

Table 1 Phenotypic traits of the domestic cat conferred by DNA variants

Locus (Alleles) OMIA entry link	MOI*	Phenotype	Gene	Gene name	Mutation
<i>Agouti (A⁺, a)⁵ 000201-9685</i>	AR	Banded fur to solid	ASIP	<i>Agouti-signaling protein</i>	c.122_123delCA
<i>Brown (B⁺, b, b')^{6,7} 001249-9685</i>	AR	Brown, light brown color variants	TYRP1	<i>Tyrosinase-related protein</i>	b' = -5IVS6 b' = c.298C>T
<i>Color (C⁺, C^b, C^s, c)⁷⁻⁹ 000202-9685</i>	AR	Burmese, Siamese color pattern, full albino	TYR	<i>Tyrosinase</i>	c ^b = c.715G>T c ^s = c.940G>A c = c.975delC
<i>Dilution (D⁺, d)¹⁰ 000206-9685</i>	AR	Black to grey/blue, orange to cream	MLPH	<i>Melanophillin</i>	c.83delT
<i>Dwarfism 000299-9685</i>	AD	Shortening of long bones	unknown	unknown	unknown
<i>Extension (E⁺, e) – Amber¹¹ 001199-9685</i>	AR	Brown/red color variant	MC1R	<i>Melanocortin receptor 1</i>	c.250G>A
<i>Fold (Fd, fd') 000319-9685</i>	AD	Ventral ear fold	unpublished	unpublished	unpublished
<i>Gloves (G⁺, g)¹² 001580-9685</i>	AR	White feet	KIT	<i>KIT</i>	c.1035_1036delinsCA
<i>Hairless (Hr⁺, hr)¹³ 001584-9685</i>	AR	Atrichia	KRT71	<i>Keratin 71</i>	c.816+1G>A
<i>Inhibitor (I, i') 001584-9685</i>	AD	Absence of pheomelanin	unknown	unknown	unknown
<i>Japanese Bobtail (J, j') 000244-9685</i>	AD	Kinked tail	unknown	unknown	unknown
<i>Kurl (K, k') 000245-9685</i>	AD	Rostral curled pinnae	unknown	unknown	unknown
<i>LaPerm 000245-9685</i>	AD	Curly hair coat	unknown	unknown	unknown
<i>Longhair (L⁺, l)^{14,15} 000439-9685</i>	AR	Long fur	FGF5	<i>Fibroblast growth factor 5</i>	c.356_367insT c.406C>T c.474delT c.475A>C
<i>Tailless (Manx) (M, m')¹⁶ 000975-9685</i>	AD	Absent/short tail	TBOX	<i>T-box</i>	c.998delT c.1169delC c.1199delC c.998_1014dup17delGCC
<i>Orange (O, o') 001201-9685</i>	X-linked	Change in pigment hue	unknown	unknown	unknown
<i>Peterbald 001866-9685</i>	AD	Hairless, brush coat	unknown	unknown	unknown
<i>Polydactyla (Pd, pd')¹⁷ 000810-9685</i>	AD	Extra toes	SHH	<i>Sonic hedgehog</i>	c.479A>G c.257G>C c.481A>T
<i>Rexing (R⁺, r)¹⁸ 001684-9685</i>	AR	Curly hair coat	LPAR6	<i>Lysophosphatidic acid receptor 6</i>	c.250_253delTTTG
<i>Rexing (Re⁺, re)¹³ 001581-9685</i>	AR	Curly hair coat	KRT71	<i>Keratin 71</i>	c.1108-4_1184del, c.1184_1185insAGTTGGAG c.1196insT
<i>Rexing (R^s, r^{s+})¹⁹ 001712-9685</i>	AD	Curly hair coat	KRT71	<i>Keratin 71</i>	c.445-1G>C
<i>Spotting (S, s')²⁰ 000214-9685</i>	Co-D	Bicolor/Van white	KIT	<i>KIT</i>	7125ins FERV1 element
<i>Tabby (T^M, t')²¹ 001429-9685</i>	AR	Blotched/classic pattern	TAQPEP	<i>Transmembrane aminopeptidase Q</i>	c.176C>A c.416C>A c.682C>A c.2522G>A
<i>Ticked (T^R, t) 001484-9685</i>	AD	No tabby pattern	unknown	unknown	unknown
<i>White (W, w')²⁰ 000209-9685</i>	AD	Loss of pigmentation	KIT	<i>KIT</i>	FERV1 LTR ins
Wide-band	AR?	Length of pheomelanin band in hair	unknown	unknown	unknown

*Mode of inheritance of the non-wild type variant. '+' implies the wild type allele when known. In reference to the mutant allele, AD = autosomal dominant, AR = autosomal recessive, co-D = co-dominant.

OMIA: Online Mendelian Inheritance in Animals (omia.angis.org.au) entries provide links to citations and clinical descriptions of the phenotypes and the diseases. Listed citations are for the causative variant discovery