GENOMIA
genetic laboratory

Result certificate #012345

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.

SAMPLE

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 diease (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: 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

verified and certified document GENOMIA genetic laboratory

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