

PROGRESSIVE RETINAL ATROPHY – GR2

Client Name:	E. RADEMEYER	REPORT NO.:	SA2019/79516/0829/03
Kennel Name:			
Client Address:	5 DE VOS STREET STELLENBOSCH 7600		
Client Tel No.:	084 504 5493		
Canine Name:	BRUCE		
Breed:	GOLDEN RETRIEVER		
Microchip No.:	900032001979297	Registration No.:	ZA051342B14
Genetic Test:	PROGRESSIVE RETINAL ATROPHY – GR2		
Result:	CLEAR		

PROGRESSIVE RETINAL ATROPHY – GR2 (PRA_GR2)

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.

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).

SAMPLE TYPE: EDTA BLOOD AMPULE
EXTRACTION METHOD: DNA EXTRACTION
TEST TYPE: SANGER SEQUENCE DETECTION

BREEDING IMPLICATIONS

		MATERNAL CANDIDATE		
		CLEAR	CARRIER	AFFECTED
PATERAL CANDIDATE	CLEAR	ALL CLEAR	50% CLEAR 50% CARRIER	ALL CARRIER
	CARRIER	50% CLEAR 50% CARRIER	25% CLEAR 50% CARRIER 25% AFFECTED	50% CARRIER 50% AFFECTED
	AFFECTED	ALL CARRIER	50% CARRIER 50% AFFECTED	ALL AFFECTED

Disclaimer: This report does not disregard the existence of any unknown or rare variant of TTC8 gene that may cause PRA, nor does it disregard the existence of any unknown or rare variant of any other gene that may cause or be associated with PRA in Golden Retrievers.



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