



Ocular disorders known or presumed to be inherited (published)

	Diagnosis	Description and comments specific to the breed	Inheritance	Gene/ marker test	References
A	Microphthalmia	Associated with other ocular anomalies (cataract, lens luxation, ciliary body dysplasia)	Unknown	NO	1,2,3
B	Retinal dysplasia 1. multifocal 2. geographic 3. total		1. presumed autosomal recessive 2. & 3. autosomal recessive	NO	1,2,3,4,5,6
C	Progressive Retinal Atrophy (PRA)	(CORD1) 1. Cone-Rod Dystrophy; night blindness between 6 months and 2 years of age 2. Late onset; night blindness at 7 years of age	1. Presumed autosomal recessive or polygenic 2. Unknown	1.RPGRIP1	1,2,7,8,9,10, 11,12
D	Glaucoma	In the breed positive association	Unknown	NO	1,2,13,14

		between PLA and glaucoma, between narrowing of the iridocorneal angle and glaucoma and effect of age on the iridocorneal angle			
E	Entropion	Lower eyelid and lateral canthus	Unknown	NO	1,2
F	Corneal dystrophy 1.stromal 2.endothelial	1. 1-5 y.o. 2. >9 y.o	Unknown	NO	1,2
G	Cataract	1. posterior subcapsular, 1-3 y.o. 2. Fibrillar nuclear cataract; >5 y.o.	Unknown	NO	1,2

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	Persistent pupillary membranes All forms	ACVO genetics committee

C	Persistent hyaloid artery	ACVO genetics committee
D	Vitreous degeneration	ACVO genetics committee

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