

# Result certificate #012345

Detection of c.460\_463delAAGA mutation in the exon 5 of RPE65 gene causing CSNB disease in Briard breed

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.460\_463delAAGA mutation in the exon 5 of RPE65 (retina pigment epitelium-specific protein, 65 kDa) causing CSNB (Congenital Stationary Night Blindness) in Briard breed was tested. CSNB disease manifests as slow retina degeneration starting in the age of about six months. During the animal's life CSNB disease can develop to total blindness.

Mutation that causes CSNB in Briards 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: SOPAgriseq\_canine, ngs, accredited method

Sensitivity (probability of correct identification of the defective form of the gene in heterozygous or mutated homozygous) is higher than 99%. Specificity (probability of correct identification of the normal form of the gene in a normal homozygous or heterozygous) is higher than 99%.

## 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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