Hobs House Bengals

Dottycat Texas Hold 'Em of Hobs House

Breed: Bengal
Colouration: Black (Brown) Spotted Tabby
Gender: King

Pedigree Of
Dottycat Texas Hold 'Em of HobsHouse

Sex: Male
Colour: Brown (Black) Spotted Tabby
SBT: .
Date of Birth: 05/02/2025

Wildnbeauty Dino of Dottycat Brown (Black) Spotted Tabby	CH Spice Zataar of Katzizi	Vom Karwendleberg Monte Carlo of Spice	CH Laristochat Des Griffes De Feu	Brownsugar Fast and Furious
				Jamanpur Des Griffes de Feu
			Burningbright First Crescent	Burningbright Halku
				Spice Tar Tar of Burningbright
		Spice Arella	GRC Leopardcats Mambo	IW SGC Fraservalley Denalis Thunder
				Leopardcats Express Myself
			CH Spice Nigella	RW SGC Kalnlkats Rocketstar
				TGC Spice Pimenta
	Pearlyqueen Lea of Wildnbeauty	Amazongold Luis Ronaldo	Batifoleurs Johan Dewitt of Amazongold	RW SGC Batifoleurs Armani
				CH Batifoleurs Mirabelle
			Amazongold Berta	RW GRC Amazongold Simba
				RW BW OGC Amazongold Russian Beauty
		Vengalcat Laia Rubia	Fraservalley Toblano of Vengalcat	RW OGC Jewelspride Solo
				Snopride Mystique of Jewelspride
			Leozeebah Rhea of Vengalcat	Leozeebah Callpso
				Leozeebha Callisto
Dottycat Clarity Brown (Black) Spotted Tabby	Dazzlingbronze Starsky of Dottycat	Urbanjaguars Kenzo	Custrogarra Niko	CH Tamerian Fleuramour of Old Race
				Olivia
			Finestbengals Panthera Regina	Murenacat Trigger
				Sharm Rose
		Pawsombengals STA of Dazzlingbronze	Brtiannia Romeo Prince of Lovekatz	Sunstorm Tiberius
				Queenanne Sandsoftime
			Noras Delight	Otus
				Snowfall Hugs and Kisses
	Dottycat Take My Time	Katzizi Soloman of Dottycat	Wildbengals Helios of Katzizi	RW OGC Batifoleurs Jabari
				Sherakan Magic Moment
			QGC Katzizi Zorina	Poshpaws Im So Awesome Look At Me
				Adthish Zuri
		Dottycat Nutkin	Jasiri Dartagnan of Dottycat	Solanaranch Serendipity of Jasari
				Jasari Golden Dawn
			Dottycat Harvest Dawn	Silverglam Ragnur of Olympusscats
				Olympusscats Clover

I certify this pedigree to be true. Signed: ..... *Colletta Waugh*

THE INTERNATIONAL CAT ASSOCIATION

certifies that under the rules of this association

Dottycat Texas Hold em of Hobshouse

is awarded the title of

Double Grand Champion

Saturday, December 13 2025

Frances Cardona
EXECUTIVE SECRETARY

THE INTERNATIONAL CAT ASSOCIATION

certifies that under the rules of this association

Dottycat Texas Hold em of Hobshouse

is awarded the title of

Grand Champion

Sunday, November 16 2025

Frances Cardona
EXECUTIVE SECRETARY

THE INTERNATIONAL CAT ASSOCIATION

certifies that under the rules of this association

Dottycat Texas Hold em of Hobshouse

is awarded the title of

Champion

Sunday, October 05

2025

Frances Cardona
EXECUTIVE SECRETARY

THE INTERNATIONAL CAT ASSOCIATION

certifies that under the rules of this association

Dottycat Texas Hold em of Hobshouse

is awarded the title of

Triple Grand Champion

Sunday, January 11 2026

Frances Cardona
EXECUTIVE SECRETARY

Veterinair Verwijscentrum HEELIX
Grouwesteenstraat 50
9170 Sint-Pauwels
+32 3 777 51 79

Cardiac Screening Report (HCM)

Patient Information

Name: **Dottycat Texas Hold Em of Hobs House,** [redacted]
Breed: cat, Bengal
Birthdate: 05-02-2025 (11 Months)
Gender: Male Not sterilized
Chip number: 953010007589750 (Transponder neck left)
Registration number: [redacted]

Owner: **Waugh Collette**
Cattery: Hobs House
Address: [redacted]
Phone number: +44 7814 398616
Email: [redacted]

Physical examination

General condition: good
Weight: 4 kg
Heartrate: .bpm

Cardiac ascultation: [8] normal
D murmur (grade: .../6)
D arrhythmia

Echocardlography

IVSd	4.5	mm
LVIDd	16.7	mm
LVFWd	4.4	mm
IVSs	6.0	mm
LVIDs	9.9	mm
LVFWs	7.9	mm
FS%	41	%
LA diam	11.0	mm
Ao diam	8.6	mm
LA/Ao	1.29	

Subjective left atrial size: [8] normal
D mild enlargement
D moderate enlargement
D severe enlargement

Papillary muscles: [8] normal
D abnormal

DLVOTO [8] no
☐yes (velocity: ... mis; pressure gradient: ... mmHg)

Assessment (based on phenotype)

[8] Normal
D Equivocal
☐ HCM (mild/moderate/severe)

☐ RCM
D Congenital heart disease:
☐ Other:

Veterinarian:

Veterina1r verwli scentrum HEELIX
Barbara Vandevelde, DVM , ECVDI
Grouwesteenstraat
9170 Sint-Gil\is-Waas, Belg1e
Tel. 0032 3 777 51 79
narbara.vandevelde@heel1x.be

01/'26

Barbara Vandevelde, DipECVDI, Medische beeldvorming - Bernac .tls· m Ryssen, OipECVSMR, Orlhopeclie en arlrosoopie • Maai Albers, European MAS-IM, Inlern1:.1l/tmt1&skuntf1J- Ju Vtumeman, Europear.t..MASaCJI.IJJ.tJa/Birtiart Cardiologie - Anneleen Gielen, Aneslhesie en assislenlie • Lisa Bena.li, dierenarlsassislenle

Veterinair Verwijscentrum HEELIX
Grouwesteenstraat 50
9170 Sint-Pauwels
+32 3 777 51 79

Ultrasonographic screening kidneys

Patient Information

Narne: **Dottycat Texas Hold Em of Hobs House,** [redacted]
Breed: cat, Bengal
Birthdate: 05-02-2025 (11 Months)
Gender: Male Not sterilized
Chip nurnber: 953010007589750 (Transponder neck left)
Registration nurnber: [redacted]

Owner: **Waugh Collette**
Cattery: Hobs House
Address: [redacted]
Phone nurnber: [redacted]
Ernail: [redacted]

Physical examination

General condition: good
Weight 4 kg
Palpation: no abnormalities

Echography

Both kidneys are present and normal in size, echogenicity and architecture.
Both kidneys are free of cysts, stones and cortical lesions.

Diagnosis

1R1 No significant abnorrnalities
☐PKO
☐CIN
D Agenesia
☐Other:

Breeding advice: positive

Veterinarian: *Barbara Vandevelde, DipECVDI*
European Specialist in Veterinary Medical Imaging

Date: 19/01/26

scentrum HEELIX
veterina1r verwiJ
bara Vandevelde,oVM,0- 1 ,p. ECVDI
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barbara.va

Barbara Vandevelde, DipECVDI, Mt=ldische bel'ldvormm!) - Bemadeltt'! Van Ryssen, D1pECVSMR, Or/hopet1ie en arlrosr.011il'! - Mani Albers, Ewn11enn MAS-IM, fnl1-1me q1lt'11..:;kunde - Jo Vnnnnmrw,
E11rt.1f.lt'!an MAS-CA in orlniding, Cardiologie - Anneft en Gielen, Aneslhesie tm assislentie

Dottycat Texas Hold'em

WISDOM PANEL™

Microchip number: 953010007589750
Kit type: Complete
ID kit: CLGQMTW
Test date: 2025-06-30

Dottycat Texas Hold'em's Profile

Pet information

Registered name
Dottycat Texas Hold'em

Date of birth
2025-02-05

Microchip number
953010007589750

Sex
M

Neutered
No

Top breeds

100% Bengal

Blood type summary

Blood type
Type A (most common)

Transfusion risk
⚠ Moderate

Health summary

At Risk 0 conditions

Carrier 1 condition

- Pyruvate Kinase Deficiency

Clear 48 conditions

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Breed ancestry

Dottycat Texas Hold’em appears to be 100% Bengal.

Asian

100 % Bengal

Dottycat Texas Hold'em

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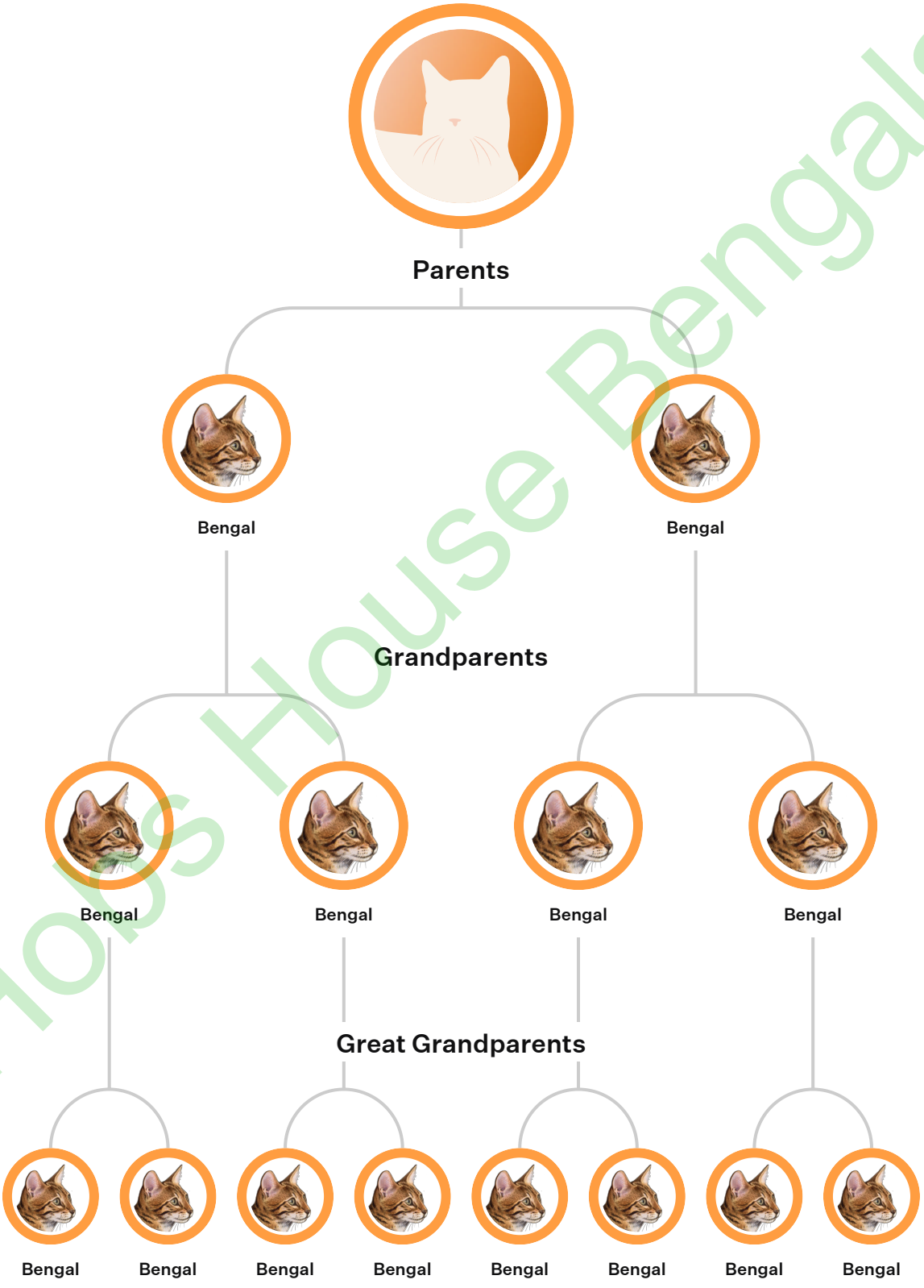
Family Tree



Dottycat Texas Hold’em

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Genetic Diversity

Heterozygosity

Dottycat Texas Hold’em’s Percentage of Heterozygosity

33%

Dottycat Texas Hold’em’s genome analysis shows an average level of genetic heterozygosity when compared with other Bengals.

Typical Range for Bengals

31% - 36%

Dottycat Texas Hold'em

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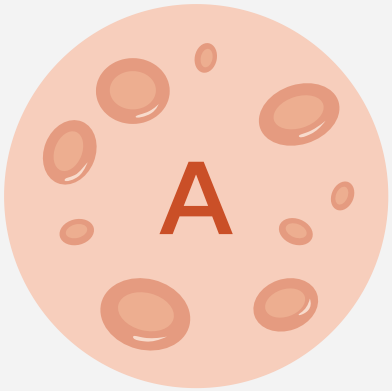
Blood Type



Blood type
Type A (Most common)

Genotype*
A/A

Transfusion risk
⚠ Moderate

Dottycat Texas Hold'em has the most common blood type. He can be transfused with Type A blood.

Blood variants tested*

Variant Tested	Description	Copies
b variant 1	(Common b variant)	0
b variant 2	(Discovered in Turkish breeds)	0
b variant 3	(Discovered in Ragdolls)	0
c variant - Causes AB Blood Type	(Discovered in Ragdolls)	0

*This test identifies three known 'b' variants and one known 'c' variant in the CMAH gene when determining a cat's genetic blood type. Blood Type A is inferred in reporting when less than two genetic blood variants are detected.

Dottycat Texas Hold'em

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Kit type: Complete
Test date: 2025-06-30

Summary of health conditions

Key Findings

We detected 1 genetic condition in Dottycat Texas Hold'em's DNA.

0	1	48
At Risk	Carrier	Clear

Genetic Condition	Gene	Risk Variant	Copies	Inheritance	Result
Pyruvate Kinase Deficiency	PKLR	G>A	1	AR	Carrier

What this means for Dottycat Texas Hold'em

Carrier **Pyruvate Kinase Deficiency**

Two copies of the Pyruvate Kinase Deficiency variant are needed for a cat to be affected by this condition, so Dottycat Texas Hold'em should not show disease signs due to this variant. Please note that similar disease signs could develop due to a different genetic or clinical cause.

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Health conditions tested

At-risk and carrier conditions (1)

Pyruvate Kinase Deficiency

Gene	Risk Variant	Copies	Inheritance	Result
PKLR	G>A	1	AR	Carrier

What is it

Pyruvate Kinase (PK) Deficiency is a disorder that causes anemia due to the breakdown of red blood cells.

What it means

Two copies of the Pyruvate Kinase Deficiency variant are needed for a cat to be affected by this condition, so Dottycat Texas Hold'em should not show disease signs due to this variant. Please note that similar disease signs could develop due to a different genetic or clinical cause.

What to do

Here's how to care for a cat with Pyruvate Kinase Deficiency

Partner with your veterinarian to make a plan regarding your cat's well-being, including any insights provided through genetic testing. If your pet is at risk or is showing signs of this disorder, then the first step is to speak with your veterinarian.

Dottycat Texas Hold'em

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Other health conditions tested

Genetic Condition	Gene	Risk Variant	Copies	Inheritance	Result
Acute Intermittent Porphyria (Variant 1)	HMBS	Deletion	0	AD	Clear
Acute Intermittent Porphyria (Variant 2)	HMBS	G>A	0	AD	Clear
Acute Intermittent Porphyria (Variant 3)	HMBS	Insertion	0	AD	Clear
Acute Intermittent Porphyria (Variant 4)	HMBS	Deletion	0	AD	Clear
Acute Intermittent Porphyria (Variant 5)	HMBS	G>A	0	AR	Clear
Autoimmune Lymphoproliferative Syndrome (Discovered in British Shorthair)	FASL	Insertion	0	AR	Clear
Burmese Head Defect (Discovered in the Burmese)	ALX1	Deletion	0	AD	Clear
Chediak-Higashi Syndrome (Discovered in the Persian)	LYST	Insertion	0	AR	Clear
Congenital Adrenal Hyperplasia	CYP11B1	G>A	0	AR	Clear
Congenital Erythropoietic Porphyria	UROS	G>A	0	AR	Clear
Congenital Myasthenic Syndrome (Discovered in the Devon Rex and Sphynx)	COLQ	G>A	0	AR	Clear
Cystinuria Type 1A	SCL3A1	C>T	0	AR	Clear
Cystinuria Type B (Variant 1)	SCL7A9	C>T	0	AR	Clear
Cystinuria Type B (Variant 2)	SCL7A9	G>A	0	AR	Clear
Cystinuria Type B (Variant 3)	SCL7A9	T>A	0	AR	Clear
Dihydropyrimidinase Deficiency	DPYS	G>A	0	AR	Clear
Earfold and Osteochondrodysplasia (Discovered in the Scottish Fold)	TRPV4	G>T	0	AD	Clear
Factor XII Deficiency (Variant 1)	F12	Deletion	0	ARa	Clear
Factor XII Deficiency (Variant 2)	F12	Deletion	0	ARa	Clear
Familial Episodic Hypokalemic Polymyopathy (Discovered in the Burmese)	WNK4	C>T	0	AR	Clear
Glutaric Aciduria Type II	ETFDH	T>G	0	AR	Clear

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Other health conditions tested

Genetic Condition	Gene	Risk Variant	Copies	Inheritance	Result
Glycogen Storage Disease (Discovered in the Norwegian Forest Cat)	GBE1	Insertion	0	AR	Clear
GM1 Gangliosidosis	GLB1	G>C	0	AR	Clear
GM2 Gangliosidosis	GM2A	Deletion	0	AR	Clear
GM2 Gangliosidosis Type II (Discovered in Domestic Shorthair cats)	HEXB	Insertion	0	AR	Clear
GM2 Gangliosidosis Type II (Discovered in Japanese domestic cats)	HEXB	C>T	0	AR	Clear
GM2 Gangliosidosis Type II (Discovered in the Burmese)	HEXB	Deletion	0	AR	Clear
Hemophilia B (Variant 1)	F9	C>T	0	XR	Clear
Hemophilia B (Variant 2)	F9	G>A	0	XR	Clear
Hyperoxaluria Type II	GRHPR	G>A	0	AR	Clear
Hypertrophic Cardiomyopathy (Discovered in the Maine Coon)	MYBPC	G>C	0	AR	Clear
Hypertrophic Cardiomyopathy (Discovered in the Ragdoll)	MYBPC	C>T	0	AD	Clear
Hypotrichosis (Discovered in the Birman)	FOXN1	Deletion	0	AR	Clear
Lipoprotein Lipase Deficiency	LPL	G>A	0	AR	Clear
MDR1 Medication Sensitivity	ABCB1	Deletion	0	AR	Clear
Mucopolysaccharidosis Type I	IDUA	Deletion	0	AR	Clear
Mucopolysaccharidosis Type VI	ARSB	T>C	0	AR	Clear
Mucopolysaccharidosis Type VI Modifier	ARSB	G>A	0	MO	Clear
Mucopolysaccharidosis Type VII (Variant 1)	GUSB	G>A	0	AR	Clear
Mucopolysaccharidosis Type VII (Variant 2)	USB	C>T	0	AR	Clear
Myotonia Congenita	CLCN1	G>T	0	AR	Clear
Polycystic Kidney Disease (PKD)	PKD1	C>A	0	AD	Clear

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Other health conditions tested

Genetic Condition	Gene	Risk Variant	Copies	Inheritance	Result
Progressive Retinal Atrophy (Discovered in the Abyssinian)	CEP290	T>G	0	AR	Clear
Progressive Retinal Atrophy (Discovered in the Bengal)	KIF3B	G>A	0	AR	Clear
Progressive Retinal Atrophy (Discovered in the Persian)	AIPL1	C>T	0	AR	Clear
Sphingomyelinosis (Variant 1)	NPC1	G>C	0	AR	Clear
Sphingomyelinosis (Variant 2)	NPC2	G>A	0	AR	Clear
Vitamin D-Dependent Rickets	CYP27B1	G>T	0	AR	Clear

Dottycat Texas Hold'em

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Traits

Coat Color

	Gene	Variant	Copies	Result
Charcoal (Discovered in the Bengal)	ASIP	A^{Pb}	0	No effect
Solid Color	ASIP	a	0	Banded hairs, tabby patterns likely
Partial and Full White	KIT	W or w^{s}	0	No effect
Amber (Discovered in the Norwegian Forest Cat)	MC1R	e	0	No effect
Russet (Discovered in the Burmese)	MC1R	e^{r}	0	No effect
Dilution	MLPH	d	0	No effect
Albinism (Discovered in Oriental breeds)	TYR	c^{a}	0	No effect
Colorpoint (Discovered in the Burmese)	TYR	c^{b}	0	No effect
Colorpoint (Discovered in the Siamese) Two copies of this variant result in a colorpoint pattern, although this can be blocked by other variants. Cats with one copy of the Colorpoint (Discovered in the Burmese) variant and one copy of the Colorpoint (Discovered in the Siamese) variant will show a darker base coat color and less contrasting colorpoint pattern than cats with two copies of the Colorpoint (Discovered in the Siamese) variant.	TYR	c^{s}	1	Colorpoints possible
Mocha (Discovered in the Burmese)	TYR	c^{m}	0	No effect
Chocolate	TYRP	b	0	No effect
Cinnamon	TYRP	b^{l}	0	No effect

Coat Type

	Gene	Variant	Copies	Result
Long Hair (Discovered in many breeds)	FGF5	M4	0	No effect
Long Hair (Discovered in the Norwegian Forest Cat)	FGF5	M2	0	No effect
Long Hair (Discovered in the Ragdoll and Maine Coon)	FGF5	M3	0	No effect

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Coat Type

	Gene	Variant	Copies	Result
Long Hair (Discovered in the Ragdoll)	FGF5	M1	0	No effect
Lykoi Coat (Variant 1)	HR	hr^{Ca}	0	No effect
Lykoi Coat (Variant 2)	HR	hr^{VA}	0	No effect
Hairlessness (Discovered in the Sphynx)	KRT71	re^{hr}	0	No effect
Rexing (Discovered in the Devon Rex)	KRT71	re^{dr}	0	No effect
Rexing (Discovered in the Cornish Rex and German Rex)	LPAR6	r	0	No effect
Glitter	Pending	gl	0	No effect

Tail Length

	Gene	Variant	Copies	Result
Short Tail (Variant 3)	HES7	jb	0	No effect
Short Tail (Variant 1)	T	C1199del	0	No effect
Short Tail (Variant 2)	T	T988del	0	No effect

Extra Toes

	Gene	Variant	Copies	Result
Polydactyly (Variant 1)	LIMBR1	HW	0	No effect
Polydactyly (Variant 2)	LIMBR1	UK1	0	No effect
Polydactyly (Variant 3)	LIMBR1	UK2	0	No effect

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Glossary of genetic terms

Test result definitions

At Risk: Based on the disorder's mode of inheritance, the cat inherited a number of genetic variant(s) which increases the cat's risk of being diagnosed with the associated disorder.

Carrier: The cat inherited one copy of a genetic variant when two copies are usually necessary to increase the cat's risk of being diagnosed with the associated disorder. While carriers are usually not at risk of clinical expression of the disorder, carriers of some complex variants may be associated with a low risk of developing the disorder.

Notable: Inheriting two copies of the genetic variant is noteworthy for specific aspects of health and breeding of the cat, but the cat should otherwise not suffer disease due to this genetic cause when in absence of other genetic variants.

Clear: The cat did not inherit the genetic variant(s) associated with the disorder and will not be at elevated risk of being diagnosed with the disorder due to this genotype. However, similar clinical signs could develop from different genetic or clinical causes.

Inconclusive: An inconclusive result indicates a confident call could not be made based on the data for that genetic variant. Health testing is performed in replicates, and on occasion the outcomes do not agree. This may occur due to an unusual sequence of DNA in the region tested, multiple cell genotypes present due to chimerism or acquired mutations, or due to quality of the DNA sample.

Inheritance mode definitions

Autosomal Recessive (AR): For autosomal recessive disorders, cats with two copies of the genetic variant are at risk of developing the associated disorder. Cats with one copy of the variant are considered carriers and are usually not at risk of developing the disorder. However, carriers of some complex variants grouped in this category may be associated with a low risk of developing the disorder. Cats with one or two copies may pass the disorder-associated variant to their kittens if bred.

Autosomal Recessive, asymptomatic (ARa): For autosomal recessive, asymptomatic disorders, cats with two copies of the variant can exhibit certain aspects of the variant-associated disorder but otherwise, they should not suffer clinical disease as typically expected with autosomal recessive disorders. Cats with one copy of the variant are called carriers and should not exhibit any aspect of the disorder. However, cats with one or two copies may pass the disorder-associated variant to their kittens if bred.

Autosomal Dominant (AD): For autosomal dominant disorders, cats with one or two copies of the genetic variant are at risk of developing the associated disorder. Inheriting two copies of the variant may increase the risk of development of the disorder or cause the condition to be more severe. These cats may pass the disorder-associated variant to their kittens if bred.

X-linked Recessive (XR): For X-linked recessive disorders, the genetic variant is found on the X chromosome. Female cats must inherit two copies of the variant to be at risk of developing the condition, whereas male cats only need one copy to be at risk. Males and females with any copies of the variant may pass the disorder-associated variant to their kittens if bred.

Modifier (MO): Genetic modifiers do not cause disease on their own but can cause disease or change the onset or severity of a disorder when combined with another disorder-associated variant. For some modifier variants only one copy is required to cause an effect, for others two copies are required. Please refer to the associated variant's breeder recommendations regarding safe breeding practices for each modifier variant.