Owner's Details

GENETIC ANALYSIS REPORT

Wade Robertson

9 Chedworth Ave

Christchurch, NZ

Animal's Details

Registered Name:

Pet Name: Wyatt

Lab ID:

12-012326

HOTWYR LOCK STOCK AND BARREL

Microchip No: Registration No.

941000014103318 04910-2010

German Wirehair Pointer

27/12/12

Date of Test:

Collected By:

Sample with Lab ID Number 12-012326 was received at ASAP Laboratories, DNA was extracted and analysed with the following results reported:

Bill Gardyne

VON WILLEBRAND'S DISEASE TYPE I (NORMAL)

TRAIT(S):

DISEASE(S):

B (BROWN) LOCUS (Bb)
D (DILUTE) LOCUS (DD)

EM-LOCUS MELANISTIC MASK ALLELE (EM) (NORMAL)

E-LOCUS (EXTENSION - YELLOW/LEMON/RED/CREAM/APRICOT) (NORMAL)

DNA PROFILE The DNA Profile below represents the genetic identification of HOTWYR LOCK STOCK AND BARREL

CSNP67CSNP68CSNP69CSNP70CSNP71CSNP72CSNP73CSNP74CSNP75CSNP76CSNP77CSNP78CSNP79CSNP80CSNP81CSNP83CSNP83CSNP83CSNP86CSNP86CSNP86CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP87CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNP88CSNPCSNP45CSNP46CSNP47CSNP48CSNP49CSNP50CSNP51CSNP52CSNP53CSNP53CSNP55CSNP55CSNP57CSNP58CSNP59CSNP60CSNP61CSNP63CSNP63CSNP63CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP66CSNP6 CSNP23 CSNP01 CSNP02 CSNP03 CSNP04 CSNP05 CSNP06 CSNP07 CSNP08 CSNP09 CSNP10 CSNP11 CSNP12 CSNP13 CSNP14 CSNP16 CSNP16 CSNP17 CSNP18 CSNP19 CSNP20 CSNP21 CSNP22 CS A CSNP25 CSNP26 CSNP27 CSNP28 CSNP29 CSNP31 CSNP31 CSNP32 CSNP33 CSNP34 CSNP35 CSNP36 CSNP36 CSNP38 CSNP39 CSNP40 CSNP41 CSNP42 CSNP43 CSNP44 CSNP45 CSNP46 CSNP47 CSNP46 CSNP47 CSNP48 CSNP47 CSNP48 CS C G SA GG 8 GA GG CC S င္ပ CC S GG 8 2 CC GG # A GG GG = = ⋛ C 8 GT G B CT



Results reviewed and confirmed by:

Dr. Noam Pik BVs MDSV

George Sofronidis BSc (Hons)

EXPLANATION OF RESULTS

NORMAL (N) - no presence of mutation, clear of disease

CARRIER (C) - one copy of the disease gene is present, will not exhibit disease symptoms

animal may show symptoms. Also referred to as Positive Heterozygous AFFECTED (1 COPY) - One copy of the disease gene is present, yet due to the dominant mode of inheritance of the disease the

Appropriate treatment should be pursued by consulting a veterinarian. **AFFECTED (A)** - 2 copies of the disease gene are present, the animal may show symptoms associated with the disease.

information together with the history submitted for the parents excludes this animal from having this disease NORMAL By PARENTAGE HISTORY (NBPH) - the sample submitted has had its parentage confirmed. By definition, this

NRA (No Results Available) - Insufficient information has been provided to provide a result for this test. Sire and Dam information and/or sample may be required. Please contact the laboratory to discuss

at CSNP1 then it must inherit either a G or C from the dam and the other base from the sire. by green, T by black and A by red. Each SNP will exhibit 2 bases (one from each parent), so if an individual shows the bases GC At each SNP an individual will inherit a DNA base of G, A, T or C. These bases are also colour coded. G is represented by blue, C the single nucleotide polymorphism (SNP) at a particular site on the chromosome with each number representing a different site. **DNA Profile** - also known as a DNA fingerprint is unique for the animal. No animal shares the same DNA profile. An individual's DNA profile is inherited from both parents and can be used for verifying parentage (pedigrees). The nomenclature CSNP identifies

dry (stand up) for at least 3 minutes prior to placing them back into the original swab packaging advise that another sample be re-collected and submitted. To minimise bacterial contamination you should allow the swab to air FAIL - The sample submitted has failed to give a conclusive result. Failures are due mainly to quality/quantity of DNA. We strongly

PARENTAGE CONFIRMATION - Parentage (pedigree confirmation) can be carried using our web based confirmation tool. (www.asaplab.com.au/parentage)



Test Result

Sample ID: Result Date:

1301-W-02053 25/01/2013

UK Office

125 Northenden Road

Manchester M33 3HF

Tel: 0161 282 30 66 Fax: 0161 973 3434

Email: Info@laboklin.co.uk Website: www.laboklin.co.uk

Veterinary Surgeon

MARNIE CRILLY BVSC TIMARU'S FAMILY VET 9 LATTER ST TIMARU

Owner

RAEWYN LI-ANNE BANK

P.O. BOX 2058 WASHDYKE TIMARU 7910

NEW ZEALAND

Animal Details

NEW ZEALAND

Animal: Dog

Name HOTWYR LOCK STOCK AND BARREL

(WYATT)

Breed GERMAN WIREHAIRED POINTER

Sex MALE

D.O.B

30/04/2010

Microchip No.

941000014103318 Tattoo No.

KC Registration

04910-2010

Sample

Sample Material EDTA BLOOD Sample Date: 05/01/2013

Test

Test Name:

80142 von Willerbrand disease type II

Result

Genotype:

Interpretation:

N/N (Clear)

The dog is homozygous normal regarding the intact gene.

The dog is a noncarrier of the defect gene (von

Willebrand Disease).

The dog is genetically clear and will not be affected by vWDII.

The dog can pass only the normal gene onto its entire offspring.

The result is only valid for the submitted sample.

Dept. Molecular Biology **LABOKLIN**

DIN ISO 17025 ACCREDITTED LABORATORY