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ITALIAN GREYHOUND GENETIC TEST REPORT

ELIAS DUARTE CAIXA POSTAL 19018 81531-990 CURITIBA PR BRAZIL	Case: NCD65518 Date Received: 03-Aug-2017 Print Date: 11-Aug-2017 Report ID: 3611-4660-8738-3135 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
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Name: GENOWEWA DO BR REINO DOB: 01/06/2013 Sex: Female Breed: Italian Greyhound Microchip: 963008000229729 Color: RED FAWN	Reg: PRG/13/00555
Sire: CH KAOS RED HOT SKY ROCKET IN FLIGH Dam: CH KINGA DO BR REINO	Reg: TR98796903 Reg: PRG/05/01541

Familial Enamel Hypoplasia¹	N/N	No copies of FEH mutation; dog is normal.
Susceptibility to PRA²	AA BB CC DD Ee	Genotype has low risk for PRA-IG1. Dog is not likely to develop PRA-IG1, but is a carrier of PRA-risk allele e.
Susceptibility to Glaucoma³	N/N	No copies of the glaucoma associated haplotype. Dog has a low risk (2%) of developing primary closed angle glaucoma.

Notes:

1. This test is specific for the mutation associated with Familial Enamel Hypoplasia (Autosomal Recessive Amelogenesis Imperfecta) in Italian Greyhounds.
2. This test is specific for 5 mutations associated with susceptibility to the major cause of PRA presently occurring in the Italian Greyhound breed. This test is not valid for other breeds. It is important to note that there may be other genetic forms of PRA in the breed.
3. This test is specific for the 2 mutations associated with susceptibility to primary closed angle glaucoma in the Italian Greyhound breed. This test is not valid for other breeds.

For more information about Familial Enamel Hypoplasia, Susceptibility to PRA and Susceptibility to Primary Closed Angle Glaucoma please go to:
www.vgl.ucdavis.edu/services/italiangreyhound.php