

Detection of c.1046T>C mutation in
SLC25A12 gene causing CDMC disease in
Dutch Shepherds

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.1046T>C mutation in SLC25A12 gene causing CDMC disease (Cerebellar degeneration-myositis complex) in Dutch Shepherds was tested. CDMC combines symptoms of cerebellar degeneration and inflammatory myopathy. Affected dogs develop muscle tremors, stiffness of the pelvic limbs, muscle weakness progressing to inability to walk and severe muscle atrophy. Cerebellar ataxia (stumbling or lack of coordination) and hypermetria (abnormal walking with front legs extending higher than normal) were also observed. Muscle weakness and generalized muscle atrophy began to manifest at 3 to 9 months of age.

Mutation that causes CDMC is inherited autosomally recessively which means that the disease develops only in those dogs who inherit mutated allele from both parents; 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: SOPagrisseq_canine, ngs, accredited method

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Approved by: Mgr. Martina Šafrová, Laboratory Manager



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