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ITALIAN GREYHOUND GENETIC TEST REPORT

HELENA SEBESTOVÁ U RADNICE 455 250 70 ODOLENA VODA CZECH REPUBLIC		Case: NCD103785 Date Received: 04-Sep-2019 Print Date: 11-Sep-2019 Report ID: 5559-6165-7975-8139 Verify report at www.vgl.ucdavis.edu/myvgl/verify.htm
Name: SARAFINA CANA DEL VENTO DOB: 03/27/2019 Sex: Female Breed: Italian Greyhound Microchip: 981100004551261 Color: Blue		Reg: LOSH1286702
Call Name: Sara		
Sire: QUIRINUS PUSTYNNY WIATR Dam: LUNA COULEURS D'AUTOMNE		Reg: CLP/IT/2679 Reg: LOSH-01114368
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 alleles b and 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 detailed information on Italian Greyhound Genetic test results, please go to:
www.vgl.ucdavis.edu/services/italiangreyhound.php