


Progressive Retinal Atrophy (GR_PR2)

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Profile:	DG2021/20150	Species:	Canis lupus familiaris / Canine / Dog
Name:	Bonyl Kilted Ochre	Microchip #:	941000026460135
Breed:	Golden Retriever	Registration #:	ZA009682B21
Test:	[GR_PR2] Progressive Retinal Atrophy (GR_PR2)		
Results:	c.699delA	AA	CLEAR

Sample Type: Whole Blood (EDTA)	Extraction Method: DNA Extraction: D4069	Test Type: Genetic Health
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[GR_PR2] Progressive Retinal Atrophy (GR_PR2)

Progressive Retinal Atrophy (PRA) is a collective of genetic eye disorders that share similar symptoms. There are numerous mutations that cause PRA in various canine breeds.

PRA_GR2 is a late onset PRA discovered in Golden Retrievers responsible for 30% of PRA cases. This test detects a single adenine deletion at c.699 of the TTC8 gene causing a shift in the reading frame.

PRA_GR2 is an autosomal recessive photoreceptor degenerative disease, where two copies of the mutation are required for an individual to be affected.

References: Downs et al 2014. A novel mutation in TTC8 is associated with progressive retinal atrophy in the golden retriever. Canine Genetics and Epidemiology 1(4).

It is the sender's responsibility to ensure the correctness of the information accompanying the samples. In no event shall Inqaba Biotechnical Industries (Pty) Ltd or its divisions be held liable for indirect, substantial or secondary damages of any kind. Results are usually made available within 7-14 days of receipt of samples. Please note that results are only released subject to payment.

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The DNA profile is based on a preliminary marker panel that is subject to modification pending additional genetic information.

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