

**Detection of c.350del mutation in SLC3A1
gene causing cystinuria type I-A
in Labrador Retrievers****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.350del mutation in SLC3A1 gene causing cystinuria type I-A in Labrador Retrievers was tested. In affected dogs, cystine accumulates in the urine, cystine crystals and eventually urinary stones are formed, which can lead to irritation of the urinary tract, stagnation of urine, susceptibility to more frequent urinary tract infections or even complete closure of the urinary tract, which can cause kidney failure. The disease affects males more because they have a longer urinary tract than females.

Mutation that causes cystinuria type I-A in Labrador Retrievers 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: SOPagriseq_canine, ngs, 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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