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ITALIAN GREYHOUND GENETIC TEST REPORT

HELENA SEBESTOVÁ U RADNICE 455 250 70 ODOLENA VODA CZECH REPUBLIC	Case: NCD26840 Date: 13-Mar-2015 Print Date: 16-Nov-2015 Report ID: 8056-7242-6127-2040 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
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Name: EARLEEN ANNAPERLA DOB: 01/13/2013 Sex: Female Breed: Italian Greyhound Microchip: 203098100329319 Color: Isabell	Reg: CLP/IT/2109/14
Sire: FIEFOERNIEK'S FEEL THE GROOVE Dam: PERLA FERITTE BUGSY	Reg: NHSB 2791324 Reg: CMKU/IT/1430/08/09

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, c 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 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