

REFERENCE NO.: 2019 - 27098

OWNER:

RICCARDO VOLERI

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ITALY

NAME/LABEL:

GOOSE DOMUS M.A.S.

SPECIES: DOG

BREED: ROTTWEILER

SEX: MALE

MICROCHIP NO.: 380260041777425

TATOO NO.: NOT PROVIDED

PEDIGREE NO.: LO1424632

GENETIC REPORT

SAMPLE: DNA ISOLATE

SAMPLE TAKEN BY: SILVIA COSTAMAGNA, DVM

REQUESTED TEST: JUVENILE LARYNGEAL PARALYSIS & POLYNEUROPATHY (JLPP)

RESULT: CLEAR

COMMENT :

The test examines presence or absence of RAB3GAP1 gene mutation (c.743delC) described as the cause of Juvenile Laryngeal Paralysis & Polyneuropathy (JLPP) in Black Russian Terrier and Rottweiler. JLPP is neuronal disease characterized by difficult breathing, difficulty swallowing, difficulty to getting up and also Ocular changes. RAB3GAP1 gene defect is inherited as an autosomal recessive trait.

Regarding to the presence of tested mutation animals are classified in three groups:

- Clear (wt/wt) - mutation is not present, normal genotype
- Carrier (mut/wt) - one of two alleles carries tested mutation, disease is not clinically manifested
- Affected (mut/mut) - both alleles carry tested mutation, disease is clinically manifested

For each group different breeding strategies should be followed. Breeding of affected and carrier animals should be avoided. If particularly valuable animal is classified as affected, it should be bred only with clear animal. In such case, all first generation siblings will be carriers. If a carrier is bred with clear animal, 50% of siblings are expected to be clear. In case two carriers are bred, 25% of siblings are expected to be clear and 50% are expected to be carriers. However, 25% of siblings are expected to be affected, therefore such breeding practice is discouraged.

AUTHORIZED SIGNATURE:

MARIBOR, 17.06.2019