UNIVERSITY OF CALIFORNIA, DAVIS

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ITALIAN GREYHOUND GENETIC TEST REPORT

ISABELL BOHMANN
SCHAFWIESENSTRASSE 66
4600 WELS
AUSTRIA

Case: **NCD66714**
Date Received: 29-Aug-2017
Print Date: 06-Sep-2017
Report ID: 2107-3475-7207-7106
Verify report at www.vgl.ucdavis.edu/myvgl/verify.html

Name: **LUISE OF MYSTICAL WOODEN HOUSE** *Reg:* **WISP 748**

DOB: **02/02/2017** *Sex:* **Female** *Breed:* **Italian Greyhound** *Microchip:* **276098106248654** *Color:* **grey**

Sire: DERVISCH DAYDREAMER *Reg:* SE 59384/2014
Dam: DARK LEGEND'S YULE *Reg:* WI 4327

Familial Enamel Hypoplasia[1]	N/N	No copies of FEH mutation; dog is normal.
Susceptibility to PRA[2]	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 c.
Susceptibility to Glaucoma[3]	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