# GENOMIA
genetic laboratory

**Result certificate #067597:**

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

**Sample**
Sample: 15-20211
Name: CRAZY YAZZ PARTY Bjellakis
Breed: Samoyed
Microchip: 752 098 100 615 889
Reg. number: CMKU/S/2486/-14/12
Date of birth: 14.2.2012
Sex: female
Date received: 22.07.2015
Sample type: buccal swab
Sample certified by Vet/Tech or witness.

**Customer**
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This document replace document dated 30.7. 2015 (microchip was corrected).

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## 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: 10.08.2015
Responsible person: Mgr. Martina Šafrová, Laboratory Manager

Genomia is accredited according to ISO/IEC 17025:2005 under #1549.

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