



Demographic Information

Call Name	Millie
Registered Name	
Breed	Maine Coon
Sex	Female
Owner	Erica Kessler
DOB	June 9, 2021
Registration #	
Tattoo	
Microchip	██████████ 6044
Laboratory #	421441
Report Date	December 12, 2023

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

Explanation of Results

Normal	A 'Normal' result means that your cat does not have the mutation that causes the associated genetic disease.
Carrier	A 'Carrier' result indicates that your cat has inherited one copy of the mutation that has been reported to cause this genetic disease. Your cat 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 cat 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. Cats with one copy of the mutation may have a milder phenotype as compared to cats with two copies of this mutation.
At-Risk / Affected	An 'At-Risk / Affected' result indicates that your cat 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 cat for this specific disease or trait and does not mean that your cat 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 cat'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. Cats with at least 90% of the test results are determined to be acceptable and reportable. If your cat 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
Cystinuria, Type B, Variant 2 (Cat)	WT/WT	Normal (Clear)
Hypertrophic Cardiomyopathy (Maine Coon Type) (Cat)	WT/WT	Normal (Clear)
Pyruvate Kinase Deficiency (Cat)	WT/WT	Normal (Clear)
Spinal Muscular Atrophy (Cat)	WT/WT	Normal (Clear)

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Coat Colors & Traits

Trait Name	Geno.	Interpretation
ABC Blood Group System	A/A	A blood group
ABC Locus - B Group Variant 1 - b ¹	0	
ABC Locus - B Group Variant 2 - b ²	0	
ABC Locus - B Group Variant 3 - b ³	0	
ABC Locus - C Group Variant - a ^c	0	
Agouti Coat Color - A Locus Solid/Charcoal	a/a	Solid - No tabby expression allowed
Amber and Russet Coat Color - E Locus	E/E	Non-amber, darkly pigmented coat color
E Locus - Amber Variant - e	0	
E Locus - Russet Variant - e ^r	0	
Brown Coat Color - B Locus	B/B	Black Coat Color
B Locus - Cinnamon Variant - b ¹	0	
B Locus - Chocolate Variant - b	0	
Coat Type - Curly (Devon Rex, Selkirk Rex Type) or Hairless (Sphynx Type) - R Locus	R/R	Straight coat

R Locus - Selkirk Rex Curly Variant - SR	0
R Locus - Devon Rex Curly Variant - re	0
R Locus - Sphynx Hairless Variant - hr	0

[Curly Coat \(Cornish Rex Type\)](#) Cu/Cu Straight coat

[Dilute Coat Color - D Locus](#) D/D Non-dilute

[Dominant White and White Spotting - W Locus](#) w/w No white spotting

[Folded Ears with Osteochondrodysplasia](#) f/f Typical (non-folded) ears

[Golden/Sunshine Coat \(Siberian Type\) - Wb Locus](#) Wb/Wb Non-sunshine tabby

[Hairlessness \(Lykoi Type\) - Hr Locus](#) Hr/Hr Normal hair

Hairlessness (Lykoi Type) - Hr Locus - hr ^{Ca}	0
Hairlessness (Lykoi Type) - Hr Locus - hr ^{Fr}	0
Hairlessness (Lykoi Type) - Hr Locus - hr ^{NC}	0
Hairlessness (Lykoi Type) - Hr Locus - hr ^{TN}	0
Hairlessness (Lykoi Type) - Hr Locus - hr ^{TX}	0
Hairlessness (Lykoi Type) - Hr Locus - hr ^{VA}	0

[Long Hair - L Locus](#) lh³/lh³ Longhaired

L Locus - Long Hair Variant 1 - M1/lh ¹	0
L Locus - Long Hair Variant 2 - M2/lh ²	0
L Locus - Long Hair Variant 3 - M3/lh ³	2
L Locus - Long Hair Variant 4 - M4/lh ⁴	0
L Locus - Long Hair Variant 5 - M5/lh ⁵	0

[Pointed Coat Color and Albinism - C Locus](#) C/C Non-pointed coat

C Locus - Siamese Variant - c ^s	0
C Locus - Burmese Variant - c ^b	0
C Locus - Albino Variant - c	0
C Locus - Albino Variant 2 - c ²	0

[Polydactyly](#) pd/pd Normal (typical) toes

Polydactyly - Variant 1 - PD ¹	0
Polydactyly - Variant 2 - PD ²	0
Polydactyly - Hemingway Variant - PD ^H	0

[Short Tail \(Bobtail\) - T Locus](#) t/t Normal length tail

Short Tail (Bobtail) - T Locus - T ¹	0
Short Tail (Bobtail) - T Locus - T ²	0
Short Tail (Bobtail) - T Locus - T ³	0

[Short Tail \(Japanese Bobtail Type\)](#) st/st Normal length tail

[Tabby Coat Color Pattern - Mc Locus](#) mc¹/mc¹ Blotched (classic) tabby coat color pattern

Cystinuria, Type B, Variant 1	WT/WT	Normal (Clear)
Cystinuria, Type B, Variant 2	WT/WT	Normal (Clear)
Cystinuria, Type B, Variant 3	WT/WT	Normal (Clear)
Cystinuria, Type B, Variant 4	WT/WT	Normal (Clear)
Cystinuria, Type B, Variant 5	WT/WT	Normal (Clear)
Dihydropyrimidinase Deficiency	WT/WT	Normal (Clear)
Epidermolysis Bullosa Simplex	WT/WT	Normal (Clear)
Factor XII Deficiency, Variant 1	WT/WT	Normal (Clear)
Factor XII Deficiency, Variant 2	WT/WT	Normal (Clear)
Factor XII Deficiency, Variant 3	WT/WT	Normal (Clear)
Feline Immunodeficiency Virus (FIV) Infection Risk Modifier	WT/WT	No Increased Resistance to FIV Infection
Feline Leukocyte Adhesion Deficiency, Type 1	WT/WT	Normal (Clear)
Feline Niemann-Pick Disease	WT/WT	Normal (Clear)
Feline Spongy Encephalopathy	WT/WT	Normal (Clear)
Forebrain Commissural Malformation	WT/WT	Normal (Clear)
Gangliosidosis GM2A	WT/WT	Normal (Clear)
Glycogen Storage Disease, Type IV	WT/WT	Normal (Clear)
GM1 Gangliosidosis	WT/WT	Normal (Clear)
GM2 Gangliosidosis, Type II (Burmese Type)	WT/WT	Normal (Clear)
GM2 Gangliosidosis, Type II	WT/WT	Normal (Clear)
GM2 Gangliosidosis, Type II (Japanese Domestic Type)	WT/WT	Normal (Clear)
GM2 Gangliosidosis, Type II (Korat Type)	WT/WT	Normal (Clear)
Hemophilia B, Variant 1	WT/WT	X-Linked Female Normal
Hemophilia B, Variant 2	WT/WT	X-Linked Female Normal
Hyperlipoproteinemia	WT/WT	Normal (Clear)
Hypertrophic Cardiomyopathy (Maine Coon Type)	WT/WT	Normal (Clear)
Hypertrophic Cardiomyopathy (Ragdoll Type)	WT/WT	Normal (Clear)
Hypertrophic Cardiomyopathy (Sphynx Type Risk Factor)	WT/WT	Normal (Clear)
Hypogonadotropic Hypogonadism	WT/WT	Normal (Clear)

Hypokalemic Periodic Paralysis	WT/WT	Normal (Clear)
Hypotrichosis with Short Life Expectancy	WT/WT	Normal (Clear)
Inflammatory Linear Verrucous Epidermal Nevus	WT/WT	X-Linked Female Normal
L-2-Hydroxyglutaric Aciduria	WT/WT	Normal (Clear)
Methemoglobinemia, Variant 1	WT/WT	Normal (Clear)
Methemoglobinemia, Variant 2	WT/WT	Normal (Clear)
Mucopolipidosis II	WT/WT	Normal (Clear)
Mucopolysaccharidosis Type I	WT/WT	Normal (Clear)
Mucopolysaccharidosis Type VI (Modifier)	WT/WT	Normal (Clear)
Mucopolysaccharidosis Type VI (Siamese Type)	WT/WT	Normal (Clear)
Mucopolysaccharidosis Type VII, Variant 1	WT/WT	Normal (Clear)
Mucopolysaccharidosis Type VII, Variant 2	WT/WT	Normal (Clear)
Multiple Drug Resistance	WT/WT	Normal (Clear)
Myotonia Congenita	WT/WT	Normal (Clear)
Neuronal Ceroid Lipofuscinosis 6	WT/WT	Normal (Clear)
Neuronal Ceroid Lipofuscinosis 7, Variant 1	WT/WT	Normal (Clear)
Neuronal Ceroid Lipofuscinosis 7, Variant 2	WT/WT	Normal (Clear)
Niemann-Pick C1 Disease, Variant 1	WT/WT	Normal (Clear)
Niemann-Pick C1 Disease, Variant 2	WT/WT	Normal (Clear)
Niemann-Pick C2 Disease	WT/WT	Normal (Clear)
Oculocutaneous Albinism	WT/WT	Normal (Clear)
Polycystic Kidney Disease	WT/WT	Normal (Clear)
Polycystic Kidney Disease (Siberian Type)	WT/WT	Normal (Clear)
Primary Congenital Glaucoma	WT/WT	Normal (Clear)
Primary Hyperoxaluria Type II	WT/WT	Normal (Clear)
Progressive Retinal Atrophy (Abyssinian Type)	WT/WT	Normal (Clear)
Progressive Retinal Atrophy (Bengal Type)	WT/WT	Normal (Clear)
Progressive Retinal Atrophy (Persian Type)	WT/WT	Normal (Clear)
Pyruvate Kinase Deficiency	WT/WT	Normal (Clear)
Rod-Cone Dysplasia	WT/WT	Normal (Clear)

Spinal Muscular Atrophy	WT/WT	Normal (Clear)
Vitamin D-Dependent Rickets Type IB	WT/WT	Normal (Clear)
Vitamin D-dependent Rickets, Type IA, Variant 1	WT/WT	Normal (Clear)
Vitamin D-dependent Rickets, Type IA, Variant 2	WT/WT	Normal (Clear)

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Helen F Smith, PhD

Associate Laboratory Director



Christina J Ramirez, PhD, DVM, DACVP

Medical Director

CatScan 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 are performed are DNA-based, rare genomic variations may interfere with the performance of some tests producing false results. If you think these results are in error, please contact the laboratory for further evaluation.