

Result certificate #012345

Detection of c.2839del mutation in RELN gene causing cerebellar hypoplasia in White Swiss Shepherd Dogs

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.2839del mutation in RELN gene causing cerebellar hypoplasia in White Swiss Shepherd Dogs was tested. Cerebellar hypoplasia means incomplete development of the cerebellum. In White Swiss Shepherds, it occurs together with lissencephaly, a developmental defect characterised by an abnormal structure of the brain tissue, without the typical folds, co-called gyri and sulci on the cerebral cortex of the brain. The condition also accompanies moderate hydrocephalus with dilatation of the cerebral ventricles. The disease manifests itself in puppies as early as 2 weeks of age with ataxia.

Mutation that causes cerebellar hypoplasia 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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Approved by: Mgr. Martina Šafrová, Laboratory Manager



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