

ORTHOPEDIC FOUNDATION FOR ANIMALS, INC.

DISCOVERY TAILS I LOVE LUCY
registered name

AUSTRALIAN LABRADOODLE
breed

254KSK
film/test/lab #

956000014958529
tattoo/microchip/DNA profile

2567395
application number

09/17/2025
date of report

RESULTS:

The 09/12/2025 exam finds this dog free of significant observable inherited eye disease. The Eye Certification Registry Number is valid for one year after the exam. Breeder option code(s): E2: Posterior Suture Tip Opacities, are potentially inherited conditions observed but not representing compromise of comfort, vision, or other ocular function.

ALAA126437, ALAA126437
registration no.

F
sex

05/14/2023
date of birth

27
age at evaluation in months



A Not-For-Profit Organization

LD-EYE10092/27F-VPI
O.F.A. NUMBER

*This number issued with the right to correct or
revoke by the Orthopedic Foundation for Animals.*

NORMAL W/BREEDER OPTIONS NOTED

owner
VICKI MCCORMACK
DISCOVERY TAILS LABRADOODLES LLC
81 EASTGATE PL
SEQUIM WA 98382



Verify QR scan

G.G. Keller, DVM

G.G. KELLER, DVM, MS, DACVR
CHIEF OF VETERINARY SERVICES

www.ofa.org

This electronic OFA certificate was generated on: 09/17/2025

This certification can be verified on the OFA website by entering the dog's registration number into the orange search box located at the top of the page or by scanning the QR code above.

If there are any errors on this certificate, please email **CORRECTIONS@OFA.ORG** to request a correction.

Orthopedic Foundation for Animals, Inc.
2300 E. Nifong Blvd.
Columbia, MO 65201-3806

OFA website: www.ofa.org
E-mail address: ofa@ofa.org
Phone number: 573-442-0418
Fax number: 573-875-5073

Orthopedic Foundation for Animals
Preliminary Hip Dysplasia Evaluation Report



A Not-for-Profit
Organization

DISCOVERY TAILS I LOVE LUCY
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AUSTRALIAN LABRADOODLE
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film/test/lab #

956000014958529

tattoo/microchip/DNA profile

2567395

application number

10/10/2024

date of report

ALAA126437, WALA00092779
registration no.

F
sex

05/14/2023
date of birth

16
age at evaluation in months

Owner

VICKI MCCORMACK
81 EASTGATE PL
SEQUIM WA 98382

Veterinarian

SEQUIM ANIMAL HOSPITAL
202 N 7TH AVE; PO BOX 668
SEQUIM WA 98382

Preliminary Hip Dysplasia Evaluation Report

No radiographic evidence of hip dysplasia is present. The consensus evaluation is: EXCELLENT

☒ EXCELLENT HIP JOINT CONFORMATION

superior hip joint conformation as compared with other individuals of the same breed and age

☐ GOOD HIP JOINT CONFORMATION

well formed hip joint conformation as compared with other individuals of the same breed and age

☐ FAIR HIP JOINT CONFORMATION

minor irregularities of the hip joint conformation as compared with other individuals of the same breed and age

☐ BORDERLINE HIP JOINT CONFORMATION

marginal hip joint conformation of indeterminate status with respect to hip dysplasia at this time -- Repeat study in six months

☐ MILD HIP DYSPLASIA

radiographic evidence of minor dysplastic changes of the hip joints

☐ MODERATE HIP DYSPLASIA

well defined radiographic evidence of dysplastic changes of the hip joints

☐ SEVERE HIP DYSPLASIA

radiographic evidence of marked dysplastic changes of the hip joints

RADIOGRAPHIC FINDINGS

- ☐ subluxation
☐ remodeling of femoral head/neck
☐ osteoarthritis/degenerative joint disease
☐ shallow acetabula
☐ acetabular rim/edge change

- ☐ unilateral ☐ left ☐ right
☐ transitional vertebra
☐ spondylosis
☐ panosteitis

G.G. KELLER, DVM, MS, DACVR
CHIEF OF VETERINARY SERVICES

Orthopedic Foundation for Animals
Preliminary Elbow Dysplasia Evaluation Report



A Not-for-Profit
Organization

DISCOVERY TAILS I LOVE LUCY
registered name

AUSTRALIAN LABRADOODLE
breed

film/test/lab #

956000014958529

tattoo/microchip/DNA profile

2567395

application number

10/10/2024

date of report

ALAA126437, WALA00092779
registration no.

F
sex

05/14/2023
date of birth

16
age at evaluation in months

Owner

VICKI MCCORMACK
81 EASTGATE PL
SEQUIM WA 98382

Veterinarian

SEQUIM ANIMAL HOSPITAL
202 N 7TH AVE; PO BOX 668
SEQUIM WA 98382

Preliminary Elbow Dysplasia Evaluation Report

✓ negative for elbow dysplasia

L ✓ R ✓

ELBOW DYSPLASIA

GRADE I

GRADE II

GRADE III

L _____ R _____

L _____ R _____

L _____ R _____

RADIOGRAPHIC FINDINGS

degenerative joint disease (DJD)

united anconeal process (UAP)

fragmented coronoid process (FCP)

osteochondrosis

L _____ R _____

L _____ R _____

L _____ R _____

L _____ R _____

G.G. Keller DVM

G.G. KELLER, DVM, MS, DACVR
CHIEF OF VETERINARY SERVICES

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Orthopedic Foundation for Animals

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 Email ofa@ofa.org | www.ofa.org
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Application for Patellar Luxation Database

Registered name: DISCOVERY TAILS I LOVE LUCY			AKC registration number:			Other registry name: ALAA		
Breed: AUSTRALIAN LABRADOODLE			Sex: F			Date of birth (MM/DD/YY): 05/14/2023		
Microchip/tattoo: 956000014958529			Registration number of sire: WALA00092779			Registration number of dam: ALAA126437		
Owner name: DISCOVERY TAILS LABRADOODLES LLC			Date of evaluation (MM/DD/YY): 10/07/2024					
Co-owner name: VICKI MCCORMACK			Examining veterinary clinic: SEQUIM ANIMAL HOSPITAL					
Mailing address: 81 EASTGATE PL			Mailing address: 202 N 7TH AVE					
City: SEQUIM	State: WA	Zip/postal code: 98382	City: SEQUIM	State: WA	Zip/postal code: 98382			
Phone: (360) 808-9800	E-mail: DISCOVERYTAILS@GMAIL.COM		Phone: (360) -683-7286	E-mail: SAH@SEQUIMANIMALHOSPITAL.COM				

I hereby certify that the information submitted is of the animal described on this application. I understand that by submitting these results to the OFA, if the animal was 12 months or older at the time of the exam, the results will be released to the public. Exams on animals under 12 months of age are considered preliminary, are not eligible for OFA certification numbers, and the results will not be released to the public.

Signature of owner or authorized representative

Vicki McCormack

Patellar Examination Results

1. Normal

☒ normal right ☒ normal left

2. Patellar Luxation

☐ bilateral ☐ right ☐ left
☐ unilateral: ☐ medial ☐ lateral
☐ luxated: ☐ intermittent ☐ permanent
 luxation is: ☐ < 2 months ☐ 2-6 months
 age of onset: ☐ 6-12 months ☐ > 12 months

3. Classification of luxation

- ☐ **Grade 1**—The patella easily luxates manually at full extension of the stifle joint, but returns to the trochlea when released.
- ☐ **Grade 2**—There is frequent patellar luxation which, in some cases, becomes more or less permanent.
- ☐ **Grade 3**—The patella is permanently luxated with torsion of the tibia and deviation of the tibial crest of between 30 degrees and 50 degrees from the cranial/caudal plane.
- ☐ **Grade 4**—The tibia is medially twisted and the tibial crest may show further deviation medially with the result that it lies 50 degrees to 90 degrees from the cranial/caudal plane.

☒ I certify that the examination was performed according to the OFA procedure.

☒ I DID verify microchip/tattoo on this dog ☐ I DID NOT verify microchip/tattoo on this dog

Heath Shurt
 Veterinarian Signature Specialty: ☒ Practitioner ☐ Specialist

10/7/24
 Date

Fees Animals over 12 months.....\$15.00 each
 A litter of 3 or more submitted together.....\$30.00 total

Exams on animals under 12 months of age are considered preliminary evaluations and are not eligible for OFA numbers

Kennel rate: Individuals submitted as a group, owned/co-owned by the same person
 Minimum of 5 individuals \$10.00 each

Payments can be made by Visa, Mastercard, check or money order (U.S. funds drawn on a U.S. bank) payable to the Orthopedic Foundation for Animals.

Do Not Submit

Card number

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07/2024

Submit thru <https://online.ofa.org> - OR - provide payment details here if mailing or emailing

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Orthopedic Foundation for Animals

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v010122

Application for Basic Cardiac Database

Registered name: DISCOVERY TAILS I LOVE LUCY		AKC registration number:		Other registry name: ALAA	
Breed: AUSTRALIAN LABRADOODLE		Sex: F		Date of birth (MM/DD/YY): 05/14/2023	
Microchip/tattoo: 956000014958529		Registration number of sire: WALA00092179		Registration number of dam: ALA A000126437	
Owner name: DISCOVERY TAILS LABRADOODLES LLC		Co-Owner name: VICKI MCCORMACK		Examining veterinary clinic: SEQUIM ANIMAL HOSPITAL	
Mailing address: 81 EASTGATE PL		Mailing address: 202 N 7TH AVE		Date of evaluation (MM/DD/YY): 10/07/2024	
City: SEQUIM	State: WA	Zip/postal code: 98382	City: SEQUIM	State: WA	Zip/postal code: 98382
Phone: (360) 808-9800	E-mail: DISCOVERYTAILS@GMAIL.COM	Phone: (360) -683-7286	E-mail: SAH@SEQUIMANIMALHOSPITAL.COM		

I hereby certify that the animal examined is the animal described on this application. I understand that by submitting these results to the OFA, if the animal was 12 months or older at the time of the exam, the results will be released to the public. Exams on animals under 12 months of age are considered preliminary, are not eligible for OFA certification numbers, and the results will not be released to the public.

Signature of owner or authorized representative

Vicki McCormack

Veterinary Exam Results

Clinical findings based on cardiac auscultation is required. (see page 2)

AUSCULTATION (REQUIRED)					
Normal <input checked="" type="checkbox"/>	Abnormal <input type="checkbox"/>	Arrhythmia <input type="checkbox"/>			
Murmur Grade: I <input type="checkbox"/>	II <input type="checkbox"/>	III <input type="checkbox"/>	IV <input type="checkbox"/>	V <input type="checkbox"/>	VI <input type="checkbox"/>
PMI: Left <input type="checkbox"/>	Right <input type="checkbox"/>	Base <input type="checkbox"/>	Apex <input type="checkbox"/>		
Timing: Systolic <input type="checkbox"/>	Diastolic <input type="checkbox"/>	Continuous <input type="checkbox"/>			
Extra Sounds: Click <input type="checkbox"/>	Gallop <input type="checkbox"/>	Split S1 <input type="checkbox"/>	Split S2 <input type="checkbox"/>		

Summary evaluation and opinion of the examiner:

- ☒ Normal cardiovascular examination—heart disease is not evident
☐ Equivocal cardiovascular examination—heart disease cannot be diagnosed nor excluded; status uncertain for breeding.
☐ Abnormal cardiovascular examination indicative of heart disease; indicate suspected diagnosis below:

☒ I certify that the standards for cardiac examination as set forth by the OFA were carefully followed in performing this examination.
☒ I DID verify microchip/tattoo on this dog ☐ I DID NOT verify microchip/tattoo on this dog

Debra Shott
Veterinarian Signature

Check one box: ☒ Practitioner, ☐ Specialist, ☐ Cardiologist

10/7/24
Date

Fees Animals Over 12 Months \$15.00 **Kennel Rate**—Individuals submitted as a group, owned/co-owned by same person.
Litter of 3 or more submitted together \$30.00 Minimum of 5 individuals \$10.00 each

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Do Not Submit
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Application for Patellar Luxation Database

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☒ I DID verify microchip/tattoo on this dog ☐ I DID NOT verify microchip/tattoo on this dog

Heath Shurt
 Veterinarian Signature Specialty: ☒ Practitioner ☐ Specialist

10/7/24
 Date

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Do Not Submit

Card number

Exp MM/YY

CW

07/2024

Submit thru <https://online.ofa.org> - OR - provide payment details here if mailing or emailing



Demographic Information

Call Name	Lucy
Registered Name	Discovery Tails I Love Lucy
Breed	Australian Labradoodle
Sex	Female
Owner	Vicki McCormack
DOB	May 14, 2023
Registration Number	WALA00092779
Tattoo	
Microchip	956000014958529
Laboratory #	441915
Report Date	March 28, 2024

These tests were developed and performed by Paw Print Genetics®, Lincoln, NE.

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 dogs with two copies of this mutation.
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	Geno.	Interpretation
Centronuclear Myopathy	WT/WT	Normal (Clear)
Cone Degeneration (Labrador Retriever Type)	WT/WT	Normal (Clear)
Congenital Myasthenic Syndrome (Labrador Retriever Type)	WT/WT	Normal (Clear)
Cystinuria (Labrador Retriever Type)	WT/WT	Normal (Clear)
Degenerative Myelopathy	WT/WT	Normal (Clear)
<div>Degenerative Myelopathy (Bernese Mountain Dog Variant) Degenerative Myelopathy (Common Variant)</div>		
Elliptocytosis	WT/WT	Normal (Clear)
Exercise-Induced Collapse	WT/WT	Normal (Clear)
Familial Nephropathy (Cocker Spaniel Type)	WT/WT	Normal (Clear)
Gallbladder Mucocoeles	WT/WT	Normal (Clear)
Glycogen Storage Disease VII, PFK Deficiency	WT/WT	Normal (Clear)
GM2 Gangliosidosis (Poodle Type)	WT/WT	Normal (Clear)
Hereditary Cataracts	WT/WT	Normal (Clear)
Hereditary Nasal Parakeratosis (Labrador Retriever Type)	WT/WT	Normal (Clear)
Hyperuricosuria	WT/WT	Normal (Clear)
Ichthyosis (Golden Retriever Type 1)	WT/WT	Normal (Clear)
Intervertebral Disc Disease Risk Factor and Chondrodystrophy (CDDY with IVDD)	WT/WT	Normal (Clear)
Macular Corneal Dystrophy (Labrador Retriever Type)	WT/WT	Normal (Clear)

Multidrug Resistance 1	WT/WT	<div>Normal (Clear)</div>
Myotonia Congenita (Labrador Retriever Type)	WT/WT	<div>Normal (Clear)</div>
Myotubular Myopathy 1 (Labrador Retriever Type)	WT/WT	<div>X-Linked Female Normal</div>
Narcolepsy (Labrador Retriever Type)	WT/WT	<div>Normal (Clear)</div>
Neonatal Encephalopathy with Seizures	WT/WT	<div>Normal (Clear)</div>
Osteochondrodysplasia	WT/WT	<div>Normal (Clear)</div>
Progressive Retinal Atrophy, Cone-Rod Dystrophy 4	WT/WT	<div>Normal (Clear)</div>
Progressive Retinal Atrophy, Golden Retriever 2	WT/WT	<div>Normal (Clear)</div>
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration (prcd)	WT/WT	<div>Normal (Clear)</div>
Progressive Retinal Atrophy, Rod-Cone Dysplasia 4	WT/WT	<div>Normal (Clear)</div>
Pyruvate Kinase Deficiency (Labrador Retriever Type)	WT/WT	<div>Normal (Clear)</div>
Retinal Dysplasia/Oculoskeletal Dysplasia 1	WT/WT	<div>Normal (Clear)</div>
Skeletal Dysplasia 2	WT/WT	<div>Normal (Clear)</div>
Stargardt Disease	WT/WT	<div>Normal (Clear)</div>
Ullrich Congenital Muscular Dystrophy (Labrador Retriever Type 1)	WT/WT	<div>Normal (Clear)</div>
Ullrich Congenital Muscular Dystrophy (Labrador Retriever Type 2)	WT/WT	<div>Normal (Clear)</div>
Von Willebrand Disease I	WT/WT	<div>Normal (Clear)</div>

WT:

wild type (normal)

 M:

mutant

 Y:

Y chromosome (male)

Coat Colors & Traits

Trait Name	Geno.	Interpretation
A Locus (Agouti)	a ^t /a ^t	Tricolor, black and tan
A^s Locus (Saddle Tan)	N/A ^s	Saddle tan/creeping tan (non saddle tan carrier)
B Locus (Brown)	B/B	Black coat, nose and foot pads (does not carry brown)
<div><div><div>B Locus (Brown) - b^a</div><div>B Locus (Brown) - b^c</div><div>B Locus (Brown) - b^d</div><div>B Locus (Brown) - b^h</div><div>B Locus (Brown) - b^e</div><div>B Locus (Brown) - b^s</div></div><div><div>0</div><div>0</div><div>0</div><div>0</div><div>0</div><div>0</div></div><div></div></div>		

Brachycephaly	BR/BR	Likely medium to long muzzle
Chondrodysplasia (CDPA)	cd/cd	Likely typical leg length
Co Locus (Cocoa, French Bulldog Type)	CO/CO	Black coat, nose and foot pads (does not carry cocoa)
Cu Locus (Curly Hair)	Cu/Cu	Straight coat
D Locus (Dilute)	D/D	Non-dilute (does not carry dilute)
<div><div>D Locus (Dilute) - d¹</div><div>D Locus (Dilute) - d²</div><div>D Locus (Dilute) - d³</div></div>		
E Locus	e ¹ /e ¹	Yellow/Red
<div><div>E Locus - E^m (Melanistic Mask)</div><div>E Locus - E^g (Grizzle, Afghan Hound Type)</div><div>E Locus - E^h (Sable, Cocker Spaniel Type)</div><div>E Locus - e^A (Ancient Red, Spitz and Scent Hound Type)</div><div>E Locus - e¹ (Yellow/Red)</div><div>E Locus - e² (Cream, Australian Cattle Dog Type)</div><div>E Locus - e³ (White, Alaskan and Siberian Husky Type)</div></div>		
H Locus (Harlequin, Great Dane Type)	h/h	No harlequin
Hairlessness	Rh/Rh	Coated
<div><div>Hairlessness (American Hairless Terrier Type) - rh¹</div><div>Hairlessness (Scottish Deerhound Type) - rh²</div></div>		
Hr Locus (FOXI3 Hairless Gene Test, Mexican Hairless, Peruvian Hairless and Chinese Crested Type)	hr/hr	Coated
I Locus (Intensity)	I/I	Normal intensity
IC Locus (Improper Coat/Furnishings)	F/F	Furnishings
K Locus (Dominant Black)	K ^B /K ^B	No agouti expression allowed
L Locus (Long Hair/Fluffy)	Lh ¹ /Lh ¹	Longhaired (carries two copies of long hair)
<div><div>L Locus (Long Hair/Fluffy) - Lh¹</div><div>L Locus (Long Hair/Fluffy) - Lh²</div><div>L Locus (Long Hair/Fluffy) - Lh³</div><div>L Locus (Long Hair/Fluffy) - Lh⁴</div></div>		
M Locus (Merle)	m/m	Non merle

Polydactyly (Common Variant)	pd/pd	Normal (typical) toes (likely no hind dewclaws)
Polydactyly (Great Pyrenees Type)	WT/WT	Normal (Clear)
R Locus (Roan/Ticked)	r/r	No roan or ticking
<div><div>R Locus (Roan/Ticked) - R^Ti</div><div>R Locus (Roan/Ticked) - R</div><div>0</div><div>0</div></div>		
S Locus (White Spotting, Parti, or Piebald)	S/s ^p	Limited white spotting, flash, parti, or piebald (carrier)
SD Locus (Shedding)	sd/SD	Moderate shedding
Sex Determination	X/X	Female
Social Behavior	WT/WT; WT/M	May demonstrate more social behavior
<div><div>Social Behavior, Variant 1</div><div>Social Behavior, Variant 2</div><div>0</div><div>1</div></div>		
T Locus (Natural Bobtail)	t/t	Normal tail

WT: **wild type (normal)** M: **mutant** Y: **Y chromosome (male)**

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.

Diseases

Disease Name	Geno.	Interpretation
Acral Mutilation Syndrome	WT/WT	Normal (Clear)
Acute Respiratory Distress Syndrome	WT/WT	Normal (Clear)
Adult Paroxysmal Dyskinesia	WT/WT	Normal (Clear)
Afibrinogenemia (Dachshund Type)	WT/WT	Normal (Clear)
Alaskan Husky Encephalopathy	WT/WT	Normal (Clear)
Alaskan Malamute Polyneuropathy	WT/WT	Normal (Clear)
Amelogenesis Imperfecta (Italian Greyhound Type)	WT/WT	Normal (Clear)
Amelogenesis Imperfecta (Parson Russell Terrier Type)	WT/WT	Normal (Clear)
Ataxia (Norwegian Buhund Type)	WT/WT	Normal (Clear)

Benign Familial Juvenile Epilepsy	WT/WT	Normal (Clear)
Bernard-Soulier Syndrome	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)
Cardiomyopathy and Juvenile Mortality	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 Ataxia 1 (Belgian Shepherd Type)	WT/WT	Normal (Clear)
Cerebellar Ataxia 2 (Belgian Shepherd Type)	WT/WT	Normal (Clear)
Cerebellar Cortical Degeneration	WT/WT	Normal (Clear)
Cerebellar Degeneration	WT/WT	Normal (Clear)
Charcot-Marie-Tooth Disease	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)	WT/WT	Normal (Clear)
Congenital Macrothrombocytopenia (Cairn and Norfolk Terrier Type)	WT/WT	Normal (Clear)
Congenital Methemoglobinemia	WT/WT	Normal (Clear)
Congenital Myasthenic Syndrome (Golden Retriever Type)	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)
Copper Storage Disease	WT/WT	Normal (Clear)
Craniomandibular Osteopathy.	WT/WT	Normal (Clear)
Cyclic Neutropenia	No Result	No Result
Cystinuria (Australian Cattle Dog Type).	WT/WT	Normal (Clear)
Cystinuria (Labrador Retriever Type).	WT/WT	Normal (Clear)
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)
<div><div>Cystinuria Type 3 (Bulldog Type Risk Factor, Variant 1)</div><div>Cystinuria Type 3 (Bulldog Type Risk Factor, Variant 2)</div><div>0</div><div>0</div><div></div></div>		
Dandy-Walker-Like Malformation	WT/WT	Normal (Clear)
Darier Disease and Associated Infundibular Cyst Formation	WT/WT	Normal (Clear)
Deafness and Vestibular Dysfunction (Doberman Pinscher Type), Variant 2	WT/WT	Normal (Clear)
Degenerative Myelopathy.	WT/WT	Normal (Clear)
<div><div>Degenerative Myelopathy (Bernese Mountain Dog Variant)</div><div>Degenerative Myelopathy (Common Variant)</div><div>0</div><div>0</div><div></div></div>		
Degenerative Myelopathy Early-Onset Risk Modifier (Pembroke Welsh Corgi Type).	WT/WT	Normal (Clear)
Dental Hypomineralization	WT/WT	Normal (Clear)
Diffuse Cystic Renal Dysplasia and Hepatic Fibrosis	WT/WT	Normal (Clear)
Dilated Cardiomyopathy_(Doberman Pinscher Type Risk Factor, Variant 1).	WT/WT	Normal (Clear)
Dilated Cardiomyopathy_(Doberman Pinscher Type Risk Factor, Variant 2).	WT/WT	Normal (Clear)

Dilated Cardiomyopathy (Schnauzer Type).	WT/WT	Normal (Clear)
Dry Eye Curly Coat Syndrome	WT/WT	Normal (Clear)
Dystrophic Epidermolysis Bullosa (Basset Hound Type).	WT/WT	Normal (Clear)
Dystrophic Epidermolysis Bullosa (Golden Retriever Type).	WT/WT	Normal (Clear)
Early Onset Adult Deafness (Rhodesian Ridgeback Type).	WT/WT	Normal (Clear)
Early Retinal Degeneration	WT/WT	Normal (Clear)
Early-Onset Epilepsy (Parson Russell Terrier Type).	WT/WT	Normal (Clear)
Ectodermal Dysplasia (Chesapeake Bay Retriever Type).	WT/WT	Normal (Clear)
Ectodermal Dysplasia, X-Linked (Dachshund Type).	WT/WT	X-Linked Female Normal
Ectodermal Dysplasia, X-Linked (Shepherd Type).	WT/WT	X-Linked Female Normal
Ehlers-Danlos Syndrome (Doberman Pinscher Type).	WT/WT	Normal (Clear)
Ehlers-Danlos Syndrome (Labrador Retriever Type), Variant 1	WT/WT	Normal (Clear)
Ehlers-Danlos Syndrome (Labrador Retriever Type), Variant 2	WT/WT	Normal (Clear)
Ehlers-Danlos Syndrome (Poodle Type, Variants 1 and 2).	WT/WT	Normal (Clear)
<div><div>Ehlers-Danlos Syndrome (Poodle Type), Variant 1</div><div>Ehlers-Danlos Syndrome (Poodle Type), Variant 2</div><div>0</div><div>0</div></div>		
Elliptocytosis	WT/WT	Normal (Clear)
Epidermolytic Hyperkeratosis	WT/WT	Normal (Clear)
Episodic Falling Syndrome	WT/WT	Normal (Clear)
Exercise-Induced Collapse	WT/WT	Normal (Clear)
Exfoliative Cutaneous Lupus Erythematosus (ECLE).	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	WT/WT	Normal (Clear)
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)
GM2 Gangliosidosis (Shiba Inu Type).	WT/WT	Normal (Clear)
Greyhound Polyneuropathy.	WT/WT	Normal (Clear)
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 A (Rhodesian Ridgeback Type).	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 Ataxia (Australian Shepherd Type).	WT/WT	Normal (Clear)
Hereditary Cataracts (Australian Shepherd Type).	WT/WT	Normal (Clear)
Hereditary Cataracts	WT/WT	Normal (Clear)
Hereditary Cataracts (Wirehaired Pointing Griffon Type).	WT/WT	Normal (Clear)
Hereditary Footpad Hyperkeratosis (Irish Terrier and Kromfohrländer Type).	WT/WT	Normal (Clear)
Hereditary Footpad Hyperkeratosis (Rottweiler Type).	WT/WT	Normal (Clear)
Hereditary Nasal Parakeratosis (Greyhound Type).	WT/WT	Normal (Clear)

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Hereditary Nasal Parakeratosis (Labrador Retriever 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 (Golden Retriever Type 2).			WT/WT		
			Normal (Clear)		
Ichthyosis (Great Dane Type).			WT/WT		
			Normal (Clear)		
Ichthyosis (Jack Russell Terrier Type).			WT/WT		
			Normal (Clear)		
Inflammatory Myopathy (Shepherd 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)		
Intestinal Cobalamin Malabsorption (Giant Schnauzer Type).			WT/WT		
			Normal (Clear)		
Intestinal Lipid Malabsorption			WT/WT		
			Normal (Clear)		
Junctional Epidermolysis Bullosa (Australian Shepherd Type).			WT/WT		
			Normal (Clear)		
Juvenile Laryngeal Paralysis and Polyneuropathy (Black Russian Terrier Type).			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)		
L-2-Hydroxyglutaric Aciduria (Yorkshire Terrier Type).			WT/WT		
			Normal (Clear)		
Lagotto Storage Disorder			WT/WT		
			Normal (Clear)		
Laryngeal Paralysis and Polyneuropathy (Leonberger Type 3).			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)
Limb-Girdle Muscular Dystrophy_(Dachshund Type)	WT/WT	Normal (Clear)
Lundehund Syndrome	WT/WT	Normal (Clear)
Macular Corneal Dystrophy_(Labrador Retriever Type)	WT/WT	Normal (Clear)
Mammary Tumors_(English Springer Spaniel Type Risk Factor)	WT/WT	Normal (Clear)
May-Hegglin Anomaly	WT/WT	Normal (Clear)
Microphthalmia_(Soft Coated Wheaten Terrier Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis I_(Boston Terrier Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis I_(Plott Hound Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis IIIA_(Dachshund Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis IIIA_(New Zealand Huntaway Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis IIIB_(Schipperke Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis VI_(Miniature Schnauzer Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis VII_(Brazilian Terrier 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	WT/WT	Normal (Clear)
Myostatin Deficiency_(Whippet Type)	WT/WT	Normal (Clear)
Myotonia Congenita_(Australian Cattle Dog Type)	WT/WT	Normal (Clear)
Myotonia Congenita_(Labrador Retriever Type)	WT/WT	Normal (Clear)
Myotonia Congenita_(Schnauzer Type)	WT/WT	Normal (Clear)
Myotubular Myopathy_1_(Boykin Spaniel Type)	WT/WT	X-Linked Female Normal
Myotubular Myopathy_1_(Labrador Retriever Type)	WT/WT	X-Linked Female Normal
Myotubular Myopathy_1_(Rottweiler Type)	WT/WT	X-Linked Female Normal

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Narcolepsy_(Dachshund Type).	WT/WT	Normal (Clear)
Narcolepsy_(Doberman Pinscher Type).	WT/WT	Normal (Clear)
Narcolepsy_(Labrador Retriever Type).	WT/WT	Normal (Clear)
Neonatal Ataxia	WT/WT	Normal (Clear)
Neonatal Cerebellar Cortical Degeneration	WT/WT	Normal (Clear)
Neonatal Encephalopathy with Seizures	WT/WT	Normal (Clear)
Neuroaxonal Dystrophy_(Giant Schnauzer Type).	WT/WT	Normal (Clear)
Neuroaxonal Dystrophy_(Papillon Type).	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).	WT/WT	Normal (Clear)
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_(Golden Retriever Type).	WT/WT	Normal (Clear)
Neuronal Ceroid Lipofuscinosis 5_(Herding Dog Type).	WT/WT	Normal (Clear)
Neuronal Ceroid Lipofuscinosis 6	WT/WT	Normal (Clear)
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)
Nonsyndromic Hearing Loss_(Rottweiler Type).	WT/WT	Normal (Clear)
Oculocutaneous Albinism_(Doberman Pinscher Type).	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).	WT/WT	Normal (Clear)

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Osteogenesis Imperfecta (Golden Retriever Type)	WT/WT	Normal (Clear)
P2RY12 Receptor Platelet Disorder	WT/WT	Normal (Clear)
Pancreatitis (Miniature Schnauzer Type Risk Factor)	WT/M	Carrier (No increased risk)
<div><div>Pancreatitis (Miniature Schnauzer Type Risk Factor), Variant 1</div><div>Pancreatitis (Miniature Schnauzer Type Risk Factor), Variant 2</div><div>Pancreatitis (Miniature Schnauzer Type Risk Factor), Variant 3</div><div>0</div><div>1</div><div>1</div></div>		
Pembroke Welsh Corgi Duchenne Muscular Dystrophy	WT/WT	X-Linked Female Normal
Persistent Müllerian Duct Syndrome	WT/WT	Normal (Clear)
Pituitary Dwarfism (Shepherd Type)	WT/WT	Normal (Clear)
Polyneuropathy_(Leonberger Type 1)	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 (Alaskan Malamute Type)	WT/WT	Normal (Clear)
Primary Ciliary Dyskinesia (Old English Sheepdog Type)	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)

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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_(Dachshund Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 1_(American Staffordshire Terrier Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 2_(American Staffordshire Terrier Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 3_(Glen of Imaal Terrier Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Cone-Rod Dystrophy 4		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Early Onset (Spanish Water Dog Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Early-Onset (Portuguese Water Dog Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Generalized (Schapendoes Type).		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, Late-Onset (Lapponian Herder Type).		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		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Syndromic Retinal Degeneration (Shetland Sheepdog Type).		WT/WT	Normal (Clear)
Progressive Retinal Atrophy, X-Linked 1 (Husky Type).		WT/WT	X-Linked Female Normal
Progressive Retinal Atrophy, X-linked 2		WT/WT	X-Linked Female Normal
Protein Losing Nephropathy.		WT/WT, WT/WT	Normal (Clear) - No Increased Risk
<div>Protein Losing Nephropathy (Variant 1) Protein Losing Nephropathy (Variant 2)</div>		<div>0 0</div>	

Pyruvate Dehydrogenase Deficiency	WT/WT	Normal (Clear)
Pyruvate Kinase Deficiency_(Basenji_Type)	WT/WT	Normal (Clear)
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	WT/WT	Normal (Clear)
Retinal Dysplasia/Oculoskeletal Dysplasia 2	WT/WT	Normal (Clear)
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_(Alpine Dachsbrake_Type)	WT/WT	Normal (Clear)
Spinocerebellar Ataxia_(Terrier_Type)	WT/WT	Normal (Clear)
Spondylocostal Dysostosis	WT/WT	Normal (Clear)
Stargardt Disease	WT/WT	Normal (Clear)
Startle Disease	WT/WT	Normal (Clear)
Subacute Necrotizing Encephalopathy_(Yorkshire Terrier_Type)	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)

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Trapped Neutrophil Syndrome	WT/WT	Normal (Clear)
Ullrich Congenital Muscular Dystrophy (Labrador Retriever Type 1).	WT/WT	Normal (Clear)
Ullrich Congenital Muscular Dystrophy (Labrador Retriever Type 2).	WT/WT	Normal (Clear)
Urolithiasis (Native American Indian Dog Type).	WT/WT	Normal (Clear)
Van Den Ende-Gupta Syndrome	WT/WT	Normal (Clear)
Von Willebrand Disease I	WT/WT	Normal (Clear)
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)

WT: **wild type (normal)** M: **mutant** Y: **Y chromosome (male)**

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 with >99% sensitivity and specificity. The results included in this report relate only to the items tested using the sample provided. The presence of mosaicism may not be detected by this test. Non-paternity may lead to unexpected results. This is not a diagnostic test. This is not a breed identification test. Because all tests are DNA-based, rare genomic variations may interfere with the performance of some individual tests producing false results. If you think any results are in error, please contact the laboratory for further evaluation.