

Result certificate #012345

Detection of c.11C>G mutation in RHO gene causing AD-PRA in English Mastiffs and Bullmastiffs

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)

Explanation

Presence or absence of c.11C>G mutation in RHO gene causing dominant progressive retinal atrophy (AD-PRA) in English Mastiffs and Bullmastiffs was tested. The first symptom is night blindness that can be identified in puppies at six weeks of age. The night blindness progresses gradually in blindness and at the age of 1 to 2 years.

Mutation that causes AD-PRA is inherited as an autosomal dominant trait. This means that it only takes one copy of the mutated gene inherited from one parent to cause symptoms of the disease (N/P genotype).

Method: SOPAgriseq_canine, ngs

Date of issue: 06.02.2021

Date of testing: 01.02.2021 - 06.02.2021

Approved by: Mgr. Martina Šafrová, Laboratory Manager



Genomia s.r.o, Republikánská 6, 31200 Plzeň, Czech Republic www.genomia.cz, laborator@genomia.cz, tel: +420 373 749 999