

Customer: Aneta Pániková, Pribinova 3, 95301 Zlaté Moravce, Slovak Republic

Sample:

Sample: 22-08802

Date received: 07.04.2022

Sample type: buccal swab

Information provided by the customer

Name: AKIRA JARVI Mały Wiedeń

Breed: Finnish Lapphund (Suomenlapinkoira)

Microchip: 616 093 901 138 210

Reg. number: SPKP39

Date of birth: 25.07.2020

Sex: male

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.5G>A mutation in PRCD gene causing PRA-prcd (Progressive Retinal Atrophy) was tested. Disease causes degeneration of retinal cells. That results in complete blindness of the animal. The age of onset of disease varies, but, generally, it cannot be recognized before the adulthood of the animal.

Mutation that causes PRA-prcd 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.

It is necessary to be aware that not all retinal diseases belong in progressive retina atrophy group of disorders and not all are a variant of PRA-prcd. In many breeds, the cause of PRA inheritance has not been still clarified. It is also possible that several mutations can be responsible for retinal atrophy in one breed. Therefore, we recommend that an eye examination by a veterinary ophthalmologist is performed every year.

The analysis was carried out by partner laboratory.

Method: SOP182-PRA, HRMA

Date of issue: 12.04.2022

Date of testing: 07.04.2022 - 12.04.2022

Approved by: Mgr. Martina Šafrová, Laboratory Manager



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