Emne **Joergensen, Keld Dog Labrador Retriever**
Afsender <befund@laboklin.de>
Modtager <hjpdyr@ofir.dk>
Dato 2018-01-15 12:52
Prioritet Normal

GODKENDT

LABOKLIN
LABOR FÜR KLINISCHE DIAGNOSTIK GMBH & CO. KG

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Report
No.: 1801-W-01253
Date of arrival: 11-01-2018
Date of report: 15-01-2018

Patient identification:	Dog	Male	* 11.07.15
	Labrador Retriever		
Owner / Animal-ID:	Joergensen, Keld		
Type of sample:	EDTA-Blood		
Date sample was taken:	04-01-2018		

Name: **Lochiness Clotworthy Dobbin**
Stud book no.: **DK 14443/2015**
Chip no.: **208210000540262**
Tattoo no.: ---

Hereditary myopathy (CNM) - PCR

Result: Genotype N/N

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

Trait of inheritance: autosomal-recessive

Scientific studies found correlation between the mutation and symptoms of the disease in the following breeds: Labrador Retriever
Other forms of myopathy cannot be excluded by this test.

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).

*** END of report ***

Hr.LM-Chemiker D. Schindelmann
Abt. Molekularbiologie