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DNA ANALYSIS PROTOCOL FOR DETECTION OF HEREDITARY DISEASES

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Dátum odobrania vzorky: 21.09.2016
Dátum doručenia: 26.09.2016
Dátum analýzy: 27.09.2016

Rasa/Meno	**Číslo čipu alebo tetovania** Certifikát pôvodu	**Kód laboratória**	**Typ Analýzy**	**Výsledok**
Papillon / D-MONARCH BLACK Babočka Admirál	956000001107330 CMKU/PAP/2566/08/10	160926/N1944	Pap-PRA1	N/N non-affected

Výsledok analýzy je uložený v databáze pod laboratórnym kódom 160926/N1944.

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 clinically without any symptoms. They are genetically considered carriers of the disease, disease is transmitted to offspring.
A/A– homozygous affected individual

Upozornenie: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 purpose and results of the analysis will be provided to third parties.

V Bratislave dňa: 27.09.2016
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