

DNA Test Report

Dog Information

Margo (Margo) Female NAME SEX

Labrador Retriever June 27th, 2023
BREED ANCESTRY DATE OF BIRTH

American Kennel Club (AKC) n/a
REGISTRATION MICROCHIP

Brittany N Pescara
OWNER NAME

Canine Genetic Health Screen

TEST

December 5th, 2023

TEST DATE

BREED HEALTH TESTS

Centronuclear Myopathy, CNM PTPLA NN Clear ♥ Degenerative Myelopathy, DM SOD1A GG Clear ♥ Exercise-Induced Collapse, EIC DNM1 GG Clear ♥ Hereditary Nasal Parakeratosis, HNPK SUV39H2 GG Clear ♥ Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU SLC2A9 (Exon 5) GG Clear ♥ Macular Corneal Dystrophy, MCD CHST6 CC Clear ♥ Narcolepsy HCRTR2 AA Clear ♥ Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear ♥ Pyruvate Kinase Deficiency PKLR Exon 7 SNP Variant 1 CC Clear ♥ Skeletal Dysplasia 2, SD2 COL11A2 GG Clear ♥ Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear ★ Alexander Disease GFAP (Exon 4) GG Clear ★ Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear ★ Colden Retriever Progressive Retinal Atrophy 2, GR-PRA2 TTGS Exon 8 NN Clea	DISEASE	GENE	GENOTYPE	RESULT	TESTING RECOMMENDED BY
Exercise-Induced Collapse, EIC Hereditary Nasal Parakeratosis, HNPK SUV39H2 GG Clear Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU Macular Corneal Dystrophy, MCD CHST6 CC Clear Narcolepsy HCRTR2 AA Clear Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear CHST6 CC Clear Conjenital Myasthenic Syndrome, CMS	Centronuclear Myopathy, CNM	PTPLA	NN	Clear	•
Hereditary Nasal Parakeratosis, HNPK SUV39H2 GG Clear Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU SLC2A9 (Exon 5) GG Clear Macular Corneal Dystrophy, MCD CHST6 CC Clear Narcolepsy HCRTR2 AA Clear Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear CC Clear Clear CO Clear Clear CO Clear Clear CO CO CO Clear CO CO CO CO CO Cl	Degenerative Myelopathy, DM	SOD1A	GG	Clear	•
Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU SLC2A9 (Exon 5) GG Clear CC Clear Macular Corneal Dystrophy, MCD HCRTR2 AA Clear Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear CC Clear CO Pyruvate Kinase Deficiency PKLR Exon 7 SNP Variant 1 CC Clear CO Skeletal Dysplasia 2, SD2 Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear Alexander Disease GFAP (Exon 4) GG Clear CO CO Clear CO CO Clear CO CO CO CO CO CO CO CO CO C	Exercise-Induced Collapse, EIC	DNM1	GG	Clear	•
Macular Corneal Dystrophy, MCD CHST6 CC Clear Marcolepsy HCRTR2 AA Clear Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear Cl	Hereditary Nasal Parakeratosis, HNPK	SUV39H2	GG	Clear	•
Narcolepsy HCRTR2 AA Clear Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear CC Clear COL11A2 COL11A2 COL11A2 COL11A2 COL2 Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear Alexander Disease GFAP (Exon 4) GG Clear COL2 COL2 COL3 C	Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU	SLC2A9 (Exon 5)	GG	Clear	•
Progressive Retinal Atrophy, prcd PRCD Exon 1 GG Clear CC Clear COL11A2 GG Clear COL11A2 CNGA3 (Exon 7 Deletion) NN Clear Achromatopsia CNGA3 (Exon 4) GG Clear COL2 COL3 COL	Macular Corneal Dystrophy, MCD	CHST6	CC	Clear	•
Pyruvate Kinase Deficiency PKLR Exon 7 SNP Variant 1 CC Clear COL11A2 GG Clear Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear Alexander Disease GFAP (Exon 4) GG Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Narcolepsy	HCRTR2	AA	Clear	•
Skeletal Dysplasia 2, SD2 Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear Alexander Disease GFAP (Exon 4) GG Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Progressive Retinal Atrophy, prcd	PRCD Exon 1	GG	Clear	•
Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear Alexander Disease GFAP (Exon 4) GG Clear Canine Elliptocytosis SPTB Exon 30 CC Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Pyruvate Kinase Deficiency	PKLR Exon 7 SNP Variant 1	CC	Clear	•
Alexander Disease GFAP (Exon 4) GG Clear Canine Elliptocytosis SPTB Exon 30 CC Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Skeletal Dysplasia 2, SD2	COL11A2	GG	Clear	•
Canine Elliptocytosis SPTB Exon 30 CC Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Achromatopsia	CNGA3 (Exon 7 Deletion)	NN	Clear	x
Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Alexander Disease	GFAP (Exon 4)	GG	Clear	x
	Canine Elliptocytosis	SPTB Exon 30	СС	Clear	x
Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2 TTC8 Exon 8 NN Clear	Congenital Myasthenic Syndrome, CMS	COLQ (Exon 14)	TT	Clear	x
	Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2	TTC8 Exon 8	NN	Clear	x





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BREED HEALTH TESTS

DISEASE	GENE	GENOTYPE	RESULT	TESTING RECOMMENDED BY
Myotubular Myopathy 1, X-linked Myotubular Myopathy, XL-MTM	MTM1 (Exon 7)	cc	Clear	솼
Progressive Retinal Atrophy, crd4/cord1	RPGRIP1 (Exon 2)	NN	Clear	x
Stargardt Disease	ABCA4 Exon 28	NI	1 Variant	ᅪ
Ullrich-like Congenital Muscular Dystrophy	COL6A3	GG	Clear	x





Canine Genetic Health Certificate™

Call Name: Margo Laboratory #: 412367

Registered Name: - Registration #:

Breed: Labrador Retriever Certificate Date: Dec. 8, 2023

Sex: Female **DOB:** June 2023

This canine's DNA showed the following genotype(s):

Disease	Gene	Genotype	Interpretation
Retinal Dysplasia/Oculoskeletal Dysplasia 1	COL9A3	WT/WT	Normal (Clear)

WT, wild type (normal); M, mutant; Y, Y chromosome (male)

Paw Print Genetics® performed the testing on the dog listed on this certificate. See the Laboratory Report for interpretation and recommendations based on these findings. The genes/diseases reported here were selected by the client. Normal results do not exclude inherited mutations not tested in these or other genes that may cause medical problems or may be passed on to offspring. The results included in this report relate only to the items tested using the sample provided. These tests were developed and their performance determined by Paw Print Genetics. This laboratory has established and verified the test(s)' accuracy and precision with >99.9% sensitivity and specificity. The presence of mosaicism may not be detected by this test. Non-paternity may lead to unexpected results. This is not a breed identification test. Because all tests performed are DNA-based, rare genomic variations may interfere with the performance of some tests producing false results. If you think these results are in error, please contact the laboratory immediately for further evaluation. In the event of a valid dispute of results claim, Paw Print Genetics will do its best to resolve such a claim to the customer's satisfaction. If no resolution is possible after investigation by Paw Print Genetics with the cooperation of the customer, the extent of the customer's sole remedy is a refund of the fee paid. In no event shall Paw Print Genetics be liable for indirect, consequential or incidental damages of any kind. Any claim must be asserted within 60 days of the report of the test results. Genetic counseling is available at Paw Print Genetics.