Registration: N/A

### 

### **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

Owner Info

First Name Charles **Last Name** Warden

Pet Info

**Registered Name** 

Bastu

Nickname (Call Name)

Bastu

Sex

Female

Date of Birth

5/1/2011

Sample ID

CMGWLZC

Registration

N/A

Microchip ID

N/A

Tattoo ID

N/A

Registration: N/A

### 

### **DNA Test Report**

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Complete

### **Ancestry Results**

#### Western

46% American Domestic Cat

14% Maine Coon

13% American Curl

12% Sphynx

9% Norwegian Forest Cat

5% Siberian

1% Highlander

Registration: N/A

## **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Genetic Diversity (Heterozygosity)

#### Bastu's Percentage of Heterozygosity

41%

Bastu's genome analysis shows an average level of genetic heterozygosity when compared with other random-bred cats.

**Typical Range for Domestic Cats** 

29 - 41%

Bastu Registration: N/A

# **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Health Conditions Tested

Genetic Condition	Gene	Risk Variant	Copies	Result
Acute Intermittent Porphyria (Variant 1)	HMBS	Deletion	0	Clear
Acute Intermittent Porphyria (Variant 2)	HMBS	G>A	0	Clear
Acute Intermittent Porphyria (Variant 3)	HMBS	Insertion	0	Clear
Acute Intermittent Porphyria (Variant 4)	HMBS	Deletion	0	Clear
Acute Intermittent Porphyria (Variant 5)	HMBS	G>A	0	Clear
Autoimmune Lymphoproliferative Syndrome	FASL	Insertion	0	Clear
Burmese Head Defect (Discovered in the Burmese)	ALX1	Deletion	0	Clear
Chediak-Higashi Syndrome (Discovered in the Persian)	LYST	Insertion	0	Clear
Congenital Adrenal Hyperplasia	CYP11B1	G>A	0	Clear
Congenital Erythropoietic Porphyria	UROS	G>A	0	Clear
Congenital Myasthenic Syndrome (Discovered in the Devon Rex and Sphynx)	COLQ	G>A	0	Clear
Cystinuria Type 1A	SCL3A1	C>T	0	Clear
Cystinuria Type B (Variant 1)	SCL7A9	C>T	0	Clear
Cystinuria Type B (Variant 2)	SCL7A9	G>A	0	Clear
Cystinuria Type B (Variant 3)	SCL7A9	T>A	0	Clear
Dihydropyrimidinase Deficiency	DPYS	G>A	0	Clear
Earfold and Osteochondrodysplasia (Discovered in the Scottish Fold)	TRPV4	G>T	0	Clear
Factor XII Deficiency (Variant 2)	F12	Deletion	0	Clear
Familial Episodic Hypokalemic Polymyopathy (Discovered in the Burmese)	WNK4	C>T	0	Clear
Glutaric Aciduria Type II	ETFDH	T>G	0	Clear
Glycogen Storage Disease (Discovered in the Norwegian Forest Cat)	GBE1	Insertion	0	Clear
GM1 Gangliosidosis	GLB1	G>C	0	Clear

Bastu Registration: N/A

# **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Health Conditions Tested (continued)

Genetic Condition	Gene	Risk Variant	Copies	Result
GM2 Gangliosidosis	GM2A	Deletion	0	Clear
GM2 Gangliosidosis Type II (Discovered in Domestic Shorthair cats)	HEXB	Insertion	0	Clear
GM2 Gangliosidosis Type II (Discovered in Japanese domestic cats)	HEXB	C>T	0	Clear
GM2 Gangliosidosis Type II (Discovered in the Burmese)	HEXB	0>0	0	Clear
Hemophilia B (Variant 1)	F9	C>T	0	Clear
Hemophilia B (Variant 2)	F9	G>A	0	Clear
Hyperoxaluria Type II	GRHPR	G>A	0	Clear
Hypertrophic Cardiomyopathy (Discovered in the Maine Coon)	MYBPC	G>C	0	Clear
Hypertrophic Cardiomyopathy (Discovered in the Ragdoll)	MYBPC	C>T	0	Clear
Hypotrichosis (Discovered in the Birman)	FOXN1	Deletion	0	Clear
Lipoprotein Lipase Deficiency	LPL	G>A	0	Clear
MDR1 Medication Sensitivity	ABCB1	Deletion	0	Clear
Mucopolysaccharidosis Type I	IDUA	Deletion	0	Clear
Mucopolysaccharidosis Type VI	ARSB	T>C	0	Clear
Mucopolysaccharidosis Type VI Modifier	ARSB	G>A	0	Clear
Mucopolysaccharidosis Type VII (Variant 1)	GUSB	G>A	0	Clear
Mucopolysaccharidosis Type VII (Variant 2)	USB	C>T	0	Clear
Myotonia Congenita	CLCN1	G>T	0	Clear
Polycystic Kidney Disease (PKD)	PKD1	C>A	0	Clear
Progressive Retinal Atrophy (Discovered in the Abyssinian)	CEP290	T>G	0	Clear
Progressive Retinal Atrophy (Discovered in the Bengal)	KIF3B	G>A	0	Clear
Progressive Retinal Atrophy (Discovered in the Persian)	AIPL1	C>T	0	Clear

Bastu Registration: N/A

**DNA Test Report** 

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Health Conditions Tested (continued)

Genetic Condition	Gene	Risk Variant	Copies	Result
Pyruvate Kinase Deficiency	PKLR	G>A	0	Clear
Sphingomyelinosis (Variant 1)	NPC1	G>C	0	Clear
Sphingomyelinosis (Variant 2)	NPC2	G>A	0	Clear
Vitamin D-Dependent Rickets	CYP27B1	G>T	0	Clear
Factor XII Deficiency (Variant 1)	F12	Deletion	_	Inconclusive

Registration: N/A

## **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Blood Type

 Blood Type
 Genotype

 A
 A/A

(Most common)

Transfusion Risk Breeding Risk

Moderate

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

Registration: N/A

# **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

#### **Coat Color**

Genetic Trait	Gene	Variant	Copies	Result
Charcoal (Discovered in the Bengal)	ASIP	APb	0	No effect
Solid Color	ASIP	а	2	Solid color hairs likely
Partial and Full White	KIT	Worw <sup>s</sup>	0	No effect
Amber (Discovered in the Norwegian Forest Cat)	MC1R	е	0	No effect
Russet (Discovered in the Burmese)	MC1R	e <sup>r</sup>	0	No effect
Dilution	MLPH	d	1	No effect
Albinism (Discovered in Oriental breeds)	TYR	c <sup>a</sup>	0	No effect
Colorpoint (Discovered in the Burmese)	TYR	c <sup>b</sup>	0	No effect
Colorpoint (Discovered in the Siamese)	TYR	c°	0	No effect
Mocha (Discovered in the Burmese)	TYR	c <sup>m</sup>	0	No effect
Chocolate	TYRP	b	0	No effect
Cinnamon	TYRP	b¹	0	No effect

### Coat Type

Genetic Trait	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	МЗ	1	Long coat possible, short coat likely
Long Hair (Discovered in the Ragdoll)	FGF5	M1	0	No effect
Lykoi Coat (Variant 1)	HR	hr <sup>Ca</sup>	0	No effect
Lykoi Coat (Variant 2)	HR	hr <sup>VA</sup>	0	No effect
Hairlessness (Discovered in the Sphynx)	KRT71	re <sup>hr</sup>	0	No effect

Registration: N/A

# **DNA Test Report**

Sample ID: CMGWLZC Test Date: 7/6/2021

Complete

### Coat Type (continued)

Genetic Trait	Gene	Variant	Copies	Result
Rexing (Discovered in the Devon Rex)	KRT71	re <sup>dr</sup>	0	No effect
Rexing (Discovered in the Cornish Rex and German Rex)	LPAR6	r	0	No effect
Glitter	Pending	gl	0	No effect