

GENETIC ANALYSIS REPORT



OWNER'S DETAILS

Troy Scaddan

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ANIMAL'S DETAILS

Registered Name: Chatjoli Kaia
Pet Name: Violet
Breed: British Shorthair

Registration No: 59W12
Microchip No: 981000300650299
Sex: Female

COLLECTION DETAILS

Case Number: 15-120704
Collected By: Troy Scaddan

Date of Test: 19/06/15
Approved Coll. Mthd.:

Sample with Lab ID Number 15-120704 was received at Orivet Genetics, DNA was extracted and analysed with the following results reported:

DISEASE(S): PYRUVATE KINASE (PK) DEFICIENCY (**NORMAL / CLEAR - NO MUTATION DETECTED**)
MUCOPOLYSACCHARADOSIS (**NORMAL / CLEAR - NO MUTATION DETECTED**)
POLYCYSTIC KIDNEY DISEASE (**NORMAL / CLEAR - NO MUTATION DETECTED**)
NEIMANN-PICK DISEASE TYPE C (**NORMAL / CLEAR - NO MUTATION DETECTED**)
FAMILIAL EPISODIC HYPOKALEMIC POLYMYOPATHY (**NORMAL / CLEAR - NO MUTATION DETECTED**)
HYPERTROPHIC CARDIOMYOPATHY - MAINE COON (**NORMAL / CLEAR - NO MUTATION DETECTED**)
HYPERTROPHIC CARDIOMYOPATHY - RAGDOLL (**NORMAL / CLEAR - NO MUTATION DETECTED**)
PROGRESSIVE RETINAL ATROPHY (PRA) CEP 290 (PRA-RDC) (**NORMAL / CLEAR - NO MUTATION DETECTED**)
PROGRESSIVE RETINAL ATROPHY (PRA) CRX (PRA-RDY) (**NORMAL / CLEAR - NO MUTATION DETECTED**)
SPINAL MUSCULAR ATROPHY (SMA) - MAINE COON (**NORMAL / CLEAR - NO MUTATION DETECTED**)
GLYCOGEN STORAGE DISEASE TYPE IV (**NORMAL / CLEAR - NO MUTATION DETECTED**)
GANGLIOSIDOSI- GM2 (**NORMAL / CLEAR - NO MUTATION DETECTED**)

TRAIT(S): ALBINISM (**ALB N / ALB N - NO ALBINO MUTATION DETECTED**)
AMBER (**E/E - NO COPIES OF AMBER MUTATION DETECTED**)
DILUTION (**d/d TWO COPIES OF DILUTE ALLELE - COAT COLOR IS DILUTED**)
BLOOD GROUP (**TYPE B (b/b) - CAT HAS A HIGH LEVEL OF anti-A ANTIBODIES**)
CHOCOLATE AND CINAMMON (**B/B (FULL COLOUR - CAT DOES NOT CARRY BROWN OR CINNAMON)**)
COLOURPOINT RESTRICTION (**C/C FULL COLOUR, DOES NOT CARRY BURMESE (SEPIA) OR SIAMESE**)
AGOUTI (**a/a NON AGOUTI SELF COLORED (SOLID COLORED)**)
LONG HAIR SHORTHAIIR (**N/N = NONE OF THE 4 LONG HAIR MUTATIONS DETECTED**)
WHITE GLOVES (**N⁹/N⁹ - NO GLOVE MUTATION DETECTED**)



15-120704

RESULTS REVIEWED AND CONFIRMED BY:


Dr. Noam Pik BVs MDSV


George Sofronidis BSc (Hons)