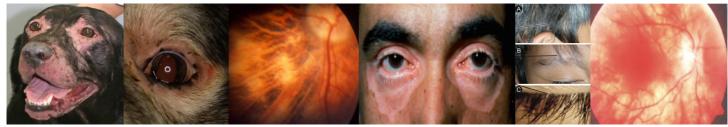
- Most of cases Japanese and northern breeds: Akita (80%), Siberian Husky and Samoyed. Others breeds described
- Age: 6 months and 6 years old
- No apparent sex predilection
- Anterior uveitis or bilateral panuveitis. OF: chorioretinitis + depigmented lesions RPE and choroid
- Vitiligo, poliosis and symmetrical alopecia: periorbital and nasal regions, eyelids and lips
- Neurological and auditory signs exceptional in dogs

### VKH disease

- Darkly pigmented races (Orientals, blacks & Hispanics)- Adults between 20 and 50 years old
- Both genders (women +++)

### 4 stages:

- Prodromal: influenza symptoms + tinnitus
- Acute: bilateral granulomatous panuveitis with RD, neurological disorders and disacusia.
- Chronic: vitiligo, poliosis & alopecia: face, neck, trunk and eyelids. OF: sunset glow fundus
- Chronic recurrent: Episodes of uveitis months or years after the acute stage



### **DIAGNOSIS**

- Clinical Dx: Association of ocular & cutaneous signs / breeds
- Skin biopsy: Lichenoid infiltrate of epidermis: histiocytes, lymphocytes, plasmocytes and giant cells & pigmentary Incontinence.
- Clinical criteria by the American Society of Uveitis in 2001
- 3 types of VKH: I o probable, II o incomplete, III o complete
- Lumbar puncture: pleocytosis in CSF

ASSOCIATION OF OCULAR, CUTANEOUS (NEUROLOGIC AND AUDITIVE IN HUMANS) SIGNS IS PREDICTIVE OF THESE SINDROMES

### TREATMENT

- Systemic and topical corticosteroid: Prednisolone/ Dexamethasone/Triamcinolone (SC, IV)
- Refractory cases and side effects: Immunosuppressive drugs: Cyc A, Azathioprine , Cyclophosphamide, Tacrolimus, Methotrexate
- Long term treatment: 6-8 w in dogs  $\,/\,$  6-12 months VKH type 1 & 2; + 1 year type 3

### **PROGNOSIS**

- Poor prognosis in dogs. Frequent recurrences
- Complications: Cataracts, secondary glaucoma, RD, BLINDNESS
- VKH: Good prognosis 50% of patients
- Cataracts, glaucoma and choroidal neovascular membranes

## **CONCLUSIONS**

- > Great similarities between both diseases (etiology, pathogenesis, clinical signs, diagnosis and treatment)
- Interest of the dog as an animal model for the study of VKH disease and vice versa
- The term UVD is the most appropriate for the dog due to differences with VKH disease
- Early diagnosis in humans and late in dogs
- Importance of early treatment to avoid complications