

# **Demographic Information**

Call Name Poison DOB Oct. 22, 2020

Registered Name - Registration # -

Breed Standard Poodle Tattoo -

Sex Female Microchip 956000013527984

Owner Beth Wolfe Laboratory # 367610 (AN-21-009836)

Report Date Dec. 30, 2022

## These tests were developed and performed by Paw Print Genetics®, Spokane WA.

## **Explanation of Results**

**Normal** A 'Normal' result means that your dog does not have the mutation that causes the associated genetic disease.

**Carrier** A 'Carrier' result indicates that your dog has inherited one copy of the mutation that has been reported to cause this genetic disease. Your dog may not be clinically affected by this mutation

because two copies of the mutation are usually required to cause disease.

Carrier / At-Risk A 'Carrier / At-Risk' result indicates that your dog inherited one copy of the mutation that has

been reported to cause this genetic disease. Based on the mode of genetic inheritance for this particular disease, inheriting one mutant copy of the gene may result in the disease. Dogs with one copy of the mutation may have a milder phenotype as compared to cats with two copies of

this mutation.

tilisiliatation

At-Risk / Affected An 'At-Risk / Affected' result indicates that your dog inherited one or two copies of the

mutation that has been reported to cause this genetic disease. Based on the mode of genetic inheritance for this particular disease, inheriting one or two mutant copies of the gene may

result in the disease.

#### No Result

'No Result' indicates that we were unable to obtain a genotype for your dog for this specific disease or trait and does not mean that your dog is a carrier or at-risk for this disease. There are a variety of reasons why a specific test may not provide a reportable result. Unique variations in the genetic code of some individuals may exist and cause certain regions of the genome to not perform properly with a specific test. In addition, suboptimal sampling of the dog's cheek cells could also result in poor sample performance due to inadequate cell counts, bacterial and fungal growth, or the presence of other test inhibitors. An acceptable level of tests with no results has been determined by Paw Print Genetics. Dogs with at least 90% of the test results are determined to be acceptable and reportable. If your dog has an unacceptable level of tests with no results, you will be contacted for a new sample to repeat the testing.

Please review our testing terms and disclaimers regarding your results.

	WT: (wild type (normal)) M	mutant Y: (Y chromosome (male)
Breed Profile		
Disease Name	Genotype	Interpretation
Degenerative Myelopathy	WT/WT	Normal (Clear)
Degenerative Myelopathy (Bernese Mountain Dog Variant)	0	
Degenerative Myelopathy (Common Variant)	0	
GM2 Gangliosidosis (Poodle Type)	WT/WT	Normal (clear)
Hereditary Cataracts	WT/WT	Normal (clear)
Intervertebral Disc Disease Risk Factor and Chondrodystrophy (CDDY with IVDD)	WT/WT	Normal (clear)
Multidrug Resistance 1	WT/WT	Normal (clear)
Neonatal Encephalopathy with Seizures	WT/WT	Normal (clear)
Osteochondrodysplasia	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration (prcd)	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Rod-Cone Dysplasia 4	No Result	No Result
Von Willebrand Disease I	No Result	No Result

Coat Colors & Traits		
Trait Name	Genotype	Interpretation
A Locus (Agouti)	A <sup>y</sup> /A <sup>t</sup>	Sable/fawn (carries tricolor/black and tan)

A <sup>s</sup> Locus (Saddle Tan)	N/A <sup>s</sup>	Saddle tan/creeping tan (non saddle tan carrier)
B Locus (Brown)	B/b	Black coat, nose and foot pads (carries brown)
B Locus (Brown) - b <sup>a</sup>	0	
B Locus (Brown) - b <sup>c</sup>	0	
B Locus (Brown) - b <sup>d</sup>	0	
B Locus (Brown) - b <sup>s</sup>	1	
Brachycephaly	BR/BR	Likely medium to long muzzle
Chondrodysplasia (CDPA)	cd/cd	Likely typical leg length
Cu Locus (Curly Hair)	Cu/Cu <sup>C</sup>	Curly/wavy coat (carrier)
D Locus (Dilute)	D/D	Non dilute
D Locus (Dilute) - d <sup>1</sup>	0	
D Locus (Dilute) - d <sup>2</sup>	0	
E Locus (Yellow/Red)	e/e	Yellow/red
E <sup>g</sup> Locus (Grizzle, Afghan Hound Type)	N/N	No grizzle
E <sup>h</sup> Locus (Sable, Cocker Spaniel Type)	N/N	No sable
E <sup>m</sup> Locus (Melanistic Mask)	N/N	No melanistic mask
H Locus (Harlequin, Great Dane Type)	h/h	No harlequin
Hr Locus (FOXI3 Hairless Gene Test, Mexican Hairless, Peruvian Hairless and Chinese Crested Type)	hr/hr	Coated
I Locus (Intensity)	1/1	Normal intensity
IC Locus (Improper Coat/Furnishings)	F/F	Furnishings

K Locus (Dominant Black)	K <sup>B</sup> /k <sup>y</sup>	No agouti expression allowed (carrier)
L Locus (Long Hair/Fluffy) - Lh <sup>1</sup> , Lh <sup>2</sup>	Lh/Lh	Longhaired
L Locus (Long Hair/Fluffy) - Lh <sup>1</sup>	2	
L Locus (Long Hair/Fluffy) - Lh <sup>2</sup>	0	
M Locus (Merle)	Fail	Non merle
Polydactyly	pd/pd	Normal (typical) toes (likely no hind dewclaws)
S Locus (White Spotting, Parti, or Piebald)	S/S	No white spotting, flash, parti, or piebald
SD Locus (Shedding)	sd/SD	Moderate shedding
Sex Determination	X/X	Female
T Locus (Natural Bobtail)	t/t	Normal tail

Determinants of coat colors and traits are complex. Many of these variants are known and many of the genes screened in the Canine HealthCheck interact. In addition, not all the genetic factors that contribute to a dog's coat color and traits are known. Because of the complexities in gene-gene interactions, the coat colors and traits reported in your Canine HealthCheck results may vary from your dog's actual appearance. Individual differences in genes throughout the canine genome, not tested in this genetic screen, may also affect the final coat color or traits seen in your dog.

WT: wild type (normal)

M: (mutant)

Y: (Y chromosome (male))

#### **Diseases Disease Name** Genotype Interpretation **Acral Mutilation Syndrome** WT/WT Normal (clear) **Acute Respiratory Distress Syndrome** WT/WT Normal (clear) **Adult Paroxysmal Dyskinesia** WT/WT Normal (clear) WT/WT Alaskan Husky Encephalopathy Normal (clear)

Alaskan Malamute Polyneuropathy	WT/WT	Normal (clear)
Amelogenesis Imperfecta	WT/WT	Normal (clear)
Benign Familial Juvenile Epilepsy	WT/WT	Normal (clear)
Canine Multiple System Degeneration (Chinese Crested Type)	WT/WT	Normal (clear)
Canine Multiple System Degeneration (Kerry Blue Terrier Type)	WT/WT	Normal (clear)
Canine Scott Syndrome	WT/WT	Normal (clear)
Catalase Deficiency	WT/WT	Normal (clear)
Centronuclear Myopathy	WT/WT	Normal (clear)
Cerebellar Ataxia (Finnish Hound Type)	WT/WT	Normal (clear)
Cerebellar Cortical Degeneration	WT/WT	Normal (clear)
Cerebellar Degeneration	WT/WT	Normal (clear)
Chondrodysplasia (Karelian Bear Dog and Norwegian Elkhound Type)	WT/WT	Normal (clear)
Cleft Palate and Syndactyly (Nova Scotia Duck Tolling Retriever Type)	WT/WT	Normal (clear)
Coagulation Factor VII Deficiency	WT/WT	Normal (clear)
Collie Eye Anomaly	WT/WT	Normal (clear)
Complement 3 Deficiency	WT/WT	Normal (clear)
Cone Degeneration	WT/WT	Normal (clear)
Cone Degeneration (German Shepherd Dog Type)	WT/WT	Normal (clear)

Cone Degeneration (German Shorthaired Pointer Type)	WT/WT	Normal (clear)
Cone Degeneration (Labrador Retriever Type)	WT/WT	Normal (clear)
Congenital Hypothyroidism with Goiter (Terrier Type)	No Result	No Result
Congenital Methemoglobinemia	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome (Jack Russell Terrier Type)	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome (Labrador Retriever Type)	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome (Old Danish Pointer Type)	WT/WT	Normal (clear)
Congenital Stationary Night Blindness	WT/WT	Normal (clear)
Craniomandibular Osteopathy	WT/WT	Normal (clear)
Cyclic Neutropenia	WT/WT	Normal (clear)
Cystinuria (Australian Cattle Dog Type)	WT/WT	Normal (clear)
Cystinuria (Labrador Retriever Type)	No Result	No Result
Cystinuria (Miniature Pinscher Type)	WT/WT	Normal (clear)
Cystinuria (Newfoundland Type)	WT/WT	Normal (clear)
Cystinuria Type 3 (Bulldog Type Risk Factor, Variant 3)	WT/WT	Normal (clear)
Cystinuria Type 3 (Bulldog Type Risk Factor, Variants 1 and 2)	WT/WT	Normal (Clear)
Cystinuria Type 3 (Bulldog Type Risk Factor, Variant 1)	0	
Cystinuria Type 3 (Bulldog Type Risk Factor, Variant 2)	0	
Dandy-Walker-Like Malformation	WT/WT	Normal (clear)
Degenerative Myelopathy	WT/WT	Normal (Clear)

Epidermolytic Hyperkeratosis	WT/WT	Normal (clear)
Elliptocytosis	WT/WT	Normal (clear)
Ehlers-Danlos Syndrome (Variant 2)	0	
Ehlers-Danlos Syndrome (Variant 1)	0	
Ehlers-Danlos Syndrome	WT/WT	Normal (Clear)
Ectodermal Dysplasia, X-Linked (Shepherd Type)	WT/WT	X-Linked Female Normal
Ectodermal Dysplasia, X-Linked (Dachshund Type)	WT/WT	X-Linked Female Normal
Ectodermal Dysplasia (Chesapeake Bay Retriever Type)	WT/WT	Normal (clear)
Early Retinal Degeneration	WT/WT	Normal (clear)
Dystrophic Epidermolysis Bullosa	WT/WT	Normal (clear)
Dry Eye Curly Coat Syndrome	No Result	No Result
Dilated Cardiomyopathy (Schnauzer Type)	WT/WT	Normal (clear)
Dilated Cardiomyopathy (Doberman Pinscher Type Risk Factor, Variant 2)	WT/WT	Normal (clear)
Dilated Cardiomyopathy (Doberman Pinscher Type Risk Factor, Variant 1)	WT/WT	Normal (clear)
Diffuse Cystic Renal Dysplasia and Hepatic Fibrosis	WT/WT	Normal (clear)
Dental Hypomineralization	WT/WT	Normal (clear)
Degenerative Myelopathy Early-Onset Risk Modifier (Pembroke Welsh Corgi Type)	WT/WT	Normal (clear)
Degenerative Myelopathy (Common Variant)	0	
Degenerative Myelopathy (Bernese Mountain Dog Variant)	0	

Episodic Falling Syndrome	WT/WT	Normal (clear)
Exercise-Induced Collapse	WT/WT	Normal (clear)
Factor XI Deficiency	WT/WT	Normal (clear)
Familial Nephropathy (Cocker Spaniel Type)	WT/WT	Normal (clear)
Familial Nephropathy (English Springer Spaniel Type)	WT/WT	Normal (clear)
Fucosidosis	No Result	No Result
Gallbladder Mucoceles	WT/WT	Normal (clear)
Glanzmann's Thrombasthenia (Great Pyrenees Type)	WT/WT	Normal (clear)
Glanzmann's Thrombasthenia (Otterhound Type)	WT/WT	Normal (clear)
Glaucoma (Border Collie Type)	WT/WT	Normal (clear)
Globoid Cell Leukodystrophy (Irish Setter Type)	WT/WT	Normal (clear)
Globoid Cell Leukodystrophy (Terrier Type)	WT/WT	Normal (clear)
Glycogen Storage Disease Ia	WT/WT	Normal (clear)
Glycogen Storage Disease IIIa	WT/WT	Normal (clear)
Glycogen Storage Disease VII (Wachtelhund Type)	WT/WT	Normal (clear)
Glycogen Storage Disease VII, PFK Deficiency	WT/WT	Normal (clear)
GM1 Gangliosidosis (Alaskan Husky Type)	WT/WT	Normal (clear)
GM1 Gangliosidosis (Portuguese Water Dog Type)	WT/WT	Normal (clear)
GM1 Gangliosidosis (Shiba Inu Type)	WT/WT	Normal (clear)
GM2 Gangliosidosis (Japanese Chin Type)	WT/WT	Normal (clear)

GM2 Gangliosidosis (Poodle Type)	WT/WT	Normal (clear)
Greyhound Polyneuropathy	No Result	No Result
Hemophilia A (Boxer Type)	WT/WT	X-Linked Female Normal
Hemophilia A (German Shepherd Dog, Type 1)	WT/WT	X-Linked Female Normal
Hemophilia A (German Shepherd Dog, Type 2)	WT/WT	X-Linked Female Normal
Hemophilia B (Cairn Terrier Type)	WT/WT	X-Linked Female Normal
Hemophilia B (Lhasa Apso Type)	WT/WT	X-Linked Female Normal
Hemophilia B (Rhodesian Ridgeback Type)	WT/WT	X-Linked Female Normal
Hereditary Cataracts (Australian Shepherd Type)	WT/WT	Normal (clear)
Hereditary Cataracts	WT/WT	Normal (clear)
Hereditary Footpad Hyperkeratosis (Irish Terrier and Kromfohrländer Type)	WT/WT	Normal (clear)
Hereditary Nasal Parakeratosis	WT/WT	Normal (clear)
Hereditary Nasal Parakeratosis (Greyhound Type)	WT/WT	Normal (clear)
Hereditary Nephritis (Samoyed Type)	WT/WT	X-Linked Female Normal
Hyperuricosuria	WT/WT	Normal (clear)
Hypomyelination (Weimaraner Type)	WT/WT	Normal (clear)
Ichthyosis (American Bulldog Type)	WT/WT	Normal (clear)
Ichthyosis (Golden Retriever Type 1)	WT/WT	Normal (clear)
Ichthyosis (Great Dane Type)	WT/WT	Normal (clear)

Inherited Myopathy of Great Danes	WT/WT	Normal (clear)
Intervertebral Disc Disease Risk Factor and Chondrodystrophy (CDDY with IVDD)	WT/WT	Normal (clear)
Intestinal Cobalamin Malabsorption (Beagle Type)	WT/WT	Normal (clear)
Intestinal Cobalamin Malabsorption (Border Collie Type)	WT/WT	Normal (clear)
Juvenile Laryngeal Paralysis and Polyneuropathy	WT/WT	Normal (clear)
Juvenile Myoclonic Epilepsy (Rhodesian Ridgeback Type)	WT/WT	Normal (clear)
L-2-Hydroxyglutaric Aciduria (Staffordshire Bull Terrier Type)	WT/WT	Normal (clear)
Lagotto Storage Disorder	WT/WT	Normal (clear)
Late Onset Ataxia	WT/WT	Normal (clear)
Lethal Acrodermatitis	WT/WT	Normal (clear)
Leukocyte Adhesion Deficiency, Type I	WT/WT	Normal (clear)
Leukocyte Adhesion Deficiency, Type III	WT/WT	Normal (clear)
Ligneous Membranitis	WT/WT	Normal (clear)
Lundehund Syndrome	WT/WT	Normal (clear)
Macular Corneal Dystrophy (Labrador Retriever Type)	WT/WT	Normal (clear)
May-Hegglin Anomaly	WT/WT	Normal (clear)
Mucopolysaccharidosis I	WT/WT	Normal (clear)
Mucopolysaccharidosis IIIA (Dachshund Type)	WT/WT	Normal (clear)
Mucopolysaccharidosis IIIA (New Zealand Huntaway Type)	WT/WT	Normal (clear)
,		

Mucopolysaccharidosis VII (Shepherd Type)	WT/WT	Normal (clear)
Multidrug Resistance 1	WT/WT	Normal (clear)
Multifocal Retinopathy 1	WT/WT	Normal (clear)
Multifocal Retinopathy 2	WT/WT	Normal (clear)
Multifocal Retinopathy 3	WT/WT	Normal (clear)
Muscular Dystrophy (Golden Retriever Type)	WT/WT	X-Linked Female Normal
Musladin-Lueke Syndrome	No Result	No Result
Myostatin Deficiency (Whippet and Longhaired Whippet Type)	WT/WT	Normal (clear)
Myotonia Congenita (Australian Cattle Dog Type)	WT/WT	Normal (clear)
Myotonia Congenita (Schnauzer Type)	WT/WT	Normal (clear)
Myotubular Myopathy 1	WT/WT	X-Linked Female Normal
Myotubular Myopathy 1 (Rottweiler Type)	WT/WT	X-Linked Female Normal
Narcolepsy (Dachshund Type)	WT/WT	Normal (clear)
Narcolepsy (Doberman Pinscher Type)	WT/WT	Normal (clear)
Narcolepsy (Labrador Retriever Type)	WT/WT	Normal (clear)
Neonatal Cerebellar Cortical Degeneration	WT/WT	Normal (clear)
Neonatal Encephalopathy with Seizures	WT/WT	Normal (clear)
Neuroaxonal Dystrophy (Rottweiler Type)	WT/WT	Normal (clear)
Neuroaxonal Dystrophy (Spanish Water Dog Type)	WT/WT	Normal (clear)

Neuronal Ceroid Lipofuscinosis (Tibetan Terrier Type)	No Result	No Result
Neuronal Ceroid Lipofuscinosis 1 (Cane Corso Type)	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 1	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 10	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 12	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 2	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 4A	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 5 (Australian Cattle Dog/Border Collie Type)	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 5 (Golden Retriever Type)	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 6	No Result	No Result
Neuronal Ceroid Lipofuscinosis 7	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 8 (Australian Shepherd Type)	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 8 (Setter Type)	WT/WT	Normal (clear)
Oculocutaneous Albinism	WT/WT	Normal (clear)
Oculocutaneous Albinism (Small Breed Type)	WT/WT	Normal (clear)
Osteochondrodysplasia	WT/WT	Normal (clear)
Osteogenesis Imperfecta (Beagle Type)	WT/WT	Normal (clear)
Osteogenesis Imperfecta (Dachshund Type)	No Result	No Result
Osteogenesis Imperfecta (Golden Retriever Type)	WT/WT	Normal (clear)

P2RY12 Receptor Platelet Disorder	WT/WT	Normal (clear)
Pembroke Welsh Corgi Duchenne Muscular Dystrophy	WT/WT	X-Linked Female Normal
Persistent Müllerian Duct Syndrome	WT/WT	Normal (clear)
Polyneuropathy (Leonberger and Saint Bernard Type)	WT/WT	Normal (clear)
Polyneuropathy (Leonberger Type 2)	WT/WT	Normal (clear)
Polyneuropathy with Ocular Abnormalities and Neuronal Vacuolation	WT/WT	Normal (clear)
Pompe Disease	WT/WT	Normal (clear)
Prekallikrein Deficiency	WT/WT	Normal (clear)
Primary Ciliary Dyskinesia	WT/WT	Normal (clear)
Primary Hyperoxaluria	WT/WT	Normal (clear)
Primary Lens Luxation	WT/WT	Normal (clear)
Primary Open Angle Glaucoma (Basset Fauve de Bretagne Type)	WT/WT	Normal (clear)
Primary Open Angle Glaucoma (Basset Hound Type)	WT/WT	Normal (clear)
Primary Open Angle Glaucoma	WT/WT	Normal (clear)
Primary Open Angle Glaucoma (Norwegian Elkhound Type)	WT/WT	Normal (clear)
Primary Open Angle Glaucoma and Primary Lens Luxation (Shar Pei Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Basenji Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Bullmastiff/Mastiff Type)	WT/WT	Normal (clear)

Progressive Retinal Atrophy (Giant Schnauzer Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Irish Setter Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Shetland Sheepdog Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Sloughi Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 1	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 2	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 3	No Result	No Result
Progressive Retinal Atrophy, Cone-Rod Dystrophy 4 (crd4/cord1)	WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Generalized	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Golden Retriever 1	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Golden Retriever 2	WT/WT	Normal (clear)
Progressive Retinal Atrophy, PRA1 (Papillon Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy, PRA3 (Tibetan Terrier and Spaniel Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration (prcd)	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Rod-Cone Dysplasia 3	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Rod-Cone Dysplasia 4	No Result	No Result
Progressive Retinal Atrophy, X-Linked 1	WT/WT	X-Linked Female Normal

Protein Losing Nephropathy	WT/WT, WT/WT	Normal (Clear) - No Increased Risk
Protein Losing Nephropathy (Variant 1)	0	
Protein Losing Nephropathy (Variant 2)	0	
Pyruvate Dehydrogenase Deficiency	WT/WT	Normal (clear)
Pyruvate Kinase Deficiency (Basenji Type)	No Result	No Result
Pyruvate Kinase Deficiency (Beagle Type)	WT/WT	Normal (clear)
Pyruvate Kinase Deficiency (Labrador Retriever Type)	WT/WT	Normal (clear)
Pyruvate Kinase Deficiency (Pug Type)	WT/WT	Normal (clear)
Pyruvate Kinase Deficiency (Terrier Type)	WT/WT	Normal (clear)
Recurrent Inflammatory Pulmonary Disease	WT/WT	Normal (clear)
Renal Cystadenocarcinoma and Nodular Dermatofibrosis	WT/WT	Normal (clear)
Retinal Dysplasia/Oculoskeletal Dysplasia 1	No Result	No Result
Retinal Dysplasia/Oculoskeletal Dysplasia 2	No Result	No Result
Sensory Neuropathy (Border Collie Type)	WT/WT	Normal (Clear)
Severe Combined Immunodeficiency Disease (Terrier Type)	WT/WT	Normal (clear)
Severe Combined Immunodeficiency Disease (Wetterhoun Type)	WT/WT	Normal (clear)
Severe Combined Immunodeficiency Disease, X-Linked (Basset Hound Type)	WT/WT	X-Linked Female Normal
Severe Combined Immunodeficiency Disease, X-Linked (Corgi Type)	WT/WT	X-Linked Female Normal

Shar-Pei Autoinflammatory Disease	WT/WT	Normal (clear)
Skeletal Dysplasia 2	WT/WT	Normal (clear)
Spinal Dysraphism	WT/WT	Normal (clear)
Spinocerebellar Ataxia	WT/WT	Normal (clear)
Spondylocostal Dysostosis	WT/WT	Normal (clear)
Stargardt Disease	WT/WT	Normal (clear)
Startle Disease	WT/WT	Normal (clear)
Thrombopathia (American Eskimo Dog Type)	WT/WT	Normal (clear)
Thrombopathia (Basset Hound Type)	WT/WT	Normal (clear)
Thrombopathia (Newfoundland Type)	WT/WT	Normal (clear)
Trapped Neutrophil Syndrome	WT/WT	Normal (clear)
Urolithiasis (Native American Indian Dog Type)	WT/WT	Normal (clear)
Van Den Ende-Gupta Syndrome	No Result	No Result
Von Willebrand Disease I	No Result	No Result
Von Willebrand Disease II	WT/WT	Normal (clear)
Von Willebrand Disease III (Kooikerhondje Type)	WT/WT	Normal (clear)
Von Willebrand Disease III (Scottish Terrier Type)	WT/WT	Normal (clear)
Von Willebrand Disease III (Shetland Sheepdog Type)	WT/WT	Normal (clear)

 $\label{eq:wild_type} \text{WT: wild type (normal)} \qquad \text{M: mutant} \qquad \text{Y: (Y chromosome (male))}$ 

Helm Smit

Chl

### Helen F Smith, PhD

Christina J Ramirez, PhD, DVM, DACVP

**Associate Laboratory Director** 

**Medical Director** 

Canine HealthCheck® is a product of Paw Print Genetics®. This test was developed and its performance determined by Paw Print Genetics®. This laboratory has established and verified the test's accuracy and precision. Because all tests performed are DNA-based, rare genomic variations may interfere with the performance of some tests producing false results. If you think any results are in error, please contact the laboratory for further evaluation.