

DNA Test Report

Health Conditions Known in This Breed

| Genetic Condition | Gene | Risk Variant | Copies | Result |
|--|--------------|--------------|--------|--------|
| Collie Eye Anomaly (CEA) | NHEJ1 | Deletion | 0 | Clear |
| Dental Hypomineralization | FAM20C | C>T | 0 | Clear |
| Early Adult Onset Deafness For Border Collies only (Linkage test) | Intergenic | Insertion | 0 | Clear |
| Hereditary Calcium Oxalate Urolithiasis, Type 1 | Confidential | — | 0 | Clear |
| Intestinal Cobalamin Malabsorption (Discovered in the Border Collie) | CUBN | Deletion | 0 | Clear |
| MDR1 Medication Sensitivity | MDR1/ABCB1 | Deletion | 0 | Clear |
| Neuronal Ceroid Lipofuscinosis 5 (Discovered in the Border Collie) | CLN5 | C>T | 0 | Clear |
| Sensory Neuropathy | FAM134B | Insertion | 0 | Clear |
| Trapped Neutrophil Syndrome | VPS13B | Deletion | 0 | Clear |

Other Conditions Tested

| Genetic Condition | Gene | Risk Variant | Copies | Result |
|--|--------------|--------------|--------|--------|
| 2,8-dihydroxyadenine (DHA) Urolithiasis | APRT | G>A | 0 | Clear |
| Acral Mutilation Syndrome | GDNF | C>T | 0 | Clear |
| Acute Respiratory Distress Syndrome | ANLN | C>T | 0 | Clear |
| Alaskan Husky Encephalopathy | SLC19A3 | G>A | 0 | Clear |
| Alexander Disease | GFAP | G>A | 0 | Clear |
| Amelogenesis Imperfecta (Discovered in the Italian Greyhound) | ENAM | Deletion | 0 | Clear |
| Amelogenesis Imperfecta (Discovered in the Lancashire Heeler) | Confidential | — | 0 | Clear |
| Amelogenesis Imperfecta (Discovered in the Parson Russell Terrier) | ENAM | C>T | 0 | Clear |
| Bandera's Neonatal Ataxia | GRM1 | Insertion | 0 | Clear |
| Benign Familial Juvenile Epilepsy | LGI2 | A>T | 0 | Clear |
| Bernard-Soulier Syndrome (Discovered in the Cocker Spaniel) | GP9 | Deletion | 0 | Clear |