

Result certificate #067598:

Detection of mutation c.1028_1032delGAGAA in RPGR gene causing XL-PRA in Siberian Husky and Samoyed by fragmentation analysis

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Sample

Sample: 15-20210 Name: NIKKA OF DEIMOS Bielkersmils Breed: Samoyed Microchip: 752 098 100 731 825 Reg. number: CMKU/S/2616/-15/15 Date of birth: 4.3.2015 Sex: female Date received: 22.07.2015 Sample type: buccal swab Sample certified by Vet/Tech or witness.

Result: Based on mutation examination genotype was determined Xn/Xn

Explanation

Presence or absence of mutation c.1028_1032delGAGAA in exon ORF15 of RPGR gene (retinitis pigmentosa GTP's regulator) was tested. This mutation causes X-linked progressive retinal atrophy diseases in Siberian Husky and Samoyed breeds. The first symptoms appear by clinical examination in 6 months. Later, rods light receptors begin to appear irregularly damaged. Cones damage arises in final stage of XL-PRA disease. In age of 4 years, affected dogs are usually completely blind.

Females have XX chromosomes so they can have following XL-PRA genotypes: **XnXn** – females with two normal X chromosomes = normal phenotype, a healthy female **XnXm** – females with one normal X (Xn) and one mutant X (Xm) = a female carrier. Clinical disability of female carriers is individual, depending on the X chromosome inactivation. **XmXm** – females with two mutated X chromosomes = an affected female

Males have XY chromosomes so they can have following XL-PRA genotypes: **XnY** – normal phenotype, a healthy male **XmY** – an affected male; he inherited mutated X chromosome from his mother

Method: SOP24, accredited method

Report date: 30.07.2015 Responsible person: Mgr. Martina Šafrová, Laboratory Manager

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