



Ocular disorders known or presumed to be inherited (published)

| | Diagnosis | Description and comments specific to the breed | Inheritance | Gene/ marker test | References |
|----------|--------------------------------|---|---------------------|-------------------|------------|
| A | Glaucoma | Associated with uveal cysts and goniodysgenesis | Unknown | NO | 1 |
| B | Canine multi-focal retinopathy | (CMR1) | Autosomal recessive | BEST1 | 2 |
| C | Neuronal Ceroid Lipofuscinosis | | Autosomal recessive | CTSD | 3 |

The ECVO's advice relating to hereditary eye disease control

Please see ECVO Manual chapter 8: VET Advice

Recommendations regarding age and frequency for eye examinations

Please see ECVO Manual chapter 7: ECVO Age and Frequency recommendations

Other ocular disorders (reported)

| | Diagnosis | Source |
|----------|---|-------------------------|
| A | Distichiasis | ACVO genetics committee |
| B | Entropion | ACVO genetics committee |
| C | Persistent pupillary membranes – iris to iris | ACVO genetics committee |

References

1. Pumphrey SA, Pizzirani S, Pirie CG & Needle DB. Glaucoma associated with uveal cysts and goniodysgenesis in American bulldogs: a case series. *Veterinary Ophthalmology*, 2013, 16,5, 377-385.
2. Guziewicz KE, Zangerl B, Lindauer SJ, et al. Bestrophin gene mutations cause canine multifocal retinopathy: A novel animal model for best disease. *Investigative Ophthalmol & Visual Science*, 48:1959-1967, 2007.
3. Awano T, Katz ML, O'Brien DP et al. A mutation in the cathepsin D gene (CTSD) in American Bulldog with neuronal ceroid lipofuscinosis. *Mol Genet Metab*, 2006; 87 (4): 341-8.