Domestic

LABGENVET (2021)

Pyruvate Kinase Deficiency
(38 tests)

N/N	92.1%
M/N	7.9%
M/M	0.0%

Anderson (2022)

Pyruvate Kinase Deficiency
(607 tests)

N/N	97.5%
M/N	2.5%
M/M	0.0%

Factor XII Deficiency (Variant 2)
(90 tests)

N/N	76.7%
M/N	23.3%
M/M	0.0%

Factor XII Deficiency (Variant 1)
(606 tests)

N/N	95.2%
M/N	4.8%
M/M	0.0%

Progressive Retinal Atrophy (Abyssinian)
(PRA-rdAc) (617 tests)

N/N	97.7%
M/N	2.3%
M/M	0.0%

Progressive Retinal Atrophy (Bengal)
(607 tests)

N/N	99.8%
M/N	0.2%
M/M	0.0%

Hyperoxaluria Type II
(617 tests)

N/N	98.7%

M/N	1.3%
M/M	0.0%

Congenital Erythropoeietic Porphyria (c.140C>T)
(617 tests)

N/N	99.0%
M/N	0.8%
M/M	0.2%

Earfold, Osteochondrodysplasia (Scottish Fold)
(617 tests)

N/N	99.0%
M/N	0.8%
M/M	0.2%

Lipoprotein Lipase Deficiency
(617 tests)

N/N	99.4%
M/N	0.3%
M/M	0.0%

MDR1 Medication Sensitivity
(617 tests)

N/N	99.1%
M/N	0.9%
M/M	0.0%

Mucopolysaccharidosis Type VI Modifier
(617 tests)

N/N	99.7%
M/N	0.3%
M/M	0.0%

Hypertrophic Cardiomyopathy
(Maine Coon) (617 tests)

N/N	99.7%
M/N	0.3%
M/M	0.0%

Hypertrophic Cardiomyopathy
(Ragdoll) (617 tests)

N/N	99.8%
M/N	0.2%

M/M	0.0%

Myotonia Congenita

(617 tests)

N/N	99.8%
M/N	0.0%
M/M	0.2%

GM2 Gangliosidosis Type II (Burmese)

(617 tests)

N/N	99.8%
M/N	0.2%
M/M	0.0%