

## Result certificate #064778:

Detection of c.8392delC mutation in the CUBN gene causing IGS in border collies by fragment analysis

Sample

Sample: 15-50104

Name: FLOWER OF OLD HILL NOT JUST A

**BANANA** 

Breed: Border Collie

Microchip: 900 108 000 883 765 Reg. number: CMKU/BOC/7636/-14/14

Date of birth: 14. 3. 2014

Sex: female

Date received: 18.05.2015 Sample type: blood

Sample certified by Vet/Tech or witness.

Customer

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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.8392delC mutation in the CUBN gene causing IGS (Imerslund-Gräsbeck syndrome) or intestinal cobalamin malabsorption in border collies was tested. IGS is metabolic disorder in border collies. Signs appear early in 6 to 12 week of dog's age and include failure to thrive and chronic loss of appetite. The affected dogs can suffer from neutropia, non-regenerative anaemia, anisocytosis and poikilocytosis, megaloblastic changes in bone marrow, reduction of Cbl level, methylmalonic aciduria and homocysteinemia.

Mutation that causes IGS in border collies 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), they are healthy but they can transmit the mutation on their offspring. In offspring of two heterozygous animals following genotype distribution can be expected: 25 % N/N, 50 % N/P and 25 % P/P.

Method: SOP159

Report date: 20.05.2015

Responsible person: Mgr. Martina Šafrová, Laboratory Manager

Saferra

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