

# GENETIC ANALYSIS REPORT



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## ANIMAL'S DETAILS

Registered Name: Nicnak Gossip Girl  
Pet Name: Nicnak Gossip Girl  
Breed: Australian Labradoodle  
Date of Birth / Age:

Registration No: Pending  
Microchip No: 900032002888475  
Sex: Female  
Colour:

## COLLECTION DETAILS

Case Number: 16-112087  
Collected By: Kaylene Steffens

Date of Test: 27/01/16  
Approved Coll. Mthd.:

Sample with Lab ID Number 16-112087 was received at Orivet Genetics, DNA was extracted and analysed with the following results reported:

### DISEASE(S):

EXERCISE INDUCED COLLAPSE - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
DEGENERATIVE MYELOPATHY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
AUTOSOMAL HEREDITARY RECESSIVE NEPHROPATHY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
CENTRONUCLEAR MYOPATHY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
CONE-ROD DYSTROPHY 1 - PRA - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
CYSTINURIA - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
GLOBOID CELL LEUKODYSTROPHY/KRABBE'S DISEASE - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
MUCOPOLYSACCHARIDOSIS - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
MYOTUBULAR MYOPATHY X LINKED - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
NARCOLEPSY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
NEONATAL ENCEPHALOPATHY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
PHOSPHOFRUCTOKINASE (PFK) DEFICIENCY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
PYRUVATE KINASE (PK) DEFICIENCY - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
PROGRESSIVE ROD CONE DEGENERATION - PRA - NORMAL BY PARENTAGE HISTORY  
VON WILLEBRAND'S DISEASE TYPE I - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
GANGLIOSIDOSIS - TYPE 1 & 2 - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
HEREDITARY NASAL PARAKERATOSIS (DRY NOSE) - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
SKELETAL DYSPLASIA 2 (COL11A2) - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)  
ELLIPTOCYTOSIS (B-SPECTRIN) - NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)

### TRAIT(S):

LONG HAIR GENE (PHENOTYPE) - **AFFECTED/POSITIVE FOR THE VARIANT**  
A-LOCUS AGOUTI - **at/a TRICOLOR/TAN POINTS (CARRYING BICOLOUR/GENE)**  
B (TYRP1 LOCUS) BROWN/CHOCOLATE - **AFFECTED bb (BROWN/CHOCOLATE, LIVER OR RED)**  
DILUTE MLPH GENE (BLUE/GREY) - **DD - NO COPY OF MLPH-D ALLELE**  
K-LOCUS (DOM BLACK/WILD TYPE) - **CARRIER - K/k ONE COPY DOM BLACK and ONE COPY NON BLACK**  
EM-LOCUS MELANISTIC MASK ALLELE - **E<sup>m</sup> E - ONE MASK AND ONE NORMAL EXTENSION ALLELE**  
E (EXTENSION) LOCUS MC1R - **Ee - CARRIES EXTENSION (YELLOW/WHITE/APRICOT/RUBY)**  
SPOTTING LOCUS (W GENE) - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
FOLLICULAR DYSPLASIA - COLOUR DILUTION ALOPECIA - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
BLACK HAIR FOLLICULAR DYSPLASIA - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**



16-112087

### RESULTS REVIEWED AND CONFIRMED BY:

  
Dr. Noam Pik BVs MDSV

  
George Sofronidis BSc (Hons)