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PROTOCOL OF DNA ANALYSIS FOR DETECTION OF INHERITED DISEASES

Submitter of analysis : Bouchand Camille
2 rue des Frênes
38320 Eybens, France

Date of sampling : 11.12.2013
Date of samples receipt: 17.12.2013
Date of analysis : 20.12.2013

Breed/name	**Tattoo or chip/** Certificate of origin	**Laboratory code**	**Type of analysis**	**Result**
Berger Australien/ GW'Mallix des Terres d'Urfé	**250268720134266**	131217/H3103	PRA-prcd	N/N Non-affected

The samples and results of analysis are stored in a database under the lab code 131217/H3103.

Hints:

Genotype PRA-prcd N/N – healthy subject – non-affected
Both genes, inherited from both male and female are unaffected. That means that the subject has both alleles healthy.

Genotype PRA-prcd N/A – carrier
Subjects with confirmed heterozygous N/A genotype are carriers. Gene mutation can be transmitted to offspring.

Genotype PRA-prcd A/A – affected subject
The subject is a homozygote with A/A genotype, which inherited the affected allele from both parents and thus is affected by the disease.

Notice: This protocol applies exclusively to the sample and the data that were supplied by the submitter. DNA analysis concerns only the above mentioned disease. No information regarding the customer as well as purpose and results of the analysis will be provided to third parties.

In Bratislava 20.12.2013
Ing. Marcela Bieliková, PhD.