

Result certificate #020675:

Detection of c.619C>T mutation in CLN5 gene causing NCL in border collies by PCR-RFLP

Sample Sample: 12-13625 Name: DEMON CAYS The Best of Gajowy Breed: Border Collie Reg. number: ---Microchip: 967000009173264 Date of birth: 13.06.2008 Sex: male Date received: 04.06.2012 Sample type: blood **Customer** Piotr Strzelecki Długa 256 42-436 Sławniów Poland

Result: Mutation was not detected (N/N)

Explanation

Mutation c.619C>T in CLN5 gene was tested. This mutation forms premature stopcodon; the mutated enzyme is 144 amino acids shorter and cannot full fill its normal function. Neuronal ceroid lipofuscinosis (NCL) is a neurodegenerative disorder that is characteristic by accumulation of lipopigments (coroid and lipofuscin) in the lysosomes. The beginning and clinical course of the disease vary greatly and are very individual. The rate of neurodegeneration increases together with the age, with psychical abnormalities and ataxy usually developing in all affected dogs. Increased restlessness, aggression, hallucinations, hyperactivity and epileptic attacks can be observed as well. Accompanying symptom is damaged retina due to lipopigment storage. Affected individuals rarely survive more 28th month of age.

Mutation that causes NCL in border collies is inherited autosomally recessively which means that the disease develops only in those dogs who inherit mutated allele from both parents; disease affects dogs with P/P (positive / positive) genotype only. The dogs with N/P (positive / negative) genotype are considered carriers of the disease (heterozygotes). In offspring of two heterozygous animals following genotype distribution can be expected: 25 % N/N (healthy non-carriers), 25 % P/P (affected), and 50 % N/P (healthy carriers).

Method: SOP39

Report date: 05.06.2012 Responsible person: Mgr. Markéta Dajbychová, Deputy Laboratory Manager

Dey hychora'

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