



Orivet

Genetic Comprehensive Report

Animal Name: Spade

Owner:

Dwight E Herron II

Membership Number : Not assigned

Member Body/Breed Club: Not assigned

Approved Collection Method: No



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Accredited and Compliant with



Members of



Harmonization of
Genetic Testing
for Dogs



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Owner's details

Name: Dwight E Herron II

Animal's Details

Registered Name : Creek Veiv Cowco's Ace In The Hole

Pet Name : Spade

Registration Number :

Breed : Australian Cattle Dog

Microchip Number :

Sex : Intact Male

Date of Birth : 20th Nov 2021

Colour : Blue

Sample Collection Details

Case Number : 22218397

Collected By :

Approved Collection : No

Sample Type : SWAB

Test Details

Test Requested : Australian Cattle Dog - Full Breed Profile

Pet Name : Spade

Date of Test : 19th Aug 2022

Authorisation

Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported:

George Sofronidis BSc (Hons)

Dr Noam Pik BVSc, MAVS



Genetic Comprehensive Report



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Animal's Details

Registered Name : Creek Veiw Cowco's Ace In The Hole

Pet Name : Spade

Registration Number :

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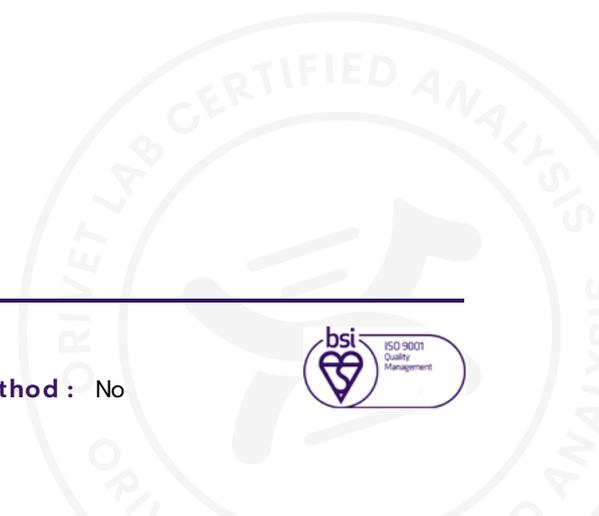
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Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No



Genetic Comprehensive Report

Animal's Details

Registered Name :	Creek Veiv Cowco's Ace In The Hole
Pet Name :	Spade
Registration Number :	
Breed :	Australian Cattle Dog
Microchip Number :	
Sex :	Intact Male
Date of Birth :	20th Nov 2021
Colour :	Blue

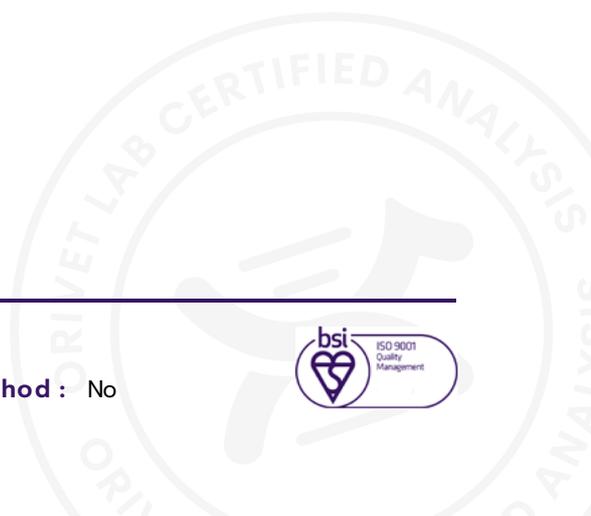
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Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No



Genetic Comprehensive Report

Animal's Details

Registered Name :	Creek Veiw Cowco's Ace In The Hole
Pet Name :	Spade
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Breed :	Australian Cattle Dog
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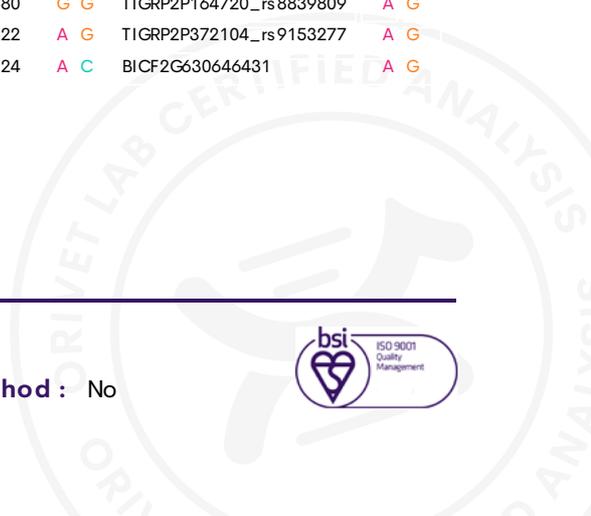
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Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No





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Genetic Comprehensive Report

Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported

Test Reported : CYSTINURIA (SLC3A1) (AUSTRALIAN CATTLE DOG TYPE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Solute carrier family 3 member 1 (SLC3A1) on chromosome 10

Variant Detected : Nucleotide Deletionc.1095-1100delp.366-367Thr 6bpdeletion (inframe)

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : DEGENERATIVE MYELOPATHY

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Superoxide dismutase 1 (SOD1) on chromosome 31

Variant Detected : Base Substitutionc.118G>Ap.Glu40Lys

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : HYPERURICOSURIA

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Solute carrier family 2 member 9 (SLC2A9) on chromosome 3

Variant Detected : Base Substitutionc.563G>Tp.Cys188Phe

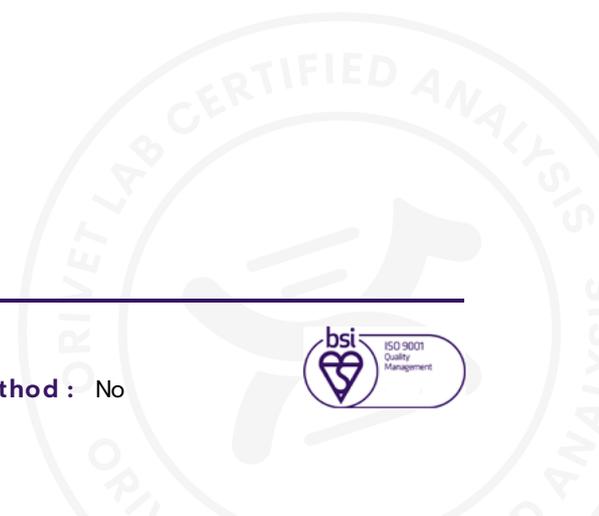
We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

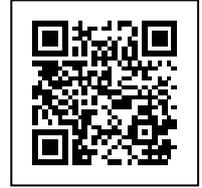
Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No





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Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported

Test Reported : IVERMECTIN SENSITIVITY MDR1 (MULTI DRUG RESISTANCE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : MDR1 on Chromosome 14

Variant Detected : Deletion 4bp AGAT

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : MYOTONIA HEREDITARIA (CATTLE DOG TYPE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Chloride voltage-gated channel 1 (CLCN1) on chromosome 16

Variant Detected : c.2703-2704 insertion Ap.Arg890Gln-frameshift888

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : NEURONAL CEROID LIPOFUSCINOSIS 5 (BORDER COLLIE TYPE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : CLN5, intracellular trafficking protein (CLN5) on Chromosome 22

Variant Detected : Base Substitutionc.619C>Tp.Glu206STOP

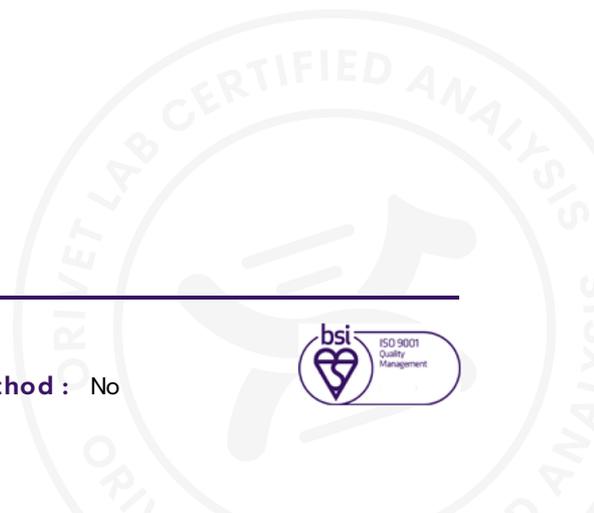
We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Owner's Name : Dwight E Herron II

Pet Name : Spade

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Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported

Test Reported : NEURONAL CEROID LIPOFUSCINOSIS NCL 12 (CATTLE DOG TYPE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : ATP

Variant Detected : c.1118C?>?T

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : PRIMARY LENS LUXATION

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : ADAM metallopeptidase with thrombospondin type 1 motif 17 (ADAMTS17) on Chromosome 3

Variant Detected : Base Substitutionc.1473+1G>Asplice-donor-site mutation at the 5' end of intron 10

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : PROGRESSIVE ROD CONE DEGENERATION (PRCD) - PRA

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Photoreceptor disc component (PRCD) on Chromosome 9

Variant Detected : Base Substitutionc.5 G>Ap.Cys2Tyr

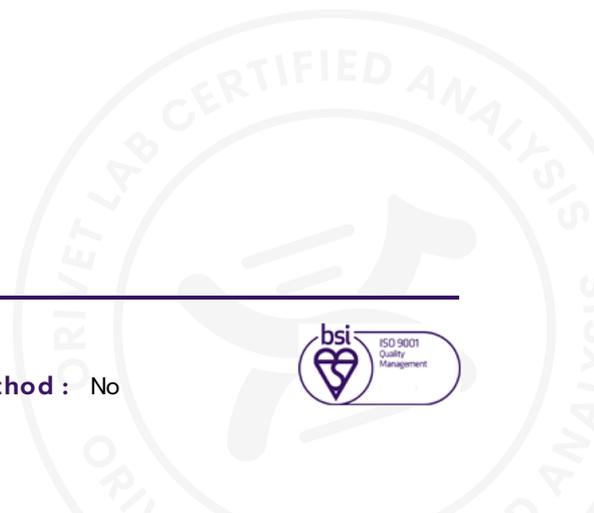
We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

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Test Reported : RCD4-PRA (LATE ONSET)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : C2orf71 on Chromosome 17

Variant Detected : c.3149_3150insCp.Cys1051ValfsX90

We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Test Reported : E LOCUS - (CREAM/RED/YELLOW)

Result : **E/E - DOMINANT BLACK DOES NOT CARRY YELLOW/RED/WHITE**¹

Gene : MC1R

Variant Detected : Em (point mutation) > E (wild type) > e (point mutation) chr5:63694334-63694334: C>T

2 copies of black E or "extension". All areas of the coat colour eumelanin will not produce any "e" offspring. The Extension loci is responsible for the majority of non-agouti patterns.

Test Reported : E LOCUS (CATTLE DOG CREAM VARIANT) E2

Result : **E²/E² - DOMINANT BLACK DOES NOT CARRY "AUSTRALIAN CATTLE DOG" TYPE CREAM**¹

Gene : MC1R

Variant Detected : c.430G>C

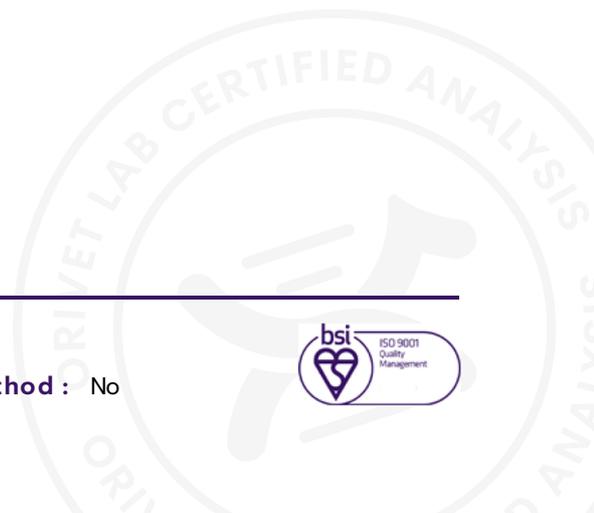
This e2 variant is associated with the pale cream coat colour seen in the Australian cattle dog and other varieties or breeds of common ancestry.

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Test Reported : EM (MC1R) LOCUS - MELANISTIC MASK

Result : Eⁿ/Eⁿ - NO MELANISTIC MASK (Eⁿ) EXTENSION ALLELE¹

Gene : MC1R

Variant Detected : Base Substitution G>A

Dog tested negative for the melanistic mask allele. The dog will not have a black mask, and cannot pass a copy on to any offspring..

Test Reported : BROWN (345DELPRO) DELETION

Result : B^d/B^d - DOES NOT CARRY BROWN/RED/LIVER or CHOCOLATE [DELETION]¹

Gene : TYRP1

Variant Detected : Base Substitution (Point Mutation)

Does not carry the brown deletion codon. Please refer to the other brown variants to clarify potential colour for offspring.

Test Reported : BROWN (GLNT331STOP) STOP CODON

Result : B^s/B^s - DOES NOT CARRY BROWN/RED/LIVER or CHOCOLATE [STOP CODON]¹

Gene : TYRP1

Variant Detected : Point Mutation

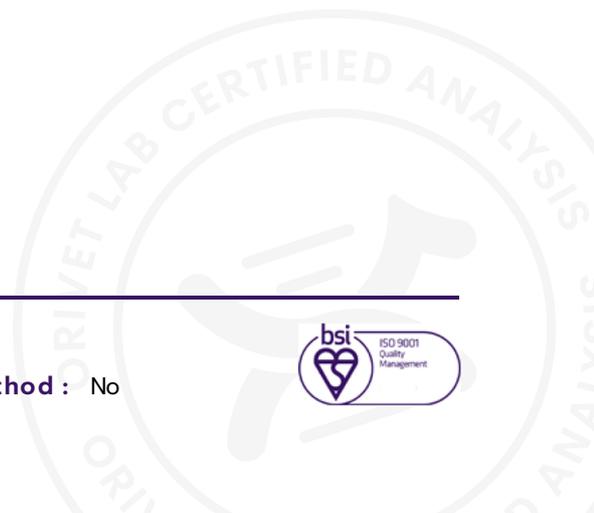
Does not carry the brown stop codon. Please refer to the other brown variants to clarify potential colour for offspring.

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Test Reported : BROWN (SER41CYS) INSERTION CODON

Result : B^c/B^c - DOES NOT CARRY BROWN/RED/LIVER or CHOCOLATE [INSERTION]¹

Gene : TYRP1

Variant Detected : Base Substitution (Point Mutation)

Does not carry the brown insertion codon. Please refer to the other brown variants to clarify potential colour for offspring.

Test Reported : LIVER [TYRP1] (LANCASHIRE HEELER TYPE)

Result : B^e/B^e - DOES NOT CARRY BROWN/LIVER [TYRP1]¹

Gene :

Variant Detected :

Test Reported : D (DILUTE) LOCUS

Result : D/D - NO COPY OF MLPH-D ALLELE (DILUTE) - PIGMENT IS NORMAL¹

Gene : MLPH

Variant Detected : Base Substitution

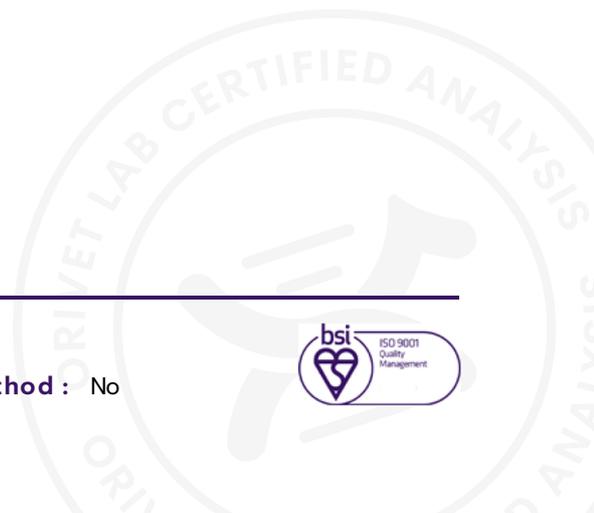
Full colour, no dilute gene present. The D allele modifies the Melanophillin (MLPH) gene. This animal cannot produce "dilute" offspring. Please Note: There are other dilute variants d2 (Sloughi, Chow Chow & Thai Ridgeback) and rare d3 (Italian Greyhound & Chihuahua) so this test/result may not identify dilute in these breeds.

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Test Reported : K LOCUS (DOMINANT BLACK)

Result : k^y/k^y - RECESSIVE NON- BLACK [COLOUR PATTERN DETERMINED BY A LOCUS]¹

Gene : CBD103

Variant Detected : Deletion of GGG

Dog does not have the dominant black mutation. Dog's coat colour will be determined by the agouti gene – may be brindled or not brindled. Any pheomelanin (red/tan) will be brindled. Can be sable/fawn, tricolour, tan points, black or brown. Will (may) have black pigment and black markings (unless the extension locus interferes).

Test Reported : A LOCUS (FAWN/SABLE;TRI/TAN POINTS)

Result : a^t/a^t - TAN POINTS/BLACK & TAN or TRICOLOUR MAY BE BRINDLED [SEE K LOCUS]¹

Gene : ASIP

Variant Detected : Base Substitution 246 G>T(A82S); G>A (R83H): C>T (p.R96C)

Homozygous for black and tan/tricolour (no hidden colours) allele. Tri factored/white factored in dogs that have white points. No Bi Factoring (Black White & Tan). Animals are primarily black and have areas of pheomelanin (tan) which tends to be seen on the leg and stomach areas, the side of the head and spots above the eyes. Please note the colour and distribution of pheomelanin "tan" will be dependent on the breed and other colour genes. Please note that any genes on the "A" series will only be expressed if the K locus is kk, kkbr or kbrkbr.

Test Reported : LONG HAIR GENE (CANINE C95F)

Result : NEGATIVE - NOT SHOWING THE PHENOTYPE¹

Gene : FGF5

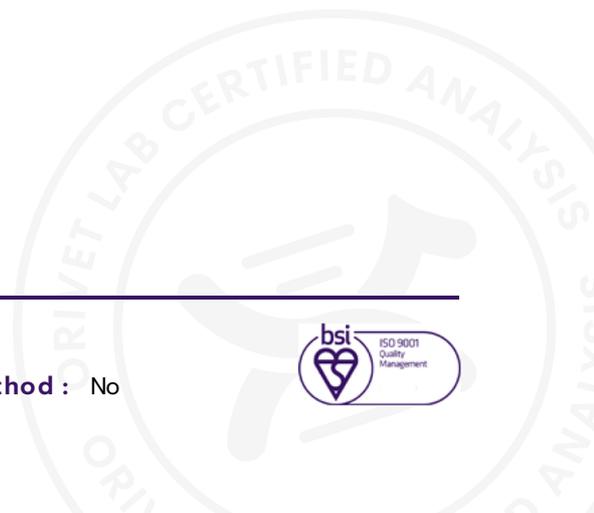
Variant Detected : p.Cys95Phe c284G>T (Point Mutation)

Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No





Scan to authenticate
this Report online

Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported

Test Reported : SHEDDING (MC5R)

Result :

shd/shd [HIGH SHEDDING] - TWO COPIES OF THE shd (MC5R) VARIANT DETECTED REFER TO R151W (IC) FOR LEVEL OF SHEDDING

1

Gene : MC5R

Variant Detected :

The dog will (may) exhibit a low level of shedding. Please Note: this level is also dependent on the furnishing allele. If the dog has no IC (R151W) phenotype will be low shedding.

Test Reported : COAT COMPOSITION CFA28 GENE (DOUBLE/SINGLE COAT)

Result : udc/udc - TWO COPIES OF THE DOUBLE COAT (DENSE UNDERCOAT) PHENOTYPE DETECTED¹

Gene : CFA28

Variant Detected :

Dog will have a shorter outer coat and will be dense and woolly in texture.

Test Reported : CURLY COAT/HAIR CURL (KRT71 R151W)

Result :

NEGATIVE FOR THE KRT71 R151W (CU/CU) VARIANT - NOT SHOWING THE CURLY COAT PHENOTYPE

1

Gene : KRT71 (R151W)

Variant Detected : chr27:2539211-2539211: c.451C>T

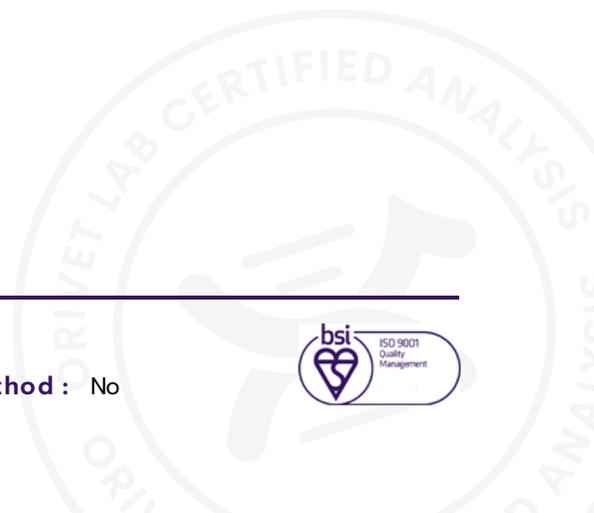
Please note there are other additional curly coat genes/variant that will impact the curly coat phenotype.

Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No





Genetic Comprehensive Report



Scan to authenticate
this Report online

Sample with Lab ID Number 22218397 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported

Test Reported : NATURAL BOB TAIL (SHORT TAIL PHENOTYPE)

Result : **NEGATIVE / CLEAR [NO VARIANT DETECTED]**¹

Gene : Gene: T on Chromosome 1

Variant Detected : Base Substitution C>G

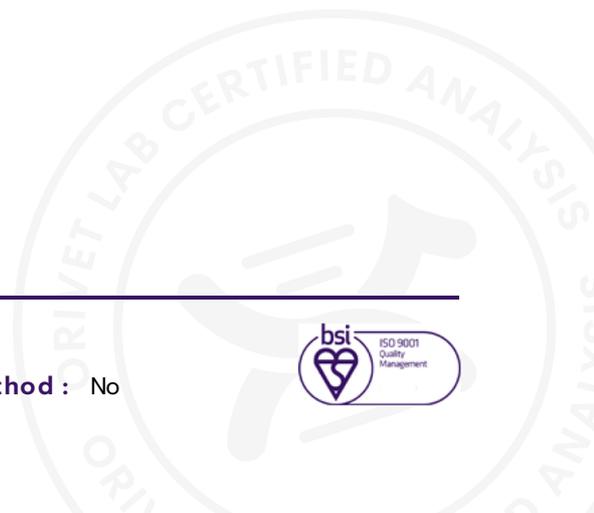
We have scanned the DNA and the genotype of this animal is NORMAL - no presence of the disease associated variant (mutation) has been detected. This result may also be referred to as NORMAL, "-/-" or "wild type (WT)" or "homozygous negative". The animal is clear of the disease and will not pass on the disease-causing variant.

Owner's Name : Dwight E Herron II

Pet Name : Spade

Microchip Number

Approved Collection Method : No



Glossary of Genetic Terms (Results)



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NEGATIVE / CLEAR [NO VARIANT DETECTED]

No presence of the variant (mutation) has been detected. The animal is clear of the disease and will not pass on any disease-causing mutation.

CARRIER [ONE COPY OF THE VARIANT DETECTED]

This is also referred to as HETEROZYGOUS. One copy of the normal gene and copy of the affected (mutant) gene has been detected. The animal will not exhibit disease symptoms or develop the disease. Consideration needs to be taken if breeding this animal - if breeding with another carrier or affected or unknown then it may produce an affected offspring.

POSITIVE / AT RISK [TWO COPIES OF THE VARIANT DETECTED]

Two copies of the disease gene variant (mutation) have been detected also referred to as HOMOZYGOUS for the variant. The animal may show symptoms (affected) associated with the disease. Appropriate treatment should be pursued by consulting a Veterinarian.

POSITIVE HETEROZYGOUS [ONE COPY OF THE DOMINANT VARIANT DETECTED]

Also referred to as POSITIVE ONE COPY or POSITIVE HETEROZYGOUS. This result is associated with a disease that has a dominant mode of inheritance. One copy of the normal gene (wild type) and affected (mutant) gene is present. Appropriate treatment should be pursued by consulting a Veterinarian. This result can still be used to produce a clear offspring.

NORMAL BY PARENTAGE HISTORY

The sample submitted has had its parentage verified by DNA. By interrogating the DNA profiles of the Dam, Sire and Offspring this information together with the history submitted for the parents excludes this animal from having this disease. The controls run confirm that the dog is NORMAL for the disease requested.

NORMAL BY PEDIGREE

The sample submitted has had its parentage verified by Pedigree. The pedigree has been provided and details (genetic testing reports) of the parents have been included. Parentage could not be determined via DNA profile as no sample was submitted.

NO RESULTS AVAILABLE

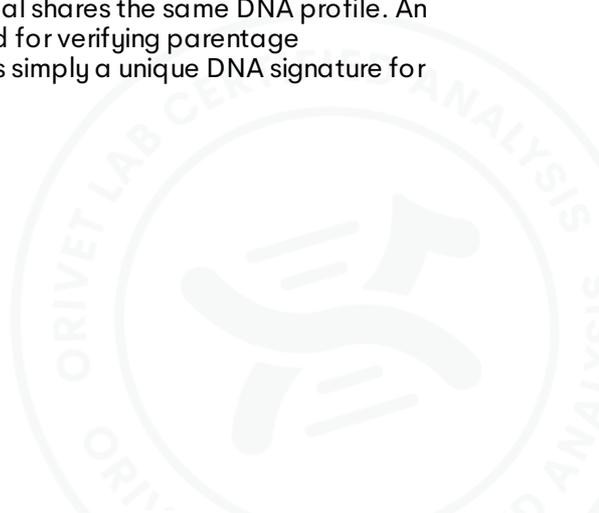
Insufficient information has been provided to provide a result for this test. Sire and Dam information and/or sample may be required. This result is mostly associated with tests that have a patent/license and therefore certain restrictions apply. Please contact the laboratory to discuss.

INDETERMINABLE

The sample submitted has failed to give a conclusive result. This result is mainly due to the sample failing to "cluster" or result in the current grouping. A recollection is required at no charge.

DNA PROFILE

Also known as a DNA fingerprint. This 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). This profile contains no disease or trait information and is simply a unique DNA signature for that animal.



Glossary of Genetic Terms (Results)



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PARENTAGE VERIFICATION/ QUALIFIES/CONFIRMED OR DOES NOT QUALIFY/EXCLUDED

Parentage is determined by examining the markers on the DNA profile. A result is generated and stated for all DNA parentage requests. Parentage confirmation reports can only be generated if a DNA profile has been carried out for Dam, Offspring and possible Sire/s.

PENDING

PENDING

TRAIT (PHENOTYPE)

A feature that an animal is born with (a genetically determined characteristic). Traits are a visual phenotype that range from colour to hair length, and also includes certain features such as tail length. If an individual is **AFFECTED** for a trait then it will show that characteristic eg. **AFFECTED** for the B (Brown) Locus or bb will be brown/chocolate.

POSITIVE – SHOWING THE PHENOTYPE

The animal is showing the trait or phenotype tested.

CLARIFICATION OF GENETIC TESTING

The goal of genetic testing is to provide breeders with relevant information to improve breeding practices in the interest of animal health. However, genetic inheritance is not a simple process, and may be complicated by several factors. Below is some information to help clarify these factors.

The goal of genetic testing is to provide breeders with relevant information to improve breeding practices in the interest of animal health. However, genetic inheritance is not a simple process, and may be complicated by several factors. Below is some information to help clarify these factors.

- 1) Some diseases may demonstrate signs of what Geneticists call "genetic heterogeneity". This is a term to describe an apparently single condition that may be caused by more than one mutation and/or gene
- 2) It is possible that there exists more than one disease that presents in a similar fashion and segregates in a single breed. These conditions -although phenotypically similar - may be caused by separate mutations and/or genes.
- 3) It is possible that the disease affecting your breed may be what Geneticists call an "oligogenic disease". This is a term to describe the existence of additional genes that may modify the action of a dominant gene associated with a disease. These modifier genes may for example give rise to a variable age of onset for a particular condition, or affect the penetrance of a particular mutation such that some animals may never develop the condition.

The range of hereditary diseases continues to increase and we see some that are relatively benign and others that can cause severe and/or fatal disease. Diagnosis of any disease should be based on pedigree history, clinical signs, history (incidence) of the disease and the specific genetic test for the disease. Penetrance of a disease will always vary not only from breed to breed but within a breed, and will vary with different diseases. Factors that influence penetrance are genetics, nutrition and environment. Although genetic testing should be a priority for breeders, we strongly recommend that temperament and phenotype also be considered when breeding.

Orivet Genetic Pet Care aims to frequently update breeders with the latest research from the scientific literature. If breeders have any questions regarding a particular condition, please contact us on (03) 9534 1544 or admin@orivet.com and we will be happy to work with you to answer any relevant questions.

