# Mastiff

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

|   | Diagnosis                                     | Description and comments specific to the breed  | Inheritance         | Gene/<br>marker test | References |
|---|---|---|---------------------|----------------------|------------|
| A | Entropion                                     |   | Unknown             | NO                   | 2          |
| В | Ectropion                                     | Often located in central portion of upper and lower eyelid in combination with macroblepharon and medial and lateral entropion. | Unknown             | NO                   | 1,2        |
| С | Distichiasis                                  |   | Unknown             | NO                   | 2          |
| D | Macroblepharon                                |   | Unknown             | NO                   | 2          |
| E | Persistent<br>pupillary<br>membranes<br>(PPM) |   | Unknown             | NO                   | 1,2        |
| F | Retinal dysplasia - folds                     |   | Unknown             | NO                   | 2          |
| G | Multifocal retinopathy                        | CMR1  | Autosomal recessive | BEST1 (VMD2)         | 2,3        |

| н | Progressive<br>Retinal Atrophy<br>(Dominant<br>PRA)   | Onset, progression and severity are modulated by light exposure. | Autosomal<br>dominant | Rhodopsin<br>(RHO) | 2,4,5,6,7 |
|---|---|--|-----------------------|--------------------|-----------|
| i | Prolapsed gland of the nictitating membrane           |  | Unknown               | NO                 | 2         |
| J | Eversion of the cartilage of the nictitating membrane |  | Unknown               | NO                 | 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                  |
|---|---------------------|-------------------------|
|   | Corneal dystrophy   | ACVO genetics committee |
| Α | -epithelial/stromal |                         |
|   | -endothelial        |                         |
| В | Cataract            | ACVO genetics committee |
| С | Uveal cysts         | ACVO genetics committee |

### **References**

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- 4. Kijas JW, Cideciyan, AV, Alemansd TS, et al. Naturally occurring rhodopsin mutation in the dog causes retinal dysfunction and degeneration mimicking human dominant retinitis pigmentosa. Proc Natl Acad Sci U S A, 2002; 99: 6328-6323.
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- 6. Cideciyan AV, Jacobson, SG, Aleman TS, et al. In vivo dynamics of retinal injury and repair in the rhodopsin mutant dog model of human retinitis pigmentosa, Proc Natl Acad Sci U S A, 2005; 102: 5233-5238
- 7. Miyadera K, Acland GM and Aguirre GD. Genetic and phenotypic variations of inherited retinal diseases in dogs: the power of within- and across-breed studies. Mamm Genome. 2012 Feb;23:40-61.