

# Ocular disorders known or presumed to be inherited (published)

	Diagnosis	Description and comments specific to the breed	Inheritance	Gene/ marker test	References
А	Strabismus/Esotropia		Autosomal recessive	NO	1-3
В	Ectopic cilia		Unknown	NO	4
С	Corneal sequestrae		Unknown	NO	1
D	Apocrine hidrocystomas		Unknown	NO	5
Е	Uveal Cyst		Unknown	NO	6
F	Iris hypoplasia		Unknown	NO	7
G	Microphakia and lens luxation		Unknown	NO	8,9
Н	Lens luxation	Usually occurs at 7-9 years of age	Unknown	NO	10
I	Cataract	Congenital	Unknown	NO	11

J	Congenital glaucoma	Associated with	Unknown	NO	12
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		iridoschisis			
		Open Angle eventually associated with PLA and	Unknown	NO	13-15
		microphakia  Associated with spherophakia and iris hypoplasia	Autosomal recessive	YES	16
K	Retinal degeneration rdAc	Blindness usually occurs at 3-5 years of age	Autosomal recessive	YES	17

## 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 diseases with ocular involvement

	Diagnosis	Source
Α	Gangliosidosis 1 (GM1)	18,19
В	Mucopolisaccaridosis VI (MP6)	20-22

### References

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OTHERS: Temperature dependent oculocutaneous albinism: autosomal recessive. known mutation. (Giebel LB et al. A tyrosinase gene missense mutation in temperature-sensitive type I oculocutaneous albinism. A human homologue to the Siamese cat and the Himalayan mouse. J Clin Invest. 87(3): 1119–1122,1991.).