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PKD1 AND PERSIAN DERIVED PRA REPORT

DANIEL BUCHS ACHERN C, 3714 FRUTIGEN ACHERN 3C SWITZERLAND	Case: CAT113072 Date Received: 25-Feb-2019 Print Date: 26-Feb-2019 Report ID: 1762-8226-9134-5157 Verify report at www.vgl.ucdavis.edu/myvgl/verify.htm
Cat: EVAN VON KANDERGARDEN Reg: FFH LO 985772 DOB: 04/15/2018 Sex: Male Breed: Ragdoll Microchip: 756098100799471 Color: seal	

PKD1 Result	PRA-pd Result
N/N	Not Requested

PKD1 Result Codes:

N/P Affected - 1 copy of the PKD1 gene, cat has or will develop PKD. Severity of symptoms cannot be predicted*

N/N Normal - Does not possess the disease-causing PKD1 gene.

The disease is inherited as an autosomal dominant trait, which means that a heterozygote (N/P) bred to a normal (N/N) will result in approximately half of the offspring being affected and half being normal. There are no observed homozygous affected (P/P), which suggests that the mutation is embryonic lethal.

*If your cat tests positive for PKD1, we recommend that you contact your veterinarian for information on disease progression and management.

PRA-pd Result Codes:

N/N N/N - Normal - no copies of the PRA-pd mutation.

N/PRApd Carrier - 1 copy of the PRA-pd mutation; vision appears normal. Breeding between carriers is expected to produce 25% affected kittens.

PRApd/PRApd Affected - 2 copies of the PRA-pd mutation; cat will go blind.

For more information on PKD1 and PRA-pd test results, please go to:

www.vgl.ucdavis.edu/services/pckd1.php
www.vgl.ucdavis.edu/services/cat/PRApd.php

IDENTITY MARKERS

LOCUS	TYPE	LOCUS	TYPE
FCA075	PT	FCA220	L
FCA223	GU	FCA678	JM
FCA698	NS		