GENOMIA
genetic laboratory

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: 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

SAMPLE

verified and certified document GENOMIA genetic laboratory

Genomia is accredited in compliance with ISO/IEC 17025:2018 under #1549
Genomia s.r.o, Republikánská 6, 31200 Plzeň, Czech Republic
www.genomia.cz, laborator@genomia.cz, tel: +420 373 749 999

ilac-MRA L 1549

Report verification code is: 12AB-CD34-GENO-MIA0-EFGH. You can verify report online at www.genomia.cz
Without a written consent by the lab, the report must not be reproduced unless as a whole.
The result refers only to the tested sample, as received. Genomia is not responsible for the accuracy of the information provided by the customer.