



Customer

JUHÁSZ CSILLA
BALLÓSZÖG TANYA 226/B.
6000 KECSKEMÉT
Hungary

Veterinarian Dr. Fáy József
confirmed sample origin - an optimal
form.

Details of animal

Sample: 10-02102
Animal: COLLIES FROM SUSY BONITA
Breed: Border Collie
Reg. number: 926/05
Microchip: -
Year of birth: 2005
Sex: female
Date received: 25.01.2010
Sample type: blood

Result: Mutation was not detected (N/N)

Explanation

7,8 kb deletion in intron 4 of canine NJEH1 gene was tested. The eye anomaly caused by this deletion affects most often collie breeds including Rough Collie, Smooth Collie, Border Collie, Shetland Sheepdog and Australian Sheepdog. CEA disease 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 (healthy non-carriers), 25 % P/P (affected), and 50 % N/P (healthy carriers).

Method: SOP02, 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%.

Report date: 27.01.2010

Responsible person: Mgr. Martina Šafrová, Laboratory Manager