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PROTOCOL OF DNA ANALYSIS FOR DETECTION OF INHERITED DISEASES

Submitter of analysis: Trotobas Amandine Date of sampling: 09.01.2014

La guillemarde

Date of samples receipt: 03.02.2014

03240 Rocles, France

Date of analysis: 13.02.2014

Breed/name	Tattoo or chip / Certificate of origin	Laboratory code	Type of analysis	Result
Berger Australien / GIVE ME	250269604523322			N/N
THE MOON DES DOLMENS D'AN ARVOR	LOF 1 B.AUS. 31937/0	140203/K0320	HSF4/HC	Non-affected

The samples and results of analysis are stored in a database under the lab code 140203/K0320.

HSF4 (**HC**) – Hereditary cataract – deletion/insertion 1 bp in HSF4 gene.

Genotype HC N/N – non-affected subject –**NORMAL (NON-AFFECTED)**

Both genes, inherited from both male and female are unaffected. That means that the subject has both alleles healthy.

Genotype HC N/A - CARRIER

Subjects with confirmed heterozygous N/A genotype are carriers. Gene mutation can be transmitted to offspring.

Genotype HC 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.

<u>Notice:</u> 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 14.02.2014 Ing. Marcela Bieliková, PhD.