Cairn Terrier	

Ocular disorders known or presumed to be inherited (published)

	Diagnosis	Description and comments specific to the breed	Inheritance	Gene/ marker test	References
Α	Ocular melanosis with and without glaucoma	Between 7 -13 y.o.; slow progression; proliferation of melanocytes within the uveal tract associated with an elevation in intraocular pressure.	Autosomal dominant	NO	1,2,3,4,5,6,7
В	Persistent pupillary membranes - iris to iris		Unknown	NO	2
С	Cataract	1. posterior cortical; young dogs 2. anterior cortical wedgeshaped; dogs between 6 -8 y.o.	Unknown	NO	1,2
D	Lens luxation	Between 4 -7 y.o.	Unknown	NO	1,2

E	Progressive Retinal Atrophy (PRA)	Young dogs; <1 y.o.	Unknown	NO	1,2
F	Retinal dysplasia -multifocal		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
Α	Distichiasis	French national panel
В	Keratoconjunctivitis sicca	French national panel
С	Persistent hyaloid artery	ACVO genetics committee
D	Vitreous degeneration	ACVO genetics committee

References

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- 5. Petersen-Jones SM, et al: Ocular melanosis in the Cairn Terrier: clinical description and investigation of mode of inheritance. Vet Ophthalmol 10:63-69, 2007.
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- 7. Van de Sandt RR, Boeve MH, Stades FC, Kik MJ: Veterinary Ophthalmology 6: 273-278, 2003.