LABOKLIN
LABORATORY FOR CLINICAL DIAGNOSTICS

Test Result

Sample ID: 2007-W-33091
Result Date: 06/07/2020

Veterinary Surgeon

Owner

MS BOZENA BIENKOWSKA
FORESIDE FARM, FERENEZE ROAD
GLASGOW
EAST RENFREWSHIRE
G78 3AQ

bozenaandmicheal@aol.com

Animal Details

Animal:	CANINE	D.O.B	25/03/2015
Name	BERTGAN DAPHNE (DAPHNE)	Microchip No.	900200000955026
Breed	MINIATURE LABRADOODLE	KC Reg	
Sex	FEMALE	Tattoo No.	

Sample

Sample Material	SWAB	Sample received:	01/07/2020
Sample Date:	29/06/2020		

Test

Test Name:	8158D Degenerative Myelopathy (EXON 2)

Result

Genotype: N/N (exon 2) (Clear)

Interpretation: The dog is homozygous normal for the intact SOD1-gene.

The dog does not carry the mutation in exon 2 of the SOD1 that is suggested to be a major risk factor for the development of Degenerative Myelopathy.

Please note: In the Bernese Mountain Dog breed the mutation in exon 1 of SOD1-gene occurs also in correlation with DM.

The dog can pass only the normal gene on to all its offspring.

The currently known mutation has been analysed.

The result is only valid for the submitted sample.

********** END OF RESULT **********

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Sample

Sample Material	SWAB	Sample received:	01/07/2020
Sample Date:	29/06/2020		

Test

Test Name:	EXERCISE INDUCED COLLAPSE (EIC)

Result

Genotype:	N / N (Clear)
Interpretation:	The dog is homozygous for the wild type allele. The dog does not carry the EIC' causative mutation in the DNM1-gene. The dog is genetically clear and will not be affected by EIC (Exercise Induced Collapse). The dog can pass only the normal gene on to its entire offspring. The currently known mutation has been analysed. The result is only valid for the submitted sample and for the following breeds: Bouvier des Flanders, Boykin Spaniel, Chesepeake Bay Retriever, Clumber Spaniel, Cocker Spaniel, Curly Coated Retriever, German Wirehaired Pointer, Labrador Retriever, Old English Sheepdog and Pembroke Welsh Corgi.

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Sample

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Sample Date:	29/06/2020		

Test

Test Name:	8225D Hereditary Nasal Parakeratosis (HNPK)

Result

Genotype:	N/N (CLEAR)
Interpretation:	The examined dog is homozygote for the intact gene. This dog does not carry the mutation described to cause Hereditary Nasal Parakeratosis (HNPK). This dog will only pass the intact healthy gene on to its entire offspring. The result is only valid for the breed: Labrador Retriever. The result is valid for the submitted sample only

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Sample

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Sample Date:	29/06/2020		

Test

Test Name: 8235D Dwarfism /Skeletal Dysplasia 2(Osteochondrodysplasia)

Result

Genotype: N/N (Clear)

Interpretation: The examined dog is homozygote for the wild type allele.

This dog does not carry the SD2 causative mutation in the COL11A2 gene.

Trait of inheritance: autosomal-recessive

This dog will only pass the healthy gene onto its offspring.

The current result is only valid for the sample submitted to our laboratory.

The sender is responsible for the correct information regarding the sample material

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Test

Test Name:	8073 Centronuclear Myopathy (CNM)

Result

Genotype:	N/N (Clear)
Interpretation:	The examined animal is homozygous for the wild type allele. It does not carry the CNM causative mutation in the PTPLA-gene. Trait of inheritance: autosomal-recessive Scientific studies found correlation between the mutation and symptoms of the disease in the Labrador Retriever breed. Other forms of Myopathy cannot be excluded by this test.

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Sex	FEMALE	Tattoo No.	

Sample

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Sample Date:	29/06/2020		

Test

Test Name:	8094P prcd – PRA (Progressive Retinal Atrophy) *

Result

Genotype:	N/ N (Clear)
Interpretation:	The examined animal is homozygous for the wild type allele. It does not carry the causative mutation for prcd-PRA in the PRCD-gene. Trait of inheritance: autosomal-recessive Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Australian cattle dog, American Cocker Spaniel, American Eskimo Dog, Australian Shepherd, Australian Stumpy Tail Cattle Dog, Bolognese, Bolonka Zwetna, Chesapeake Bay Retriever, Chihuahua, Chinese Crested, English Cocker Spaniel, English Shepherd, Entlebucher Mountain Dog, Finnish Lapphund, German Spitz, Giant Schnauzer, Golden Retriever, Karelian Beardog, Kuvasz, Lagotto Romagnolo, Lapponian Herder, Labrador Retriever, Markiesje, Norwegian Elkhound, Nova Scotia Duck tolling Retriever, Portugese Water Dog, Poodle, Schipperke, Swedish Lapphound, Silky Terrier, Spanish Water Dog, Swedish Lapphund, Wäller, Yorkshire Terrier. The current result is only valid for the sample submitted to our laboratory. * test performed by partner lab

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Sample

Sample Material	SWAB	Sample received:	01/07/2020
Sample Date:	29/06/2020		

Test

Test Name: 8392 Retinal Dysplasia (RD)/ Oculo-Skeletal Dysplasia (OSD)*

Result

Genotype: N / N (Clear)

Interpretation: The examined animal is homozygous for the wild type allele. It does not carry the OSD causative mutation.

Trait of inheritance: autosomal-dominant.

Scientific studies found correlation between the mutation and symptoms of the disease in the Labrador Retriever breed.

The current result is only valid for the sample submitted to our laboratory.

* tested by partner lab

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