

Västriku Loomakliinik OÜ
Västriku 2 A
11312 Tallinn / Harjumaa
Estonia

LABOKLIN GmbH&CoKG
Steubenstraße 4
DE-97688 Bad Kissingen
Fax-Nr.: +49 971 68546
Tel.: +49 971 72020

Report

No.: 2005-W-22227
Date of arrival: 13-05-2020
Testing started: 13-05-2020
Date of report: 27-05-2020
Testing completed: 27-05-2020

```
+-----+
| Patient identification: Dog           Female           * 30.01.2017 |
|                               Staffordshire Bull Terrier |
| Owner / Animal-ID:           Vilba, Kadri           |
| Type of sample:              EDTA-Blood             |
| Date sample was taken:       11-05-2020            |
+-----+
```

Parameter	Value	Reference value
-----------	-------	-----------------

Name:	Roxy (Grey Cardinal Madeira)	
-------	-------------------------------------	--

Stud book no.:	---	
----------------	-----	--

Chip no.:	981098106504611	
-----------	------------------------	--

Tattoo no.:	---	
-------------	-----	--

*Hereditary Cataract (HSF4) - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for cataract in the HSF4-gene.

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Boston Terrier, French Bulldog, Staffordshire Bull Terrier

L-2-Hydroxyglutaric Aciduria (L2HGA) - PCR

Result: Genotype N/N

Interpretation: The examined animal is homozygous for the wildtype-allele. It does not carry the causative mutation for L-2-HGA in the L2HGDH-gene.

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Staffordshire Bull Terrier

Referring vet: Kaysa Kokamägi (1132)

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).

These results are based on the sample material submitted to our laboratory.

This was suitable if not stated otherwise. The submitter is responsible for the accuracy of the information regarding the sample. This report can only be transmitted in toto and unchanged. Doing otherwise requires written permission from Laboklin GmbH & Co. KG.

*** END of report ***

Fr.Dipl.-Biol. Bärbel Gunreben
Abt. Molekularbiologie

*: test performed by partnerlaboratory