

Result certificate #012345

Detection of c.754C>T mutation in PDP1 gene causing [pyruvatedehydrogenase] phosphatase deficiency in Clumber and Sussex Spaniels

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 mutation c.754C>T in PDP1 gene was tested. The mutation forms a premature stopkodon that leads to [pyruvatedehydrogenase] phosphatase 1 deficiency disease in Clumber and Sussex Spaniels.

Clinically, PDP1 deficiency is manifested as exercise intolerance that can cause total collapse of the affected dog. The disease is often accompanied by cardiac, pulmonary and neurological problems. So the affected dogs often die prematurely. Disease symptoms can be reduced by the timely initiation of ketogenic diet.

Mutation that causes PDP1 deficiency 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). In offspring of two heterozygous animals following genotype distribution can be expected: 25% N/N, 25% P/P and 50% N/P.

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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