

Result certificate #012345

Detection of g.23380074_23483377del mutation in ACSL5 gene causing ILM in Australian Kelpies

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 g.23380074_23483377del mutation in ACSL5 gene causing Lipid malabsorption (ILM) in Australian Kelpies was tested. Lipid malabsorption is a disorder of fat absorption by the intestinal mucosa. The disease manifests itself in young dogs that are stunted in stature and suffer from steatorrhea (fatty stools containing excessive amounts of fat) or bloating. The condition leads to inflammation of the small intestine and abscesses. Another manifestation of the disease is a poor-quality rough coat.

Mutation that causes ILM 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

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Date of testing: 01.02.2021 - 06.02.2021

Approved by: Mgr. Martina Šafrová, Laboratory Manager



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