

DogCheck 4.0

Analysis Overview

19.02.2024

| Category | Abbreviation | Disorders, Coat Colors and Traits | Patent |
|-------------------|---------------|---|--------|
| Coat Color Traits | ay, aw, at, a | A-Locus | |
| Coat Color Traits | at-Loc | A-Locus Agouti (black and tan) | |
| Coat Color Traits | a-Loc | A-Lokus Agouti (recessive black) | |
| Coat Color Traits | ay-Loc | A-Lokus Agouti (sable) | |
| Coat Color Traits | ay-Loc | A-Lokus Agouti (sable) | |
| Disorder | AMS | Acral Mutilation Syndrom | |
| Disorder | ARDS | Acute Respiratory Distress Syndrome (Dalmatian) | |
| Coat Color Traits | OCA4 | Albinism, oculocutaneous (Small Breeds) | |
| Disorder | AxD | Alexander Disease (Labrador Retriever) | |
| Disorder | AI | Amelogenesis Imperfecta (Italian Greyhound) | |
| Disorder | CaCa | Ataxia, CNS Atrophy with Cerebellar Ataxia (Belgian Shepherd) | |
| Disorder | SPAID | Autoinflammatory Disease (Shar-Pei) | |
| Coat Color Traits | bc, bd, bs | B-Locus | |
| Coat Color Traits | ba-Loc | B-Locus Brown ba | |
| Coat Color Traits | bc-Loc | B-Locus Brown bc | |
| Coat Color Traits | bd-Loc | B-Locus Brown bd | |
| Coat Color Traits | bs-Loc | B-Locus Brown bs | |
| Disorder | BFJE | Benign Familial Juvenile Epilepsy (Lagotto Romagnolo) | |
| Traits | Body Size | Body Size (GHR1) | |
| Traits | Body Size | Body Size (GHR2) | |
| Traits | Body Size | Body Size (HMGA2) | |
| Traits | Body Size | Body Size (IGF1) | |
| Traits | Body Size | Body Size (IGFR1) | |
| Traits | Body Size | Body Size (STC2) | |
| Traits | Dental | Body Size-Dental Anomaly 1 (Shetland Sheepdog) | |
| Traits | Body Size | Body Size-Dental Anomaly 2 (Shetland Sheepdog) | |
| Traits | | Brachycephaly, Shortened Skull Shape | |
| Disorder | CMSD | Canine Multiple Systemdegeneration (Chinese Crested) | |
| Disorder | CMSD | Canine Multiple Systemdegeneration (Kerry Blue Terrier) | |
| Disorder | CSS | Canine Scott Syndrome (German Shepherd dog) | |
| Disorder | DCM | Cardiomyopathy, dilated (Schnauzer) | |
| Disorder | DCM | Cardiomyopathy, dilated, Risk Variant 2 (Doberman Pinscher) | |
| Disorder | CJM | Cardiomyopathy, Juvenile Mortality (Belgian Malinois) | |
| Disorder | | Catalase Deficiency (Beagle) | |
| Disorder | CNM | Centronuclear Myopathy (Great Dane) | |
| Disorder | CA1 | Cerebellar Ataxia (Belgian Shepherd) | |
| Disorder | | Cerebellar Cortical Degeneration (Vizsla) | |
| Disorder | HA/CA | Cerebellar Degeneration (Setter) | |
| Disorder | CDPA | Chondrodysplasia (Karelian Bear Dog) | |

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| Coat Color Traits | Co-Loc | Co-Locus Cocoa Brown (French Bulldog) | |
| Disorder | CEA | Collie Eye Anomaly | *xP |
| Disorder | C3 Def | Complement 3 Deficiency (Brittany) | |
| Disorder | CD | Cone Degeneration | |
| Disorder | CD | Cone Degeneration (German Shepherd Dog) | |
| Disorder | CD | Cone Degeneration (Labrador Retriever) | |
| Disorder | | Congenital Methaemoglobinaemia (Pomeranian) | |
| Disorder | CMS | Congenital Myasthenic syndrome (Labrador Retriever) | |
| Disorder | CSNB | Continental Stationary Night Blindness (Briard) | |
| Disorder | | Copper Toxicosis (Labrador Retriever) | |
| Other | | Copper Toxicosis-protective Modifier (Labrador Retriever) | |
| Disorder | CMO | Cranio-mandibular Osteopathy | |
| Coat Traits | Cu1-Loc | Cu1-Locus Curly Coat (Common Variant) | |
| Disorder | Cyst-2a | Cystinuria (Australian Cattle Dog) | |
| Disorder | Cyst-1a | Cystinuria (Labrador Retriever) | |
| Disorder | Cyst-2 | Cystinuria (Miniature Pinscher) | |
| Disorder | Cyst-1a | Cystinuria (Newfoundland) | |
| Disorder | Cyst3-1 | Cystinuria Risk Factor Type 3 Variant 1 (Bulldog) | |
| Disorder | Cyst3-2 | Cystinuria Risk Factor Type 3 Variant 2 (Bulldog) | |
| Disorder | Cyst3-3 | Cystinuria Risk Factor Type 3 Variant 3 (Bulldog) | |
| Coat Color Traits | d1-Loc | D-Locus d1 Dilute (Common variant) | |
| Coat Color Traits | d2-Loc | D-Locus d2 Dilute | |
| Disorder | DWLM | Dandy-Walker-Like Syndrome (Eurasier) | |
| Disorder | EOAD | Deafness (Rhodesian Ridgeback) | |
| Disorder | Deafness | Deafness (Rottweiler) | |
| Disorder | DM-2 | Degenerative Myelopathy (Bernese Mountain dog) | |
| Disorder | DM | Degenerative Myelopathy (Classic Variant) | *xP |
| Disorder | DM-Modifikator | Degenerative Myelopathy, Early-Onset Risk Modifier (Pembroke Welsh Corgi) | |
| Disorder | | Dental Hypomineralization (Border Collie) | |
| Disorder | CKCSID | Dry eye curly coat syndrome (Cavalier King Charles Spaniel)* | |
| Traits | Body Size | Dwarfism, Growth-Hormone Deficiency (Chihuahua) | |
| Coat Color Traits | e-Loc | E-Locus e1 Rezessive Red (Common Variant) | |
| Coat Color Traits | e3-Loc | E-Locus e3 Rezessive Red (Husky) | |
| Coat Color Traits | Ea-Loc | E-Locus Ea Ancient Domino | |
| Coat Color Traits | Eg-Loc | E-Locus Eg Grizzle Domino | |
| Coat Color Traits | Eh-Loc | E-Locus Eh Cocker Sable | |
| Coat Color Traits | Em-Loc | E-Locus Em Melanistic Mask | |
| Traits | | Ear Erectness, Pricked Ears vs. Floppy Ears | |

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|-------------------|--------------|--|--------|
| Disorder | ED-SFS | Ectodermal Dysplasia (Chesapeake Bay Retriever) | |
| Disorder | XHED | Ectodermal Dysplasia (Dachshund) | |
| Disorder | EDS | Ehlers-Danlos Syndrome Variant 1 (Poodle) | |
| Disorder | EDS | Ehlers-Danlos Syndrome Variant 2 | |
| Disorder | HE | Elliptocytosis (Labrador Retriever) | |
| Disorder | AHE | Encephalopathy (Alaskan Husky) | |
| Disorder | DEB | Epidermolysis Bullosa, dystrophic (Golden Retriever) | |
| Disorder | JEB | Epidermolysis Bullosa, junctionalis (Australian Shepherd) | |
| Disorder | | Epilepsy with Mitochondrial Dysfunction and Neurodegeneration (Parson Russell Terrier) | |
| Disorder | EIC | Exercise Induced Collapse | *xP |
| Disorder | F7 | Factor VII Deficiency | |
| Disorder | FN | Familial Nephropathy (Cocker Spaniel) | |
| Disorder | | Familial Nephropathy (Springer Spaniel) | |
| Disorder | | Gallbladder Mucoceles | |
| Disorder | GM1 | Gangliosidosis GM1 (Portuguese Water Dog) | |
| Disorder | GM1 | Gangliosidosis GM1 (Shiba Inu) | |
| Disorder | GM2 | Gangliosidosis GM2 (Poodle) | |
| Disorder | GM2-1b | Gangliosidosis GM2 Type 1b (Japanese Chin) | |
| Disorder | GT | Glanzmann Thrombasthenia (Otterhound) | |
| Disorder | GG/PCAG | Glaucoma and Goniodysgenesis (Border Collie) | |
| Disorder | GLD | Globoid Cell Leukodystrophy (Terrier) | |
| Disorder | GSD Ia | Glycogen Storage Disease Ia (Maltese) | |
| Disorder | GSD IIIa | Glycogen Storage Disease IIIa (Curly Coated Retriever) | |
| Disorder | GSD VII | Glycogen Storage Disease VII (English Springer Spaniel) | |
| Disorder | GSD VII | Glycogen Storage Disease VII (Wachtelhund) | |
| Coat Color Traits | H-Loc | H-Locus Harlequin (Great Dane) | |
| Disorder | F8 | Hemophilia A (Boxer) | |
| Disorder | HC1 | Hereditary Cataracts | |
| Disorder | HC2 | Hereditary Cataracts (Australian Shepherd Type) | |
| Disorder | HFH | Hereditary Footpad Hyperkeratosis (Irish Terrier & Kromfohländer) | |
| Disorder | HNPk | Hereditary Nasal Parakeratosis (Greyhound) | |
| Disorder | HNPk | Hereditary Nasal Parakeratosis (Labrador Retriever) | *xP |
| Disorder | XLHN | Hereditary XL Nephritis (Samoyed) | |
| Disorder | HUU | Hyperuricosuria | |
| Disorder | HYM | Hypomyelination (Weimaraner) | |
| Disorder | CHG | Hypothyroidism (French Bulldog) | |
| Disorder | CHG | Hypothyroidism (Toy Fox Terrier) | |
| Other | Hypoxia | Hypoxia (Altitude Adaption) | |

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| Coat Color Traits | I-Loc | I-Locus Intensity (Pheomelanin Dilution) | |
| Coat Traits | IC-Loc | IC-Locus Improper Coat (Furnishing) | |
| Disorder | ICH | Ichthyosis (American Bulldog) | |
| Disorder | ICH | Ichthyosis (Great Dane) | |
| Disorder | ICH-GR1 | Ichthyosis 1 (Golden Retriever) | |
| Disorder | | Inflammatory Myopathy (Hollands Herder) | |
| Disorder | IPD | Inflammatory Pulmonary Disease (Rough Collie) | |
| Disorder | IGS-Beagle | Intestinal Cobalamin Malabsorption (Beagle) | |
| Disorder | IGS-BC | Intestinal Cobalamin Malabsorption (Border Collie) | |
| Disorder | IGS-KOM | Intestinal Cobalamin Malabsorption (Komondor) | |
| Disorder | JME | Juvenile Myoclonic Epilepsy (Rhodesian Ridgeback) | |
| Coat Color Traits | K-Loc | K-Locus Dominant Black | |
| Disorder | L-2-HGA | L-2-Hydroxyglutaricacidemia (Staffordshire Bull Terrier) | |
| Coat Traits | L1-Loc | L1-Locus Long Hair (Common Variant) | |
| Coat Traits | L2-Loc | L2-Locus Long Hair (Akita) | |
| Coat Color Traits | L3-Loc | L3-Locus long hair (Eurasier) | |
| Coat Traits | L4-Loc | L4-Locus Long Hair (Afghan Hound, French Bulldog) | |
| Disorder | LSD | Lagotto Storage Disease (Lagotto Romagnolo) | |
| Disorder | LPPN3 | Laryngeal Paralysis with Polyneuropathy Type 3 (Leonberger) | |
| Disorder | CLAD-I | Leukocyte Adhesion Deficiency Type I (Irish Setter) | |
| Disorder | CLAD-III | Leukocyte Adhesion Deficiency Type III (German Shepherd Dog) | |
| Disorder | LD | Leukodystrophy (Schnauzer) | |
| Disorder | LEMP | Leukoencephalomyelopathy (Leonberger) | |
| Disorder | LS | Lundehund Syndrome (Norwegian Lundehund) | |
| Disorder | MTCP | Macrothrombocytopenia (Cavalier King Charles Spaniel) | |
| Disorder | MTCP | Macrothrombocytopenia (Jack Russell Terrier) | |
| Disorder | MCD | Macular Corneal Dystrophy (Labrador Retriever) | |
| Disorder | MPS IIIA | Mucopolysaccharidosis IIIA (Dachshund) | |
| Disorder | MPS VII | Mucopolysaccharidosis VII (German Shepherd Dog) | |
| Disorder | MDR1 | Multidrug-Resistance 1 | |
| Disorder | CMR1 | Multifocal Retinopathy 1 | |
| Disorder | CMR2 | Multifocal Retinopathy 2 (Coton de Tulear) | |
| Disorder | MD | Muscular Dystrophy 1 (Labrador Retriever) | |
| Disorder | MD | Muscular Dystrophy 2 (Labrador Retriever) | |
| Disorder | DMD | Muscular Dystrophy Duchenne (Golden Retriever) | |
| Disorder | DMD | Muscular Dystrophy Duchenne (Norfolk Terrier) | |
| Disorder | DMD | Muscular Dystrophy Duchenne 1 (Cavalier King Charles Spaniel) | |
| Disorder | DMD | Muscular Dystrophy Duchenne 2 (Cavalier King Charles Spaniel) | |

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| Disorder | LGMD | Muscular Dystrophy, Limb-Girdle (Dachshund) | |
| Disorder | MLS | Musladin-Lueke Syndrome (Beagle) | |
| Disorder | | Myostatin Deficiency (Whippet) | |
| Disorder | MTM-XL | Myotubular Myopathy (Labrador Retriever) | |
| Disorder | NEM | Nemaline Myopathy (American Bulldog) | |
| Disorder | NEWS | Neonatal Encephalopathy with Seizures (Poodle) | |
| Disorder | NAD | Neuroaxonal Dystrophy (Papillon) | |
| Disorder | NAD | Neuroaxonal Dystrophy (Spanish Water Dog) | |
| Disorder | NCL-10 | Neuronal Ceroid Lipofuscinosis 10 (American Bulldog) | |
| Disorder | NCL-4A | Neuronal Ceroid Lipofuscinosis 4A (American Staffordshire Terrier) | |
| Disorder | NCL-7 | Neuronal Ceroid Lipofuscinosis 7 (Chihuahua, Chinese Crested) | |
| Disorder | NCL-8 | Neuronal Ceroid Lipofuscinosis 8 (Saluki) | |
| Disorder | OCD | Osteochondrodysplasia (Miniature Poodle) | |
| Disorder | | Osteochondromatosis (American Staffordshire Terrier) | |
| Disorder | OI | Osteogenesis Imperfecta (Dachshund) | |
| Disorder | OI | Osteogenesis Imperfecta Type 3 (Golden Retriever) | |
| Disorder | P2Y12 | P2RY12-Receptor Platelet Disorder (Greater Swiss Mountain Dog) | |
| Disorder | PMDS | Persistent Mullerian duct syndrome (Schnauzer) | |
| Disorder | PKD | Polycystic Kidney Disease (Bull Terrier) | |
| Traits | PPD | Polydactyly (Asian Breeds) | |
| Disorder | AMPN | Polyneuropathy (Alaskan Malamute) | |
| Disorder | | Polyneuropathy (Greyhound) | |
| Disorder | LPN2 | Polyneuropathy 2 (Leonberger) | |
| Disorder | POANV | Polyneuropathy with Ocular Abnormalities and Neuronal Vacuolation (Rottweiler) | |
| Disorder | | Prekallikrein Deficiency (Hairless Terrier, Shih-Tzu) | |
| Disorder | PLL | Primary Lens Luxation | |
| Disorder | PRA-Schnauzer | Progressive Retinal Atrophy (Giant Schnauzer) | |
| Disorder | GR-PRA1 | Progressive Retinal Atrophy 1 (Golden Retriever)* | |
| Disorder | PRA-Pap1 | Progressive Retinal Atrophy 1 (Papillon) | |
| Disorder | PRA-GR2 | Progressive Retinal Atrophy 2 (Golden Retriever) | |
| Disorder | PRA-AD | Progressive Retinal Atrophy, Autosomal Dominant (Mastiff) | |
| Disorder | PRA-crd1 | Progressive Retinal Atrophy, Cone-Rod Dystrophy 1 (American Pit Bull Terrier) | |
| Disorder | PRA-crd1/crd4 | Progressive Retinal Atrophy, Cone-Rod Dystrophy 4/crd1 | |
| Disorder | PRA-EO | Progressive Retinal Atrophy, Early Onset (Spanish Waterdog) | |
| Disorder | PRA-g | Progressive Retinal Atrophy, Generalized (Shapendoe) | |
| Disorder | PRA-PRCD | Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration | *xP |
| Disorder | PRA-rcd3 | Progressive Retinal Atrophy, Rod-cone dysplasia 3 (Cardigan Welsh Corgi) | |
| Disorder | PRA-rcd4 | Progressive Retinal Atrophy, Rod-cone dysplasia 4 | |

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|-------------------|--------------|---|--------|
| Disorder | PLN1 | Protein Losing Nephropathy 1 | |
| Disorder | PKDef | Pyruvate Kinase Deficiency (Labrador Retriever) | |
| Disorder | HRFCD | Renal Dysplasia and Hepatic Fibrosis (Norwich Terrier) | |
| Traits | | Screw Tail (Tail Curl) | |
| Coat Traits | SD-Loc | SD-Locus (Shedding) | |
| Traits | Sex | Sex Marker Amelogenin | |
| Disorder | SD2 | Skeletal Dysplasia 2 | |
| Coat Color Traits | Sp2-Loc | Sp2-Locus Piebald (White Spotting, Parti) | |
| Disorder | SDCA1 | Spongy Degeneration with Cerebellar Ataxia 1 (Belgian Malinois) | |
| Disorder | SDCA1 | Spongy Degeneration with Cerebellar Ataxia 1 (Jack Russell Terrier) | |
| Disorder | SDCA2 | Spongy Degeneration with Cerebellar Ataxia 2 (Belgian Malinois) | |
| Disorder | STGD | Stargardt Disease, Morbus Stargardt (Labrador Retriever) | |
| Disorder | SSADHD | Succinic Semialdehyde Dehydrogenase Deficiency (Saluki) | |
| Traits | T-Loc | T-Locus Brachyury (Bobtail, Naturel Short Tail) | |
| Disorder | TNS | Trapped Neutrophil Syndrome (Border Collie) | |
| Disorder | IVA | Ventricular Arrhythmias (Rhodesian Ridgeback) | |
| Disorder | HVDRR | Vitamin D-Deficiency Rickets (Pomeranian) | |
| Disorder | vWD I | Von Willebrand Disease I | |
| Disorder | vWD II | Von Willebrand Disease II (German Shorthaired Pointer) | |
| Disorder | vWD III | Von Willebrand Disease III (Kooiker Dog) | |
| Disorder | vWD III | Von Willebrand Disease III (Shetland Sheepdog) | |

* Results will only be provided if the dog's breed is one of the breeds mentioned in brackets.

*xP = requires paid patent panel