

**Result certificate #024459:**

**Sample**

Sample: 12-24630  
Name: Bring me to life of Dashing Dawn  
Breed: Border Collie  
Reg. number: 2590/11  
Microchip: 972274000073652  
Date of birth: 24 August 2011  
Sex: female  
Date received: 07.09.2012  
Sample type: buccal swab

**Detection of c.227\_230delATAG mutation in the MDR1 gene causing drug sensitivity in dogs by fragmentation analysis of PCR product**

**Customer**

Diana Krassovari  
Lonyai utca 1.  
2330 Dunaharaszti  
Hungary

**Result: Mutation was not detected (N/N)**

**Explanation**

Presence or absence of c.227\_230delATAG mutation in MDR1 gene was tested. This mutation causes a frame shift and formation of a stop codon during P-glycoprotein synthesis. P-glykoprotein, an ATP-dependent transporter of various substrates, is contained in cells lining the blood vessels in the brain. In P-glycoprotein defective animals, administering of ivermectin or similar drug can lead to elevated levels of drug in the CNS, resulting in potentially lethal neurotoxic reaction. These drugs include, but are not limited to: Acepromazine, Butorphanol, Doramectin, Doxorubicin, Ivermectin, Loperamide, Milbemycin, Moxidectin, Selamectin, Vinblastine, Vincristine.

MDR1 related drug hypersensitivity is inherited as an autosomal recessive trait. That means the defect affects dogs with P/P genotype only. The dogs with N/P genotype are considered carriers of the deletion (heterozygotes). The dogs with N/N genotype are not endangered with MDR1 related drug hypersensitivity. Multiple drug hypersensitivity based on MDR1 gene mutation was proved in following breeds: Rough Collie, Smooth Collie, Shetland Sheepdog, Australian Sheepdog, White Swiss Shepherd Dog, Wäller, Bobtail, Border Collie and others.

Method: SOP04, 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: 14.09.2012

Responsible person: Mgr. Martina Šafrová, Laboratory Manager

Genomia is accredited according to ISO 17025 under #1549.  
Genomia s.r.o, Janáčkova 51, 32300 Plzeň, Czech Republic, VAT#: CZ25212991  
www.genomia.cz, laborator@genomia.cz, tel: +420 373 749 999

