



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

	<b>Diagnosis</b>	<b>Description and comments specific to the breed</b>	<b>Inheritance</b>	<b>Gene/ marker test</b>	<b>References</b>
<b>A</b>	Microphthalmia	Associated with multiple ocular anomalies; uni or bilateral	Unknown	NO	1, 2
<b>B</b>	Distichiasis		Suspected autosomal dominant	NO	2
<b>C</b>	Ectopic cilia	Responsible for corneal ulcers	Unknown	NO	2
<b>D</b>	Trichiasis	Nasal fold and caruncula; responsible for corneal medial pigmentation	Polygenic	NO	1, 2
<b>E</b>	Pigmentary keratitis	Exposure. Corneal ulcers with corneal thickening observed	Polygenic	NO	1, 2
<b>F</b>	Kerato-conjunctivitis sicca	Uni or bi-lateral, dogs > 5 y.o.	Unknown	NO	3, 4

<b>G</b>	Chronic superficial keratitis	2-5 y.o. dogs, bilateral, corneal lateral/inferior superficial	Unknown	NO	2
<b>H</b>	Corneal dystrophy-stromal	9m. - 2 y.o.dogs, bilateral, subepithelial,microcrystalline	Unknown	NO	2
<b>I</b>	Iris atrophy	Stromaldogs > 8 y.o.	Unknown	NO	2
<b>J</b>	Glaucoma		Unknown	SRBD1	5
<b>K</b>	Cataract -subcapsular -nuclear	Posterior cortical along suture lines or triangular polar, 1,5-3 y.o. dogs  Peripheral posterior	Unknown	NO	1
<b>L</b>	Persistent hyperplastic tunica vasculosa lentis/Persistent hyperplastic primary vitreous (PHTVL/PHPV)	Grades 1 and 2 observed	Unknown	NO	2
<b>M</b>	Retinal detachment	From 2 y.o., bilateral, secondary glaucoma possible	Unknown	NO	1, 2
<b>N</b>	Progressive Retinal Atrophy (PRA)	Night blindness in 2-5 y.o. dogs, slowly progressive	Suspected autosomal recessive	NO	1, 2
<b>O</b>	Retinal dysplasia	Folds or geographical	Unknown	NO	2

<b>P</b>	Micropapilla/Optic nerve hypoplasia		Unknown	NO	1
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### **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)**

	<b>Diagnosis</b>	<b>Source</b>
<b>A</b>	Euryblepharon	ACVO Genetics Committee
<b>B</b>	Entropion, lower medial	ACVO Genetics Committee
<b>C</b>	Persistent pupillary membranes	ACVO Genetics Committee
<b>D</b>	Vitreous degeneration	ACVO Genetics Committee
<b>E</b>	Persistent hyaloid artery	ACVO Genetics Committee

### **References**

1. Rubin LF (1989) Inherited eye diseases in the purebred dogs. Williams & Wilkins, Baltimore, 266-268.
2. Chaudieu G, Chahory S (2013) Affections oculaires héréditaires ou à prédisposition héréditaire chez le chien. Rueil-Malmaison : Ed. du Point Vétérinaire, Wolters Kluwer France, 438-41.
3. Sanchez RF et al (2007) Canine keratoconjunctivitis sicca: disease trends in a review. J Small Anim Pract 48:211-217.
4. Kaswan RL, Salisbury MA (1990) A new perspective on Canine keratoconjunctivitis sicca. Treatment with ophthalmic Cyclosporine. Vet Clin North Amer Small Anim Pract 20 : 583-613.
5. Kanemaki N, Tchedre KT, Imayasu M, et al. Dogs and humans share a common susceptibility gene SRBD1 for glaucoma risk. PloSone. 2013;8:e74372.