



Ocular disorders known or presumed to be inherited (published)

	Diagnosis	Description and comments specific to the breed	Inheritance	Gene/ marker test	References
A	Microphthalmia	Often associated with other ocular malformations, uni- or bilateral (all varieties)	Unknown	NO	1
B	Distichiasis	Common (all varieties)	Unknown	NO	1
C	Lacrimal punctum atresia		Unknown	NO	1
D	Corneal dystrophy - epithelial	Clinically affected about 1 year of age (toy, miniature)	Presumed autosomal recessive	NO	1
	-epithelial	6 to 10 y.o; dogs (miniature, medium)	Unknown	NO	21
E	Persistent pupillary membranes	Miniature, medium, giant, standard	Unknown	NO	1, 21, 22
F	Glaucoma	Narrow angle often associated with pectinate ligament abnormality. (miniature, toy, medium)	Unknown	NO	1,2,17

G	Cataract	Cortical/nuclear (all varieties)	Unknown	NO	1,8,16, 21, 23
H	Progressive Retinal Atrophy (P RA)	Early (from 3 y.o. dogs) (Toy, Miniature, and Standard)	Autosomal recessive	prcd	1,4,5,6,10- 14
I	Progressive Retinal Atrophy (PRA)(rcd4)	Rcd4-PRA Late onset; clinical signs from 10-12 y.o. (Miniature, Medium and Standard)	Autosomal recessive	C2orf71	18
J	Cone degeneration: Day Blindness/Retina l Degeneration (achromatopsia)	Day blindness in puppies which evol ves over years in retinal degeneration (Standard Poodle)	Autosomal recessive	DB/RD test	19
K	Optic nerve hypoplasia/ micropapilla	Blindness and abnormal pupil response (miniature, toy)	Unknown	NO	1,7,9,15,20

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	Trichiasis	ACVO genetics committee French National Panel
B	Vitreous degeneration	ACVO genetics committee French National Panel

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