

CatCheck test

List of approved analyses that are included in our cat test

Disease	Gene	Breed if applicable
Acute Intermittent Porphyria, Version 1	HMBS	-
Acute Intermittent Porphyria, Version 3	HMBS	-
Acute Intermittent Porphyria, Version 4, Siamese Type 1	HMBS	Siamés
Acute Intermittent Porphyria, Version 5, Siamese Type 2	HMBS	Siamés
A-Locus Agouti	ASIP	-
Alpha-Mannosidosis	MAN2B1	Persa
Autoimmune Lymphoproliferative Syndrome	FASLG	British Shorthair
B-Locus, Chocolate	b	-
B-Locus, Cinnamon	b1	-
Blood Group System ABC	CMAH, ac	Ragdoll
Blood Group System ABC	CMAH, b1	-
Blood Group System ABC	CMAH, b2	-
C-Locus, Complete Albinism	TYR	(male)
Cystinuria 1a	SLC3A1	-
Cystinuria Bv1	SLC7A9	-
Cystinuria Bv2	SLC7A9	Maine Coon, Sphynx, Siamés
Cystinuria Bv3	SLC7A9	-
Cystinuria Bv4	SLC7A9	-
Dihydropyrimidinase Deficiency	DPYS	-
D-Locus, Dilution	MLPH	-
E-Locus, Amber and Russet	e	Norwegian Forest Cat
E-Locus, Amber and Russet	er	Burmese
Epidermolysis Bullosa Simplex	KRT14	(male)
Erythropoietic Porphyria, Congenital Variant 1	-	-
Factor XII Deficiency, Version 1	-	(male)
Factor XII Deficiency, Version 2	-	-
Forebrain Commissure Malformation	GDF7	-
Gangliosidosis GM1	GLB1	Korat, Siamés

Gangliosidosis GM2	HEXB	Burmese
Gangliosidosis GM2	HEXB	Japonés
Folded Ears with Osteochondrodysplasia	TRPV4	Scottish Fold
Gender	ZFXY	-
Hr-Locus, Hairlessness	-	-
Hypertrophic Cardiomyopathy	ALMS1	Sphynx
Hypertrophic Cardiomyopathy	MYBPC3	Maine Coon (male)
Hypertrophic Cardiomyopathy	MYBPC3	Ragdoll
Hypertrophic Cardiomyopathy	MYH7	Gato doméstico
Hypokalemic Periodic Paralysis	WNK4	Burmese
Hypotrichosis with Short Life Expectancy	FOXN1	Birman
L-Locus, Long Hair	I1	Ragdoll
L-Locus, Long Hair	I3	Ragdoll, Maine Coon (male)
L-Locus, Long Hair	I2	Norwegian Forest Cat
L-Locus, Long Hair	I4	-
Multi-Drug Resistance	ABCB1	-
Myasthenic Syndrome, Congenital Muscular Dystrophy	COLQ	Devon Rex, Sphynx
Polycystic Kidney Disease	PKD1	-
Primary Congenital Glaucoma	LTBP2	(male)
Progressive Retinal Atrophy	KIF3B	Bengal
Progressive Retinal Atrophy, Leber Congenital Amaurosis	AIPL1	Persa
Progressive Retinal Atrophy, Retinal Degeneration II	CEP290	Abisinio
Pyruvate Kinase Deficiency of Erythrocytes	PKLR	-
Vitamin D-Dependent Rickets Type IA Variant 1	CYP27B1	-
White Gloves	KIT	Birman