

# Amasing Star's Hakon Of Milo/FI (HOW...

Birth date: 2024-11-28

Test date: 2026-01-13

ID kit: FKFHSWFXQQ

## Amasing Star's Hakon Of Milo/FI (HOWDY)'s Profile

### Pet information

**Registered name**

Amasing Star's Hakon Of Milo/FI (HOWDY)

**Sex**

M

**Date of birth**

2024-11-28

### Genetic Diversity

**Amasing Star's Hakon Of Milo/FI ...'s Percentage of Heterozygosity**

35%

### Health summary

At Risk 0 conditions

Carrier 0 conditions

Clear 50 conditions

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## Genetic Diversity

### Heterozygosity

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#### Amasing Star's Hakon Of Milo/FI ...'s Percentage of Heterozygosity

35%

Amasing Star's Hakon Of Milo/FI (HOWDY)'s genome analysis shows an average level of genetic heterozygosity when compared with other random-bred cats.

#### Typical Range for Random-Bred Cats

29% - 41%

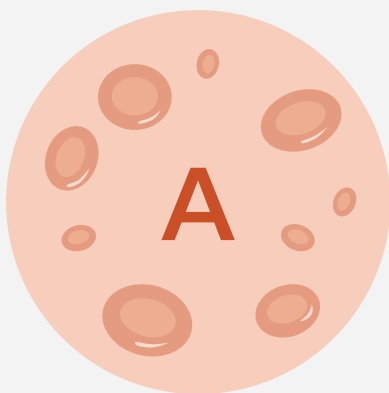
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## Blood Type



### Blood type

Type A (Most common)

### Genotype\*

A/A

### Transfusion risk

⚠ Moderate

Amasing Star's Hakon Of Milo/FI (HOWDY) has the most common blood type. He can be transfused with Type A blood.

### Blood variants tested\*

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

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## Interpreting feline blood types

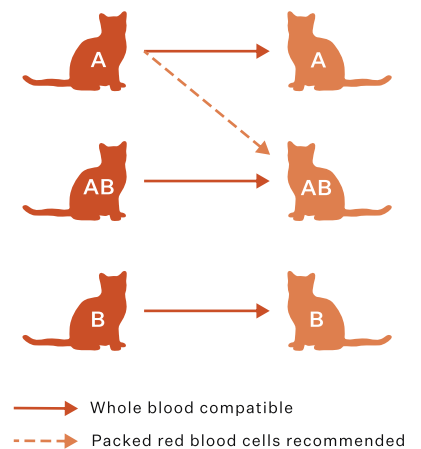
### About blood type determination

The three important feline blood types of A, B, and AB are governed primarily by variants in the CMAH gene. A cat's blood type can be determined by its genotype, which consists of two gene variants – one inherited from each parent – that should be interpreted together. When determining blood type based on genotype, the A variant associated with blood type A is most dominant while the b variants associated with blood type B are most recessive. The c variant associated with blood type AB is intermediate between the A and b variants, meaning it is recessive to the A variant but dominant to b variants. Therefore, a genotype with at least one A variant will result in blood type A. For a cat to have blood type B, the genotype must consist of two b variants. Because the c variant is intermediate, a cat with blood type AB can either have a genotype consisting of two c variants or one c variant and one b variant.

### About transfusion risk

Similar to humans, the different cat blood types will express different antigens on the surface of their red blood cells. This is significant because both type A and B cats are born with antibodies against other blood cell antigens. Notably, type B cats have high levels of antibodies against type A antigens. Cats with the rare blood type AB are most versatile as they express both red cell antigen types and, thus, can receive both type A and type AB blood transfusions.

Unlike humans, there is no cat blood type that can act as a universal blood donor. If a cat receives a non-compatible blood type during a transfusion, it may cause a severe, life-threatening reaction including fever, kidney failure, and widespread destruction of red blood cells. Prior to all transfusions, cats should be serologically typed and crossmatched to ensure compatibility.



### About breeding risk

During pregnancy, kittens are shielded from their mother's immune system. However, when kittens begin nursing, they receive some of their mother's antibodies in colostrum. Type B cats have high levels of antibodies against type A blood, so when blood type A or AB kittens are born to a blood type B mother, these antibodies, when absorbed by the newborn kitten, cause neonatal isoerythrolysis, a potentially fatal destruction of the kitten's red blood cells. Kittens of type B mothers with fathers of unknown or type A blood should be bottle fed or foster-nursed, and separated from their mother for the first 24 hours to avoid this reaction, unless blood typing performed immediately following birth shows the kitten to have a compatible blood type to the mother.

Although some blood types are less common and require additional planning when breeding, they represent normal genetic variation and should not be selected against when choosing breeding pairs.

### Current limits of this test

This test identifies 4 variants ( b variants c.269T>A, c.179G>T, c.1233delT and c variant c.346C>T) in the CMAH gene discovered in the domestic cat population and has been confirmed 99% concordant with serologic blood typing<sup>1</sup>. Mik antigens also play a role in blood type compatibility, and are not included in this test. Cats carrying undetermined, new, or undiscovered variants in CMAH or other genes may have a different blood type compatibility than that reported by this test. Accuracy of this test at predicting blood type in wildcats or wildcat hybrid breeds has not been determined.

1. Anderson H, Davison S, Lytle KM, Honkanen L, et al. Genetic epidemiology of blood type, disease and trait variants, and genome-wide genetic diversity in over 11,000 domestic cats (2022) PLOS Genetics.

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## Traits

### Coat Color

	Gene	Variant	Copies	Result
<b>Charcoal (Discovered in the Bengal)</b>	ASIP	A <sup>Pb</sup>	0	No effect
<b>Solid Color</b> <p>Two copies of the Solid Color variant are needed for a cat to have solid colored hair. However, orange coloration overrides this effect, meaning that cats with partial or full orange coats can show tabby patterning in orange areas. Cats with zero or one copy of this variant are likely to have a tabby pattern due to color banding of the hairs.</p>	ASIP	a	2	Solid color hairs likely
<b>Gloving (Discovered in the Birman)</b> <p>Two copies of the Gloving variant and Birman ancestry are needed for a cat to show white feet due to this variant. The Gloving variant can be found in non-Birman cats without having an effect on appearance and cats can have white feet without this variant suggesting other causes are yet to be discovered.</p>	KIT	w <sup>g</sup>	1	No effect
<b>Partial and Full White</b> <p>One or two copies of this variant will cause a part white or a full white appearance with blue coloration of one or both eyes possible.</p>	KIT	W or w <sup>s</sup>	1	Partly or fully white coat likely
<b>Amber (Discovered in the Norwegian Forest Cat)</b>	MC1R	e	0	No effect
<b>Russet (Discovered in the Burmese)</b>	MC1R	er	0	No effect
<b>Dilution</b> <p>Two copies of the Dilution variant are required to have a lightening effect on the coat.</p>	MLPH	d	2	Lightened coat color likely
<b>Albinism (Discovered in Oriental breeds)</b>	TYR	c <sup>a</sup>	0	No effect
<b>Colorpoint (Discovered in the Burmese)</b>	TYR	c <sup>b</sup>	0	No effect
<b>Colorpoint (Discovered in the Siamese)</b> <p>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.</p>	TYR	c <sup>s</sup>	2	Siamese colorpoint pattern likely

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## Coat Color

	Gene	Variant	Copies	Result
<b>Mocha (Discovered in the Burmese)</b>	TYR	c <sup>m</sup>	0	No effect
<b>Chocolate</b>	TYRP	b	0	No effect
<b>Cinnamon</b>	TYRP	b <sup>l</sup>	0	No effect

## Coat Type

	Gene	Variant	Copies	Result
<b>Long Hair (Discovered in many breeds)</b>	FGF5	M4	0	No effect
<b>Long Hair (Discovered in the Norwegian Forest Cat)</b>	FGF5	M2	0	No effect
<b>Long Hair (Discovered in the Ragdoll and Maine Coon)</b> Two copies of any Long Hair variant must be inherited for a cat to have a long coat. This can either be two copies of a particular variant, such as this one, or two of any combination of Long Hair variants.	FGF5	M3	2	Long coat likely
<b>Long Hair (Discovered in the Ragdoll)</b>	FGF5	M1	0	No effect
<b>Lykoi Coat (Variant 1)</b>	HR	hr <sup>Ca</sup>	0	No effect
<b>Lykoi Coat (Variant 2)</b>	HR	hr <sup>VA</sup>	0	No effect
<b>Hairlessness (Discovered in the Sphynx)</b>	KRT71	re <sup>hr</sup>	0	No effect
<b>Rexing (Discovered in the Devon Rex)</b>	KRT71	re <sup>dr</sup>	0	No effect
<b>Rexing (Discovered in the Cornish Rex and German Rex)</b>	LPAR6	r	0	No effect
<b>Glitter</b>	Pending	gl	0	No effect

## Tail Length

	Gene	Variant	Copies	Result
<b>Short Tail (Variant 3)</b>	HES7	jb	0	No effect
<b>Short Tail (Variant 1)</b>	T	C1199del	0	No effect

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## Tail Length

	Gene	Variant	Copies	Result
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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## 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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## Health conditions tested

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

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## Health conditions tested

Genetic Condition	Gene	Risk Variant	Copies	Inheritance	Result
<b>Progressive Retinal Atrophy (Discovered in the Abyssinian)</b>	CEP290	T>G	0	AR	Clear
<b>Progressive Retinal Atrophy (Discovered in the Bengal)</b>	KIF3B	G>A	0	AR	Clear
<b>Progressive Retinal Atrophy (Discovered in the Persian)</b>	AIP1	C>T	0	AR	Clear
<b>Pyruvate Kinase Deficiency</b>	PKLR	G>A	0	AR	Clear
<b>Sphingomyelinosis (Variant 1)</b>	NPC1	G>C	0	AR	Clear
<b>Sphingomyelinosis (Variant 2)</b>	NPC2	G>A	0	AR	Clear
<b>Spinal Muscular Atrophy (Discovered in the Maine Coon)</b>	LIX1	Deletion	0	AR	Clear
<b>Vitamin D-Dependent Rickets</b>	CYP27B1	G>T	0	AR	Clear

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## Glossary of genetic terms

### Test result definitions

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**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

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**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.