

<b>Cardigan Welsh Corgi</b>	
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**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>	Lens luxation		Unknown	NO	1,2,3,4
<b>B</b>	Progressive Retinal Atrophy (PRA)	Rod-cone dysplasia type 3 (rcd3); night blindness at 3 months, complete blindness by 1 y.o.	Autosomal recessive	PDE6A	1,2,5,6,7,8,9

**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>	Distichiasis	ACVO genetics committee
<b>B</b>	Corneal dystrophy - epithelial	French national panel

<b>C</b>	Persistent pupillary membranes	ACVO genetics committee
<b>D</b>	Cataract	ACVO genetics committee
<b>E</b>	RPED	ACVO genetics committee
<b>F</b>	Retinal dysplasia	ACVO genetics committee

### References

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