

LABOKLIN GmbH&CoKG . Postfach 1810 .DE-97688 Bad Kissingen

Mr.
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Report

No.: 1911-W-76039
Date of arrival: 06-11-2019
Testing started: 06-11-2019
Date of report: 07-11-2019
Testing completed: 07-11-2019

Patient identification:	Dog	Female	* 25.11.15
	Landseer		
Owner / Animal-ID:	Fors, Kare		
Type of sample:	EDTA-Blood		
Date sample was taken:	05-11-2019		

Name: **Cherry vom Graf Toggenburg**
ZB-Nummer: **SE21422/2016**
Chip-Nummer: **756093900027681**
Tattoo-Nummer: **---**

Cystinuria - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for Cystinuria in the SLC3A1-gene.

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Newfoundland, Landseer

Degenerative Myelopathy - PCR

Result: Genotype N/N (exon 2)

Interpretation: The examined animal is homozygous for the

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wildtype-allele. It does not carry the high-risk factor for DM in exon 2 of the SOD1-gene.

Trait of inheritance: autosomal-recessive

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

Hyperuricosuria - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for HUU in the SLC2A9-gene.

Trait of inheritance: autosomal-recessive

Muscular Dystrophy - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for MD in the COL6A1-gene.

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Landseer

Thrombopathia - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for thrombopathia in the Caldag-GEFI-gene.

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Landseer

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D-locus D1 (dilution)

Result: Genotype D/D

Interpretation: The examined animal does not possess the d1 allele. If no other d variant is present, the examined animal is homozygous for the D-allele.

The test detects the alleles D and d1
Allelic series: D dominant over d1

Please note: Additional d variants have to be considered to fully evaluate the characteristic of dilution.

Please note:

A further causative mutation for dilution (d2) has been found in the foll Chow Chow, Sloughi, Thai Ridgeback
The additional mutation might be responsible for dilution in further bree

Sampling:

The following impartial person (veterinarian, breed warden, or similar) signed the form for the sampling and identity check of the animal:

Ingrid Berger Engström

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. The laboratory can not be made liable. Furthermore, any obligation for compensation is limited to the value of the tests performed.

There is a possibility that other mutations may have caused the disease/phenotype. The analysis was performed according to the latest knowledge and technology.

The laboratory is accredited for the performed tests according to DIN EN ISO/IEC 17025:2005. (except partner lab tests).

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LABOKLIN
LABOR FÜR KLINISCHE DIAGNOSTIK GMBH & CO. KG

*** END of report ***

Fr. MSc Michelle Meißler
Abt. Molekularbiologie