Demographic Information

Call Name	Gent
Registered Name	
Breed	Ragdoll
Sex	Male
Owner	Pamela Banta
DOB	January 30, 2022
Registration #	
Tattoo	
Microchip	
Laboratory #	306658
Report Date	May 25, 2022

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

Explanation of Results

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

ABC Blood Group System	A/a^{c}	A blood group
ABC Locus - B Group Variant 1 - b^{1}	0	
ABC Locus - B Group Variant 2 - b^{2}	0	
ABC Locus - C Group Variant - a^{c}	1	
Agouti Coat Color - A Locus	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^{1}	Brown coat color
B Locus - Cinnamon Variant - b^{1}	1	
B Locus - Chocolate Variant - b	1	
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	Dilute
Dominant White and White Spotting - W Locus	w/w	No white spotting
Folded Ears with Osteochondrodysplasia	f/f	Typical (non-folded) ears
Long Hair - L Locus	lh/lh	Longhaired
L Locus - Long Hair Variant 1 - lh^{1}	0	
L Locus - Long Hair Variant 2 - lh^{2}	0	
L Locus - Long Hair Variant 3 - lh^{3}	0	
L Locus - Long Hair Variant 4 - lh^{4}	2	

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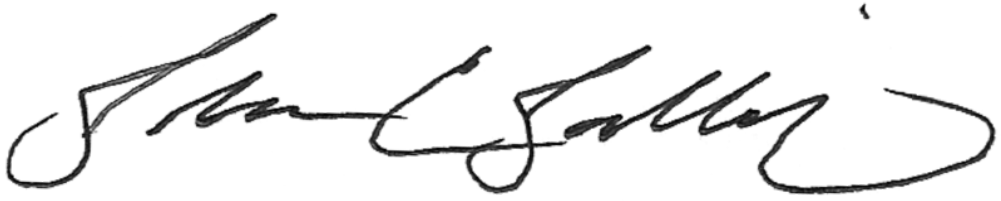
Acute Intermittent Porphyria, Variant 2	WT/WT	Normal (clear)
Acute Intermittent Porphyria, Variant 3	WT/WT	Normal (clear)
Acute Intermittent Porphyria, Variant 4 (Siamese Type 1)	WT/WT	Normal (clear)
Acute Intermittent Porphyria, Variant 5 (Siamese Type 2)	WT/WT	Normal (clear)
Acute Intermittent Porphyria, Variant 6	WT/WT	Normal (clear)
Autoimmune Lymphoproliferative Syndrome	WT/WT	Normal (clear)
Brachycephaly (Burmese Type)	WT/WT	Normal (clear)
Congenital Adrenal Hyperplasia	WT/WT	Normal (clear)
Congenital Erythropoietic Porphyria, Variant 1	WT/WT	Normal (clear)
Congenital Erythropoietic Porphyria, Variant 2	WT/WT	Normal (clear)
Congenital Hypothyroidism	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome	WT/WT	Normal (clear)
Cystinuria, Type 1A	WT/WT	Normal (clear)
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)
Factor XII Deficiency, Variant 1	WT/WT	Normal (clear)
Feline Leukocyte Adhesion Deficiency, Type 1	WT/WT	Normal (clear)

Myotonia Congenita	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)
Polycystic Kidney Disease	WT/WT	Normal (clear)
Primary Hyperoxaluria Type II	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Abyssinian Type)	WT/WT	Normal (clear)
Progressive Retinal Atrophy (Persian Type)	WT/WT	Normal (clear)
Pyruvate Kinase Deficiency	WT/WT	Normal (clear)
Spinal Muscular Atrophy	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)

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



Blake C Ballif, PhD

Laboratory & Scientific Director

Christina J Ramirez, PhD, DVM, DA

Medical Dir