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DNA ANALYSIS PROTOCOL FOR DETECTION OF HEREDITARY DISEASES

Číslo Protokolu D2101008909

Zákazník: Hedvika ŽIVNÍČKOVÁ
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Typ vzorky: Krv (EDTA)
Dátum narodenia: 30.12.2019
Pohlavie: F

Dátum odberu: 04.01.2021
Dátum doručenia: 08.01.2021
Dátum analýzy: 12.01.2021

Identita jedinca bola overená. Odber vykonaný veterinárom: MVDr. Jiří Durmon, číslo KVL: 3854

Rasa/Meno	Číslo čipu alebo tetovania Certifikát pôvodu	Kód laboratória	Typ Analýzy	Výsledok
Papillon / Aurora ot Romanovs Rose	112060000038836 CMKU/PAP/4853/-20/19	210108/X0076	Pap-PRA1	N/N non-affected

Výsledok analýzy je uložený v databáze pod laboratórnym kódom 210108/X0076.

Poznámky:

pap-PRA1 - progressive retinal atrophy, causal indel mutation (1bp deletion followed by 6 bp insertions) in CNGB1 gene that is responsible for PRA in Papillon and Phalène breeds

N/N– homozygous individual non-affected are genetically clear.

N/A– heterozygous carrier are clinical without any symptoms. They are genetically considered carriers of the disease, disease is transmitted to offspring.

A/A– homozygous affected individual

Upozornenie: Notice: This protocol applies exclusively to the sample and the data that were supplied by the submitter. DNA analysis concerns only the above-mentioned disease. No information regarding the customer as well as the purpose and results of the analysis will be provided to third parties.

V Bratislave dňa: 12.01.2021

Ing. Marcela Bieliková, PhD.