

# GENETIC ANALYSIS REPORT

## OWNER'S DETAILS

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## ANIMAL'S DETAILS

Registered Name: Tallai Kirra Lee  
Pet Name: Tallai Kirra Lee  
Breed: Australian Labradoodle  
Date of Birth / Age:

Registration No: 0068-068-01  
Microchip No: 953010001058526  
Sex: Female  
Colour: Not Supplied By Client

## COLLECTION DETAILS

Case Number: 17-157993  
Collected By: Dr. Lisa Butler  
Date of Test: 11/07/17  
Approved Coll. Mthd.: Yes

Sample with Lab ID Number 17-157993 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)**  
CANINE HYPERURICOSURIA - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
PROGRESSIVE RETINAL ATROPHY - CORD1/RCD-4 - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
GLOBOID CELL LEUKODYSTROPHY/KRABBE'S DISEASE - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
MERMECTIN SENSITIVITY MDR1 - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
MUCOPOLYSACCHARIDOSIS - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
MYOTUBULAR MYOPATHY X LINKED - **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 / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
VON WILLEBRAND'S DISEASE TYPE I - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
PROGRESSIVE RETINAL ATROPHY - RCD4 - **INDETERMINABLE - RESULT OBTAINED IS INCONCLUSIVE**  
GENERALISED PRA 1 - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
MALIGNANT HYPERTHERMIA - **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)**  
GM2 GANGLIOSIDOSIS - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
HEREDITARY NASAL PARAKERATOSIS (DRY NOSE) - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
NARCOLEPSY - **NORMAL / CLEAR / NEGATIVE (NO VARIANT DETECTED)**  
**TRAIT(S):**  
LONG HAIR GENE (PHENOTYPE) - **POSITIVE - SHOWING THE PHENOTYPE**  
A-LOCUS AGOUTI - **at/at - TAN POINTS MAY BE BRINDLED (SEE K LOCUS)**  
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 - **CARRIER- EmE ONE COPY OF MASK ALLELE DETERMINED BY A and K**  
E (EXTENSION) LOCUS MC1R - **EE - DOMINANT BLACK DOES NOT CARRY YELLOW/RED/WHITE**  
BLACK HAIR FOLLICULAR DYSPLASIA - **CARRIER (ONE COPY OF VARIANT DETECTED)**  
BLACK AND TAN/SADDLE COAT - **CARRIER (ONE COPY OF VARIANT DETECTED)**  
BROWN (GLNT331STOP) STOP CODON (**b<sup>s</sup>**) - **Bb - CARRIER OF BROWN / LIVER / CHOCOLATE**  
BROWN (345DELPRO) DELETION (**b<sup>d</sup>**) - **BB - DOES NOT CARRY BROWN or CHOCOLATE**  
BROWN (SER41CYS) INSERTION CODON (**b<sup>c</sup>**) - **Bb - CARRIER OF BROWN / LIVER / CHOCOLATE**

PENDING TEST(S): PARENTAGE VERIFICATION



17-157993