Kategori	Sjukdomar, Pälsfärg, Pälsstruktur & Kroppsegenskaper (eng)
Egenskap	A-locus agouti, ASIP, solid
Egenskap	ABC Blood group system, CMAH, ac, Ragdoll
Egenskap	ABC Blood group system, CMAH, b1
Egenskap	ABC Blood group system, CMAH, b2
Sjukdom	Acute intermittent porphyria, version 1, HMBS
Sjukdom	Acute intermittent porphyria, version 3, HMBS
Sjukdom	Acute intermittent porphyria, version 4, HMBS, Siamese type 1
Sjukdom	Acute intermittent porphyria, version 5, HMBS, Siamese type 2
Sjukdom	Alpha-mannosidosis, MAN2B1, Persian
Sjukdom	Autoimmune lymphoproliferative syndrome, FASLG, British Shorthair
Egenskap	B-locus, chocolate (b)
Egenskap	B-locus, cinnamon (b1)
Egenskap	C-locus, complete albinism, c, TYR (x-linked)
Sjukdom	Cystinuria 1a, SLC3A1
Sjukdom	Cystinuria Bv1, SLC7A9
Sjukdom	Cystinuria Bv2, SLC7A9, Maine Coon, Sphynx, Siamese
Sjukdom	Cystinuria Bv3, SLC7A9
Sjukdom	Cystinuria Bv4, SLC7A9
Egenskap	D-locus, dilution, MLPH
Sjukdom	Dihydropyrimidinase deficiency, DPYS
Egenskap	E-Locus, Amber and Russet, e, Norwegian Forest Cat
Egenskap	E-locus, Amber and Russet, er, Burmese
Sjukdom	Epidermolysis bullosa simplex, KRT14 (x-linked)
Sjukdom	Erythropoietic porphyria, congenital variant 1
Sjukdom	Factor XII deficiency, version 1 (x-linked)
Sjukdom	Factor XII deficiency, version 2
Egenskap	Folded ears with osteochondrodysplasia, TRPV4, Scottish Fold
Sjukdom	Gangliosidosis GM1, GLB1, Korat, Siamese
Sjukdom	Gangliosidosis GM2, HEXB, Burmese
Egenskap	Gangliosidosis GM2, HEXB, Japanese
Egenskap	Hr-locus, Hairlessness
Sjukdom	Hypertrophic cardiomyopathy, ALMS1, Sphynx
Sjukdom	Hypertrophic cardiomyopathy, MYBPC3, Maine Coon (x-linked)
Sjukdom	Hypertrophic cardiomyopathy, MYBPC3, Ragdoll

Sjukdom	Hypertrophic cardiomyopathy, MYH7, Domestic cat
Sjukdom	Hypokalemic periodic paralysis, WNK4, Burmese
Sjukdom	Hypotrichosis with short life expectancy, FOXN1, Burmese
Egenskap	L-Locus, Longhair, l1, Ragdoll
Egenskap	L-Locus, Longhair, l2, Norwegian Forest Cat
Egenskap	L-Locus, Longhair, l3, Ragdoll, Maine Coon (x-linked)
Egenskap	L-Locus, Longhair, l4
Sjukdom	Malformation of the forebrain commissure, GDF7
Sjukdom	Multiresistance, ABCB1
Egenskap	Myasthenic syndrome, congenital muscular dystrophy, COLQ, Devon Rex, Sphynx
Sjukdom	Polycystic kidney disease, PKD1
Sjukdom	Primary congenital glaucoma, LTBP2 (x-linked)
Sjukdom	Progressive retinal atrophy, KIF3B, Bengal
Sjukdom	Progressive retinal atrophy, Leber congenital amaurosis, AIPL1, Persian
Sjukdom	Progressive retinal atrophy, retinal degeneration II, CEP290, Abyssinian
Sjukdom	Pyruvate kinase deficiency of erythrocytes, PKLR
Egenskap	Sex, ZFXY
Sjukdom	Vitamin D-dependent rickets type IA variant 1, CYP27B1
Egenskap	White gloves, KIT, Birman