

Result certificate #012345

Detection of c.1288G>A ATG4D gene causing LSD in Lagotto Romagnolo

Customer: Jan Novák, Dlouhá 1, 30000 Plzeň, Czech Republic

Sample:

Sample: 21-12345 Date received: 01.02.2021 Sample type: blood

Information provided by the customer

Name: Lassie DEMO Breed: Plemeno

Tattoo number: 1392013 Microchip: 123 456 789 012 345 Reg. number: REGQ12345 Date of birth: 1.1.2020

Sex: female

Date of sampling: 01.02.2021

The identity of the animal has been checked.

Result: Mutation was not detected (N/N)

Legend: N/N = wild-type genotype. N/P = carrier of the mutation. P/P = mutated genotype (individual will be most probably affected with the disease). (N = negative, P = positive)

Explanation

Presence or absence of c.1288G>A mutation in ATG4D gene causing Lysosomal Storage Disease (LSD) in Lagotto Romagnolo was tested. LSD is a neurodegenerative disorder. The occurrence of a neurodegenerative disease is associated with malfunction of degradation processes and subsequent accumulation of various types of cellular materials in lysosomes of cell. Symptoms are progressive cerebellar ataxia, slight atrophy of cerebellum and forebrain, abnormal involuntary eye movement and behavioural ganges.

Mutation that causes LSD in Lagotto Romagnolo is inherited as an autosomal recessive trait. That means the disease affects dogs with P/P genotype only. The dogs with N/P genotype are considered carriers of the disease (heterozygotes), they are healthy but they can transmit the mutation on their offspring. In offspring of two heterozygous animals following genotype distribution can be expected: 25 % N/N (healthy non-carriers), 50 % N/P (healthy carriers) and 25 % P/P (affected).

Method: SOP188-MPS-canine, MPS, accredited method

Date of issue: 06.02.2021

Date of testing: 01.02.2021 - 06.02.2021

Approved by: Mgr. Martina Šafrová, Laboratory Manager



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