



Genetic Comprehensive Report

Animal Name: Archie

Owner:

Maria Bryan

Membership Number : 072019

Member Body/Breed Club : Orivet Breeders Club

Approved Collection Method : YES



orivet.com

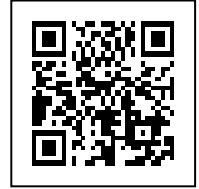
Accredited and Compliant with



Members of



Harmonization of
Genetic Testing
for Dogs



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Owner's details

Name : Maria Bryan

Animal's Details

Registered Name :

Pet Name : Archie

Registration Number :

Breed : Boston Terrier

Microchip Number : 953010005543300

Sex : Male

Date of Birth : 10th Apr 1921

Colour : Brindle and white

Sample Collection Details

Case Number : 22M050006

Collected By : LT4014

Approved Collection : YES

Sample Type : SWAB

Test Details

Test Requested : Boston Terrier – Full Breed Profile

Pet Name : Archie

Date of Test : 1st Apr 2022

Authorisation

Sample with Lab ID Number 22M050006 was received at Orivet Genetics, DNA was extracted and analysed with the following result reported:

George Sofronidis BSc (Hons)

Dr Noam Pik BVSc, MAVS





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Animal's Details

Registered Name :

Pet Name : Archie

Registration Number :

Breed : Boston Terrier

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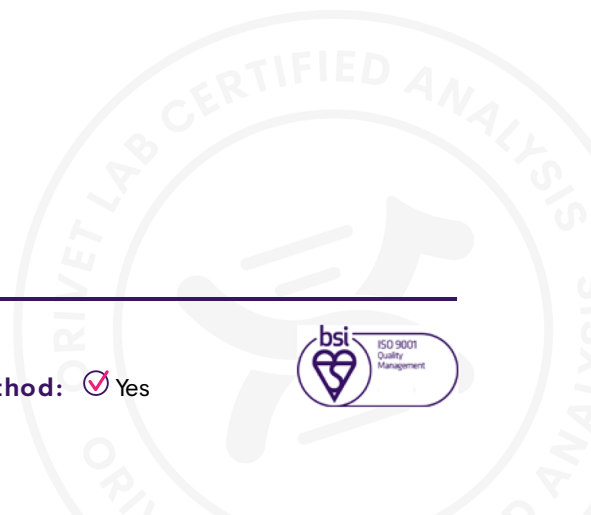
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 P19_1 A T P19_2 G G P5_2 G G P19_3 G G P2_1 G G P2_3 A C P27_3 T T P20_1 A A
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Owner's Name : Maria Bryan

Pet Name : Archie

Microchip Number : 953010005543300

Approved Collection Method: Yes



Genetic Comprehensive Report

Animal's Details

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| Registered Name : | |
| Pet Name : | Archie |
| Registration Number : | |
| Breed : | Boston Terrier |
| Microchip Number : | 953010005543300 |
| Sex : | Male |
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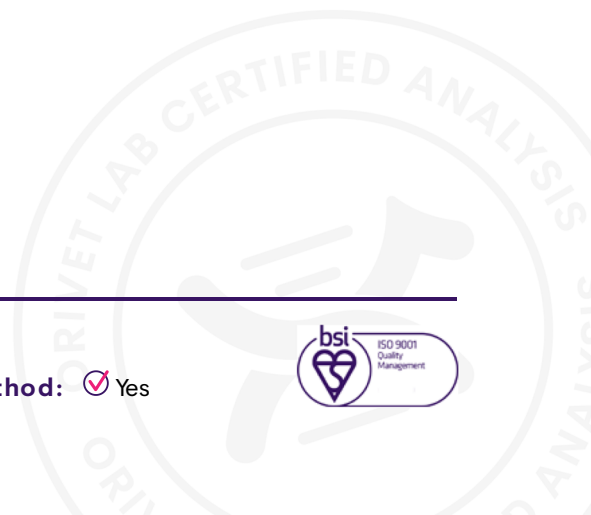
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| BICF2S23214514 | A C | BICF2S23326150 | G G | BICF2S23329382 | C C | BICF2S23357186 | G G |
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| TIGRP2P362535_rs9130694 | A G | TIGRP2P389035_rs9038546 | A G | | | | |

Owner's Name : Maria Bryan

Pet Name : Archie

Microchip Number : 953010005543300

Approved Collection Method: Yes



Genetic Comprehensive Report

Animal's Details

| | |
|-----------------------|-------------------|
| Registered Name : | |
| Pet Name : | Archie |
| Registration Number : | |
| Breed : | Boston Terrier |
| Microchip Number : | 953010005543300 |
| Sex : | Male |
| Date of Birth : | 10th Apr 1921 |
| Colour : | Brindle and white |

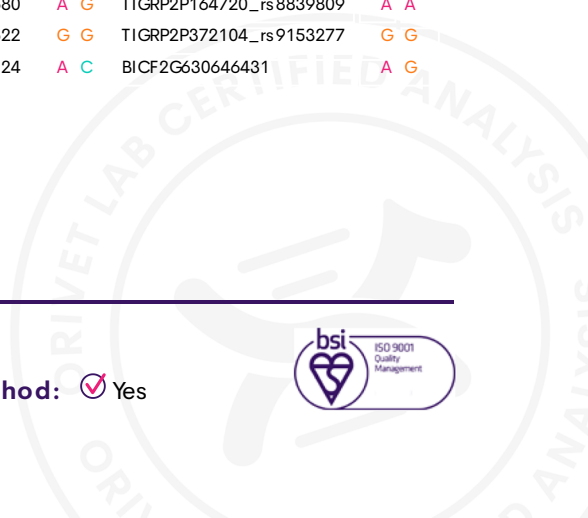
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| BICF2P878175 | G G | BICF2P935470 | A G | BICF2P990814 | G G | BICF2S22910736 | A A |
| BICF2S22913753 | G G | BICF2S22928800 | A A | BICF2S22943825 | G G | BICF2S23028732 | A T |
| BICF2S23031254 | A C | BICF2S23049416 | A A | BICF2S23057560 | A A | BICF2S23124313 | G G |
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| TIGRP2P177606_rs8886563 | C C | TIGRP2P215708_rs8686029 | T T | TIGRP2P316532_rs8597522 | G G | TIGRP2P372104_rs9153277 | G G |
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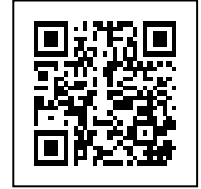
Owner's Name : Maria Bryan

Pet Name : Archie

Microchip Number : 953010005543300

Approved Collection Method: Yes





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Sample with Lab ID Number 22M050006 was received at Orivet Genetics,
DNA was extracted and analysed with the following result reported

Test Reported : HEREDITARY CATARACT

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

Gene : Heat shock transcription factor 4 (HSF4) on Chromosome 5

Variant Detected :

Nucleotide Insertion and Nucleotide Deletion c.971-972insC (Staffordshire Bull Terrier and Boston Terrier, French Bulldogs) c.971-972delC (Australian Shepherd) p.Pro324Profs27X (Staffordshire Bull Terrier and Boston Terrier, French Bulldogs) p.Pro324Hisfs86X (Australian Shepherd)

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 : 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.

Owner's Name : Maria Bryan

Pet Name : Archie

Microchip Number : 953010005543300

Approved Collection Method: Yes



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Sample with Lab ID Number 22M050006 was received at Orivet Genetics,
DNA was extracted and analysed with the following result reported

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.

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 :

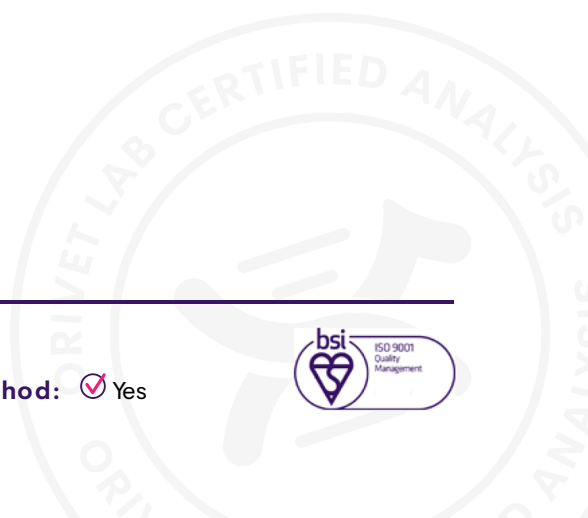
Variant Detected :

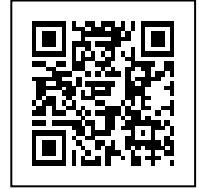
Owner's Name : Maria Bryan

Pet Name : Archie

Microchip Number : 953010005543300

Approved Collection Method: Yes





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Sample with Lab ID Number 22M050006 was received at Orivet Genetics,
DNA was extracted and analysed with the following result reported

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.

Test Reported : K LOCUS (DOMINANT BLACK)**Result :****KB / k^y or k^{br} - ONE COPY DOMINANT BLACK (KB) and ONE COPY OF NON-BLACK (k^y) dog MAY be brindled****1****Gene : CBD103****Variant Detected : Deletion of GGG**

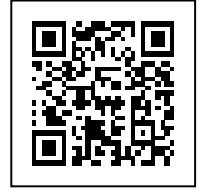
One copy of non black and one copy of ky or kbr is present. This KB will cover the A locus and all you will visualise is the base colour. Dog will express the alleles on the A locus but any and all phaeomelanin (red) in the coat will be brindled. This allele overrides the ASIP (A) locus. The agouti phenotype may be altered for some breeds and therefore be brindle. There are three alleles at the K Locus with the following dominance hierarchy KB > Kbr > k. The first KB represents dominant black, the second allele Kbr represents brindling and may display A locus gene. Brindle in most breeds appears as black stripes on a red base. Please Note: At this stage no commercial genetic testing can distinguish brindle so breeders should rely on their pedigree or breed standard to exclude or include brindle phenotype.

Test Reported : A LOCUS (FAWN/SABLE; TRI/TAN POINTS)**Result : a^y/a^y - FAWN/RED or SABLE only PRODUCE ay OFFSPRING¹****Gene : ASIP****Variant Detected : Base Substitution 246 G>T(A82S); G>A (R83H): C>T (p.R96C)**

Homozygous for fawn/sable (no hidden colour). Also referred to as "clear red". Pure factoring/no white factoring. Please note that the colour will be dependent on the breed and other colour genes. The colour shown is dependent on the E, K and B Locus.

Owner's Name : Maria Bryan**Pet Name :** Archie**Microchip Number :** 953010005543300**Approved Collection Method:** Yes

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Sample with Lab ID Number 22M050006 was received at Orivet Genetics,
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Test Reported : LONG HAIR GENE (CANINE C95F)

Result : **NEGATIVE - NOT SHOWING THE PHENOTYPE**¹

Gene : FGF5

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

Test Reported : SHEDDING (MC5R)

Result :

SHD/shd [MODERATE SHEDDING] - ONE COPY OF THE SHD (MC5R) VARIANT DETECTED [REFER TO R151W (IC) FOR LEVEL]

1

Gene : MC5R

Variant Detected :

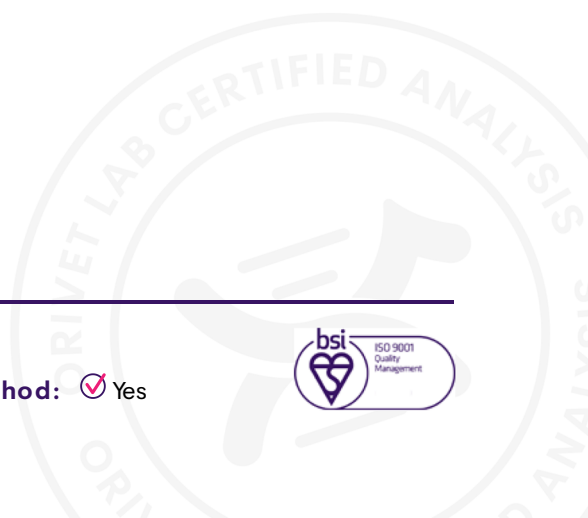
The dog will (may) exhibit a moderate (average) level of shedding. Please Note: this level is also dependent on the furnishing allele. If the dog has no IC (R151W) phenotype will be low shedding.

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Glossary of Genetic Terms (Results)



The terms below are provided to help clarify certain results phrases on your genetic report. The phrases below are those as reported by Orivet and may vary from one laboratory to the other.

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

Results for this test are still being processed. Some tests are run independently and are reported at a later date. When completed, the result will be emailed. APPROVED COLLECTION METHOD (NO) The sample submitted for testing HAS NOT met the requirements recommended by member bodies for the DNA collection process.

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.

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- 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.