

## Result certificate #130106

Detection of insertion in exon 4 of RD3 gene causing RCD2 disease in collies

## Sample

Sample: 19-13527

Name: Adina Dvůr Bažantnice

Breed: Collie Smooth

Microchip: 972 273 000 004 700 Reg. number: CMKU/CK/1079/16/18

Date of birth: 24.04.2016

Sex: female

Date received: 16.05.2019 Sample type: blood

The identity of the animal has been checked by

MVDr. Michaela Oravská

#### Customer

Ing. Lucie Glaserová Doubek 109 25101 Doubek Czech Republic

# 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 22 bp insertion in exon 4 of RD3 gene causing RCD2 disease (rode-cone dysplasia 2) in rough and smooth collies was tested. RCD2 is manifested by retina degeneration that leads to early onset night blindness in puppies about 6 weeks of age. RCD2 affected dogs are usually totally blind in one year of age.

Insertion in RD3 gene is inherited as an autosomal recessive trait. That means the 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: SOP171-RCD2, fragment analysis

Report date: 23.05.2019

Responsible person: Mgr. Martina Šafrová, Laboratory Manager



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