

DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### **BREED MIX**

German Shepherd Dog: 100.0%

### **GENETIC STATS**

Wolfiness: 0.3 % **LOW**Predicted adult weight: **72 lbs** 

#### **TEST DETAILS**

Kit number: EM-49349762 Swab number: 31211050111420

Registration: N/A IHR 2231041 ★embark Microchip: 250268780939543

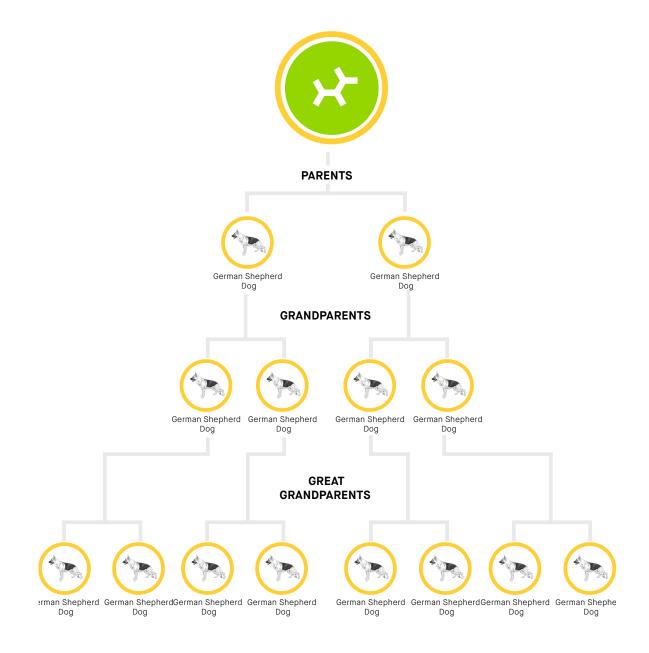


**DNA Test Report** 

Test Date: January 17th, 2024

embk.me/uandmeuriolesgriffesdesterressauva

### **FAMILY TREE**





DNA Test Report Test Date: January 17th, 2024

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#### **GERMAN SHEPHERD DOG**

The German Shepherd dog is the second most popular dog breed in the United States, and the fourth most popular in the United Kingdom (where it is known as the Alsatian). This breed was standardized in Germany at the end of the 19th century from local dogs used for herding and livestock guarding. Their confidence, courageousness and keen sense of smell coupled with their notable intelligence make them highly suited to police work, military roles, and search and rescue. German Shepherds require regular physical and mental exercise and have a heavy shedding coat that comes in both short and long varieties. They were first recognized by the AKC in 1908 and later became fashionable as soldiers returning from WWI spoke highly of the German dogs and Hollywood popularized the breed with stars like Strongheart and Rin Tin Tin.

#### **Fun Fact**

Despite being sometimes called the "Alsatian wolf dog", German Shepherds are not true wolf dogs— they are 100% dog. Nevertheless, German shepherds were crossed with wolves in the past to form the Czechoslovakian and Saarloos wolfdog breeds. German Shepherds, along with other breeds and sled dogs, were also used in the creation of the Chinook breed.

#### **RELATED BREEDS**



White Shepherd Sibling breed



Belgian Sheepdog Cousin breed



Belgian Tervuren Cousin breed



Belgian Malinois
Cousin breed



**Dutch Shepherd**Cousin breed

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DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### MATERNAL LINE



Through U and me Urio Les Griffes des Terres Sauvages's mitochondrial DNA we can trace his mother's ancestry back to where dogs and people first became friends. This map helps you visualize the routes that his ancestors took to your home. Their story is described below the map.

#### HAPLOGROUP: A1b

This female lineage was very likely one of the original lineages in the wolves that were first domesticated into dogs in Central Asia about 15,000 years ago. Since then, the lineage has been very successful and travelled the globe! Dogs from this group are found in ancient Bronze Age fossils in the Middle East and southern Europe. By the end of the Bronze Age, it became exceedingly common in Europe. These dogs later became many of the dogs that started some of today's most popular breeds, like German Shepherds, Pugs, Whippets, English Sheepdogs and Miniature Schnauzers. During the period of European colonization, the lineage became even more widespread as European dogs followed their owners to farflung places like South America and Oceania. It's now found in many popular breeds as well as village dogs across the world!

#### HAPLOTYPE: A361/409/611

Part of the A1b haplogroup, this haplotype occurs most frequently in German Shepherd Dogs, Poodles, and Shiloh Shepherds.

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DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### PATERNAL LINE



Through U and me Urio Les Griffes des Terres Sauvages's Y chromosome we can trace his father's ancestry back to where dogs and people first became friends. This map helps you visualize the routes that his ancestors took to your home. Their story is described below the map.

#### HAPLOGROUP: A1a

Some of the wolves that became the original dogs in Central Asia around 15,000 years ago came from this long and distinguished line of male dogs. After domestication, they followed their humans from Asia to Europe and then didn't stop there. They took root in Europe, eventually becoming the dogs that founded the Vizsla breed 1,000 years ago. The Vizsla is a Central European hunting dog, and all male Vizslas descend from this line. During the Age of Exploration, like their owners, these pooches went by the philosophy, "Have sail, will travel!" From the windy plains of Patagonia to the snug and homey towns of the American Midwest, the beaches of a Pacific paradise, and the broad expanse of the Australian outback, these dogs followed their masters to the outposts of empires. Whether through good fortune or superior genetics, dogs from the A1a lineage traveled the globe and took root across the world. Now you find village dogs from this line frolicking on

#### **HAPLOTYPE: H1a.15**

Part of the large A1a haplogroup, this haplotype is found in village dogs from across the globe (outside of Asia). As for breeds, it is primarily seen in German Shepherds, Labrador Retrievers, Nova Scotia Duck Tolling Retriever. It is by far the most common haplotype in German Shepherds.

Registration: N/A IHR 2231041 \rightarrow\text{embark} Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### TRAITS: BASE COAT COLOR

TRAIT RESULT

Dark or Light Fur | E (Extension) Locus | Gene: Melanocortin Receptor 1 (MC1R) | Genetic Result: E<sup>m</sup>E

This gene helps determine whether a dog can produce dark (black or brown) hairs or lighter yellow or red hairs. Any result except for **ee** means that the dog can produce dark hairs. An **ee** result means that the dog does not produce dark hairs at all, and will have lighter yellow or red hairs over their entire body.

Can have dark fur

**Did You Know?** If a dog has a **ee** result then the fur's actual shade can range from a deep copper to yellow/gold to cream - the exact color cannot be predicted solely from this result, and will depend on other genetic factors.

Dark brown pigment | Cocoa | Gene: HPS3 | Genetic Result: NN

Dogs with the **coco** genotype will produce dark brown pigment instead of black in both their hair and skin. Dogs with the **Nco** genotype will produce black pigment, but can pass the **co** variant on to their puppies. Dogs that have the **coco** genotype as well as the **bb** genotype at the B locus are generally a lighter brown than dogs that have the **Bb** or **BB** genotypes at the B locus.

No impact on fur and skin color

**Did You Know?** The **co** variant and the dark brown "cocoa" coat color have only been documented in French Bulldogs. Dogs with the cocoa coat color are sometimes born with light brown coats that darken as they reach maturity.

Red Pigment Intensity LINKAGE | I (Intensity) Loci | Genetic Result: Intermediate Red Pigmentation

Intensity refers to the concentration of red pigment in the coat. Dogs with more densely concentrated (intense) pigment will be a deeper red, while dogs with less concentrated (dilute) pigment will be tan, yellow, cream, or white. Five locations in the dog genome explain approximately 70% of red pigmentation intensity variation across all dogs. Because the locations we test may not directly cause differences in red pigmentation intensity, we consider this to be a linkage test.

Any light fur likely yellow or tan

**Did You Know?** One of the genes that influences pigment intensity in dogs, TYR, is also responsible for intensity variation in domestic mice, cats, cattle, rabbits, and Ilamas. In dogs and humans, more genes are involved.

Registration: 

→ Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### TRAITS: BASE COAT COLOR (CONTINUED)

TRAIT RESULT

Brown or Black Pigment | B (Brown) Locus | Gene: Tyrosinase Related Protein 1 (TYRP1) | Genetic Result: BB

This gene helps determine whether a dog produces brown or black pigments. Dogs with a **bb** result produce brown pigment instead of black in both their hair and skin, while dogs with a **Bb** or **BB** result produce black pigment. Dogs that have **ee** at the E (Extension) Locus and **bb** at this B (Brown) Locus are likely to have red or cream coats and brown noses, eye rims, and footpads, which is sometimes referred to as "Dudley Nose" in Labrador Retrievers.

Black or gray fur and skin

**Did You Know?** "Liver" or "chocolate" is the preferred color term for brown in most breeds; in the Doberman Pinscher it is referred to as "red".

Color Dilution | D (Dilute) Locus | Gene: Melanophilin (MLPH) | Genetic Result: DD

This gene helps determine whether a dog has lighter "diluted" pigment. A dog with a **Dd** or **DD** result will not be dilute. A dog with a **dd** result will have all their black or brown pigment lightened ("diluted") to gray or light brown, and may lighten red pigment to cream. This affects their fur, skin, and sometimes eye color. The D locus result that we report is determined by three different genetic variants that can work together to cause diluted pigmentation. These are the common **d** allele, also known as "**d1**", and the less common alleles known as "**d2**" and "**d3**". Dogs with two **d** alleles, regardless of which variant, are typically dilute.

Dark (non-dilute) fur and skin

**Did You Know?** There are many breed-specific names for these dilute colors, such as "blue", "charcoal", "fawn", "silver", and "Isabella". Dilute dogs, especially in certain breeds, have a higher incidence of Color Dilution Alopecia which causes hair loss in some patches.



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### TRAITS: COAT COLOR MODIFIERS

TRAIT RESULT

**Hidden Patterning** | K (Dominant Black) Locus | Gene: Canine Beta-Defensin 103 (CBD103) | Genetic Result: kyky

This gene helps determine whether the dog has a black coat. Dogs with a  $k^yk^y$  result will show a coat color pattern based on the result they have at the A (Agouti) Locus. A  $K^BK^B$  or  $K^BK^Y$  result means the dog is dominant black, which overrides the fur pattern that would otherwise be determined by the A (Agouti) Locus. These dogs will usually have solid black or brown coats, or if they have ee at the E (Extension) Locus then red/cream coats, regardless of their result at the A (Agouti) Locus. Dogs who test as  $K^Bk^Y$  may be brindle rather than black or brown.

More likely to have patterned fur

**Did You Know?** Even if a dog is "dominant black" several other genes could still impact the dog's fur and cause other patterns, such as white spotting.

Body Pattern | A (Agouti) Locus | Gene: Agouti Signalling Protein (ASIP) | Genetic Result: ata

This gene is responsible for causing different coat patterns. It only affects the fur of dogs that do not have **ee** at the E (Extension) Locus and do have **k**<sup>y</sup>**k**<sup>y</sup> at the K (Dominant Black) Locus. It controls switching between black and red pigment in hair cells, which means that it can cause a dog to have hairs that have sections of black and sections of red/cream, or hairs with different colors on different parts of the dog's body. Sable or Fawn dogs have a mostly or entirely red coat with some interspersed black hairs. Agouti or Wolf Sable dogs have red hairs with black tips, mostly on their head and back. Black and tan dogs are mostly black or brown with lighter patches on their cheeks, eyebrows, chest, and legs. Recessive black dogs have solid-colored black or brown coats.

Black/Brown and tan coat color pattern

**Did You Know?** The ASIP gene causes interesting coat patterns in many other species of animals as well as dogs.

Facial Fur Pattern | E (Extension) Locus | Gene: Melanocortin Receptor 1 (MC1R) | Genetic Result: EmE

In addition to determining if a dog can develop dark fur at all, this gene can give a dog a black "mask" or "widow's peak," unless the dog has overriding coat color genetic factors. Dogs with one or two copies of  $\mathbf{E}^{\mathbf{m}}$  in their result will have a mask, which is dark facial fur as seen in the German Shepherd and Pug. Dogs with no  $\mathbf{E}^{\mathbf{m}}$  in their result but one or two copies of  $\mathbf{E}^{\mathbf{g}}$  will instead have a "widow's peak", which is dark forehead fur.

Can have black masking (dark facial fur)

**Did You Know?** The widow's peak is seen in the Afghan Hound and Borzoi, where it is called either "grizzle" or "domino".

Registration: 

→ Cembark Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### TRAITS: COAT COLOR MODIFIERS (CONTINUED)

TRAIT RESULT

Saddle Tan | Gene: RALY | Genetic Result: NN

The "Saddle Tan" pattern causes the black hairs to recede into a "saddle" shape on the back, leaving a tan face, legs, and belly, as a dog ages. The Saddle Tan pattern is characteristic of breeds like the Corgi, Beagle, and German Shepherd. Dogs that have the II genotype at this locus are more likely to be mostly black with tan points on the eyebrows, muzzle, and legs as commonly seen in the Doberman Pinscher and the Rottweiler. This gene modifies the A Locus at allele, so dogs that do not express at are not influenced by this gene.

Likely saddle tan patterned

**Did You Know?** The Saddle Tan pattern is characteristic of breeds like the Corgi, Beagle, and German Shepherd.

White Spotting | S (White Spotting) Locus | Gene: MITF | Genetic Result: SS

This gene is responsible for most of the white spotting observed in dogs. Dogs with a result of **spsp** will have a nearly white coat or large patches of white in their coat. Dogs with a result of **Ssp** will have more limited white spotting that is breed-dependent. A result of **SS** means that a dog likely has no white or minimal white in their coat. The S Locus does not explain all white spotting patterns in dogs and other causes are currently being researched. Some dogs may have small amounts of white on the paws, chest, face, or tail regardless of their result at this gene.

Likely to have little to no white in coat

**Did You Know?** Any dog can have white spotting regardless of coat color. The colored sections of the coat will reflect the dog's other genetic coat color results.

Roan LINKAGE | R (Roan) Locus | Gene: USH2A | Genetic Result: Rr

This gene, along with the S Locus, regulates whether a dog will have roaning. Dogs with at least one copy of **R** will likely have roaning on otherwise uniformly unpigmented white areas created by the S Locus. Roan may not be visible if white spotting is limited to small areas, such as the paws, chest, face, or tail. The extent of roaning varies from uniform roaning to non-uniform roaning, and patchy, non-uniform roaning may look similar to ticking. Roan does not appear in white areas created by other genes, such as a combination of the E Locus and I Locus (for example, Samoyeds). The roan pattern can appear with or without ticking.

Likely no impact on coat pattern

**Did You Know?** Roan, tick, and Dalmatians' spots become visible a few weeks after birth. The R Locus is probably involved in the development of Dalmatians' spots.

Registration: 

→ Cembark Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### TRAITS: COAT COLOR MODIFIERS (CONTINUED)

TRAIT RESULT

Merle | M (Merle) Locus | Gene: PMEL | Genetic Result: mm

This gene is responsible for mottled or patchy coat color in some dogs. Dogs with an **M\*m** result are likely to appear merle or could be "non-expressing" merle, meaning that the merle pattern is very subtle or not at all evident in their coat. Dogs with an **M\*M\*** result are likely to have merle or double merle coat patterning. Dogs with an **mm** result are unlikely to have a merle coat pattern.

Unlikely to have merle pattern

**Did You Know?** Merle coat patterning is common to several dog breeds including the Australian Shepherd, Catahoula Leopard Dog, and Shetland Sheepdog.

Harlequin | Gene: PSMB | Genetic Result: hh

This gene, along with the M Locus, determines whether a dog will have harlequin patterning. This pattern is recognized in Great Danes and causes dogs to have a white coat with patches of darker pigment. A dog with an **Hh** result will be harlequin if they are also **M\*m** or **M\*M\*** at the M Locus and are not **ee** at the E locus. Dogs with a result of **hh** will not be harlequin.

No impact on coat pattern

**Did You Know?** While many harlequin dogs are white with black patches, some dogs have grey, sable, or brindle patches of color, depending on their genotypes at other coat color genes.

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**DNA Test Report** Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### TRAITS: OTHER COAT TRAITS

**RESULT TRAIT** 

Furnishings LINKAGE | Gene: RSP02 | Genetic Result: II

This gene is responsible for "furnishings", which is another name for the mustache, beard, and eyebrows that are characteristic of breeds like the Schnauzer, Scottish Terrier, and Wire Haired Dachshund. A dog with an FF or FI result is likely to have furnishings. A dog with an II result will not have furnishings. We measure this result using a linkage test.

Likely unfurnished (no mustache, beard, and/or eyebrows)

Did You Know? In breeds that are expected to have furnishings, dogs without furnishings are the exception - this is sometimes called an "improper coat".

Coat Length | Gene: FGF5 | Genetic Result: TT

This gene is known to affect hair/fur length in many different species, including cats, dogs, mice, and humans. In dogs, a TT result means the dog is likely to have a long, silky coat as seen in the Yorkshire Terrier and the Long Haired Whippet. A GG or GT result is likely to mean a shorter coat, like in the Boxer or the American Staffordshire Terrier.

Likely long coat

Did You Know? In certain breeds, such as Corgi, the long coat is described as "fluff."

Shedding | Gene: MC5R | Genetic Result: CC

This gene affects how much a dog sheds. Dogs with furnishings or wire-haired coats tend to be low shedders regardless of their result for this gene. In other dogs, a CC or CT result indicates heavy or seasonal shedding, like many Labradors and German Shepherd Dogs. Dogs with a TT result tend to be lighter shedders, like Boxers, Shih Tzus and Chihuahuas.

Likely heavy/seasonal shedding

Coat Texture | Gene: KRT71 | Genetic Result: CC

For dogs with long fur, dogs with a TT or CT result will likely have a wavy or curly coat like the coat of Poodles and Bichon Frises. Dogs with a CC result will likely have a straight coat—unless the dog has a "Likely Furnished" result for the Furnishings trait, since this can also make the coat more curly.

Likely straight coat

Did You Know? Dogs with short coats may have straight coats, whatever result they have for this gene.

Hairlessness (Xolo type) LINKAGE | Gene: FOX/3 | Genetic Result: NN

Registration: Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### TRAITS: OTHER COAT TRAITS (CONTINUED)

TRAIT RESULT

Hairlessness (Terrier type) | Gene: SGK3 | Genetic Result: NN

This gene is responsible for Hairlessness in the American Hairless Terrier. Dogs with the **DD** result are likely to be hairless. Dogs with the **ND** genotype will have a normal coat, but can pass the **D** variant on to their offspring.

Very unlikely to be hairless

Oculocutaneous Albinism Type 2 LINKAGE | Gene: SLC45A2 | Genetic Result: NN

This gene causes oculocutaneous albinism (OCA), also known as Doberman Z Factor Albinism. Dogs with a **DD** result will have OCA. Effects include severely reduced or absent pigment in the eyes, skin, and hair, and sometimes vision problems due to lack of eye pigment (which helps direct and absorb ambient light) and are prone to sunburn. Dogs with a **ND** result will not be affected, but can pass the mutation on to their offspring. We measure this result using a linkage test.

Likely not albino

**Did You Know?** This particular mutation can be traced back to a single white Doberman Pinscher born in 1976, and it has only been observed in dogs descended from this individual.

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DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### TRAITS: OTHER BODY FEATURES

TRAIT RESULT

Muzzle Length | Gene: BMP3 | Genetic Result: CC

This gene affects muzzle length. A dog with a **AC** or **CC** result is likely to have a medium-length muzzle like a Staffordshire Terrier or Labrador, or a long muzzle like a Whippet or Collie. A dog with a **AA** result is likely to have a short muzzle, like an English Bulldog, Pug, or Pekingese.

**Did You Know?** At least five different genes affect snout length in dogs, with BMP3 being the only one with a known causal mutation. For example, the muzzle length of some breeds, including the long-snouted Scottish Terrier or the short-snouted Japanese Chin, appear to be caused by other genes. This means your dog may have a long or short snout due to other genetic factors. Embark is working to figure out what these might be.

Likely medium or long muzzle

Tail Length | Gene: T | Genetic Result: CC

This is one of the genes that can cause a short bobtail. Most dogs have a **CC** result and a long tail. Dogs with a **CG** result are likely to have a bobtail, which is an unusually short or absent tail. This can be seen in many "natural bobtail" breeds including the Pembroke Welsh Corgi, the Australian Shepherd, and the Brittany Spaniel. Dogs with **GG** genotypes have not been observed, suggesting that dogs with such a result do not survive to birth.

Likely normal-length tail

**Did You Know?** While certain lineages of Boston Terrier, English Bulldog, Rottweiler, Miniature Schnauzer, Cavalier King Charles Spaniel, and Parson Russell Terrier, and Dobermans are born with a natural bobtail, it is not always caused by this gene. This suggests that other unknown genetic effects can also lead to a natural bobtail.

Hind Dew Claws | Gene: LMBR1 | Genetic Result: CC

This is one of the genes that can cause hind dew claws, which are extra, nonfunctional digits located midway between a dog's paw and hock. Dogs with a **CT** or **TT** result have about a 50% chance of having hind dewclaws. Hind dew claws can also be caused by other, still unknown, genes. Embark is working to figure those out.

Unlikely to have hind dew claws

Did You Know? Hind dew claws are commonly found in certain breeds such as the Saint Bernard.

Registration: 

Microchip: 250268780939543

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**DNA Test Report** Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### TRAITS: OTHER BODY FEATURES (CONTINUED)

**RESULT TRAIT** 

Back Muscling & Bulk (Large Breed) | Gene: ACSL4 | Genetic Result: CC

This gene can cause heavy muscling along the back and trunk in characteristically "bulky" large-breed dogs including the Saint Bernard, Bernese Mountain Dog, Greater Swiss Mountain Dog, and Rottweiler. A dog with the TT result is likely to have heavy muscling. Leaner-shaped large breed dogs like the Great Dane, Irish Wolfhound, and Scottish Deerhound generally have a CC result. The TC result also indicates likely normal muscling.

Likely normal muscling

Did You Know? This gene does not seem to affect muscling in small or even mid-sized dog breeds with lots of back muscling, including the American Staffordshire Terrier, Boston Terrier, and the English Bulldog.

Eye Color LINKAGE | Gene: ALX4 | Genetic Result: NN

This gene is associated with blue eyes in Arctic breeds like Siberian Husky as well as tri-colored (nonmerle) Australian Shepherds. Dogs with a DupDup or NDup result are more likely to have blue eyes, although some dogs may have only one blue eye or may not have blue eyes at all; nevertheless, they can still pass blue eyes to their offspring. Dogs with a NN result may have blue eyes due to other factors, such as merle or white spotting. We measure this result using a linkage test.

Less likely to have blue eyes

Did You Know? Embark researchers discovered this gene by studying data from dogs like yours. Who knows what we will be able to discover next? Answer the questions on our research surveys to contribute to future discoveries!

Chondrodysplasia (Leg Length) | Gene: Chr. 18 FGF4 Retrogene | Genetic Result: NN

This variant is associated with a type of disproportionate dwarfism known as chondrodysplasia (CDPA). CDPA is a breed-defining characteristic of many breeds exhibiting a "short-legged, long-bodied" appearance, such as Corgis, Dachshunds, Basset Hounds, and others. Dogs with the II result display the largest reduction in leg length. Dogs with the NI genotype will have an intermediate leg length, while dogs with the NN result will not exhibit leg shortening due to this variant.

Likely to have normal leg length

Did You Know? A similar genetic variant called the chondrodystrophy (CDDY) variant also plays an important role in shortening the leg length of many breeds. Dog breeds with the shortest legs, like the Corgi, Dachshund, and Basset Hound generally have one or two copies of the CDDY and CDPA variants. CDDY (but not CDPA) is also associated with an increased risk of Type I Intervertebral Disc Disease (IVDD). You can see the CDDY result in the health test results under "Intervertebral Disc Disease Type I".

Registration: Microchip: 250268780939543



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### **TRAITS: BODY SIZE**

| TRAIT   | RESULT |
|---|--------|
| Body Size 1   Gene: IGF1   Genetic Result: NN  This is one of several genes that influence the size of a dog. A result of II for this gene is associated with smaller body size. A result of NN is associated with larger body size.        | Larger |
| Body Size 2   Gene: IGFR1   Genetic Result: GG  This is one of several genes that influence the size of a dog. A result of AA for this gene is associated with smaller body size. A result of GG is associated with larger body size.       | Larger |
| Body Size 3   Gene: STC2   Genetic Result: TT  This is one of several genes that influence the size of a dog. A result of AA for this gene is associated with smaller body size. A result of TT is associated with larger body size.        | Larger |
| Body Size 4   Gene: GHR - E191K   Genetic Result: GG  This is one of several genes that influence the size of a dog. A result of AA for this gene is associated with smaller body size. A result of GG is associated with larger body size. | Larger |
| Body Size 5   Gene: GHR - P177L   Genetic Result: CC  This is one of several genes that influence the size of a dog. A result of TT for this gene is associated with smaller body size. A result of CC is associated with larger body size. | Larger |



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#### TRAITS: PERFORMANCE

TRAIT RESULT

Altitude Adaptation | Gene: EPAS1 | Genetic Result: GG

This gene causes dogs to be especially tolerant of low oxygen environments, such as those found at high elevations. Dogs with a AA or GA result will be less susceptible to "altitude sickness."

Normal altitude tolerance

**Did You Know?** This gene was originally identified in breeds from high altitude areas such as the Tibetan Mastiff.

Appetite LINKAGE | Gene: POMC | Genetic Result: NN

This gene influences eating behavior. An **ND** or **DD** result would predict higher food motivation compared to **NN** result, increasing the likelihood to eat excessively, have higher body fat percentage, and be more prone to obesity. Read more about the genetics of POMC, and learn how you can contribute to research, in our blog post (https://embarkvet.com/resources/blog/pomc-dogs/). We measure this result using a linkage test.

Normal food motivation

**Did You Know?** POMC is actually short for "proopiomelanocortin," and is a large protein that is broken up into several smaller proteins that have biological activity. The smaller proteins generated from POMC control, among other things, distribution of pigment to the hair and skin cells, appetite, and energy expenditure.



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#### **HEALTH REPORT**

#### How to interpret U and me Urio Les Griffes des Terres Sauvages's genetic health results:

If U and me Urio Les Griffes des Terres Sauvages inherited any of the variants that we tested, they will be listed at the top of the Health Report section, along with a description of how to interpret this result. We also include all of the variants that we tested U and me Urio Les Griffes des Terres Sauvages for that we did not detect the risk variant for.

#### A genetic test is not a diagnosis

This genetic test does not diagnose a disease. Please talk to your vet about your dog's genetic results, or if you think that your pet may have a health condition or disease.

#### Summary

U and me Urio Les Griffes des Terres Sauvages is not at increased risk for the genetic health conditions that Embark tests.

**⊘** Clear results

Breed-relevant (12)

Other (243)

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### **BREED-RELEVANT RESULTS**

Research studies indicate that these results are more relevant to dogs like U and me Urio Les Griffes des Terres Sauvages, and may influence his chances of developing certain health conditions.

| Anhidrotic Ectodermal Dysplasia (EDA Intron 8)   | Clear |
|--|-------|
| Canine Leukocyte Adhesion Deficiency Type III, CLAD III (FERMT3, German Shepherd Variant)    | Clear |
| Oay Blindness (CNGA3 Exon 7, German Shepherd Variant)  | Clear |
| Obegenerative Myelopathy, DM (SOD1A)   | Clear |
|  | Clear |
| Hemophilia A (F8 Exon 1, German Shepherd Variant 2)  | Clear |
| O Ichthyosis (ASPRV1 Exon 2, German Shepherd Variant)  | Clear |
| Mucopolysaccharidosis Type VII, Sly Syndrome, MPS VII (GUSB Exon 3, German Shepherd Variant) | Clear |
| Multiple Drug Sensitivity (ABCB1)  | Clear |
| Platelet Factor X Receptor Deficiency, Scott Syndrome (TMEM16F)                              | Clear |
| Renal Cystadenocarcinoma and Nodular Dermatofibrosis (FLCN Exon 7)                           | Clear |
|  | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### **OTHER RESULTS**

Research has not yet linked these conditions to dogs with similar breeds to U and me Urio Les Griffes des Terres Sauvages. Review any increased risk or notable results to understand his potential risk and recommendations.

| ② 2-DHA Kidney & Bladder Stones (APRT)  | Clear |
|---|-------|
| Acral Mutilation Syndrome (GDNF-AS, Spaniel and Pointer Variant)  | Clear |
| Alaskan Husky Encephalopathy (SLC19A3)  | Clear |
| Alaskan Malamute Polyneuropathy, AMPN (NDRG1 SNP)   | Clear |
|   | Clear |
|   | Clear |
| <ul> <li>Autosomal Dominant Progressive Retinal Atrophy (RHO)</li> </ul>  | Clear |
| Bald Thigh Syndrome (IGFBP5)  | Clear |
| Bernard-Soulier Syndrome, BSS (GP9, Cocker Spaniel Variant)   | Clear |
| Bully Whippet Syndrome (MSTN)   | Clear |
| Canine Elliptocytosis (SPTB Exon 30)  | Clear |
| Canine Fucosidosis (FUCA1)  | Clear |
| Canine Leukocyte Adhesion Deficiency Type I, CLAD I (ITGB2, Setter Variant)   | Clear |
| Oanine Multifocal Retinopathy, cmr1 (BEST1 Exon 2)  | Clear |
| Oanine Multifocal Retinopathy, cmr2 (BEST1 Exon 5, Coton de Tulear Variant)   | Clear |
| Canine Multifocal Retinopathy, cmr3 (BEST1 Exon 10 Deletion, Finnish and Swedish Lapphund,<br>Lapponian Herder Variant) | Clear |
| Oanine Multiple System Degeneration (SERAC1 Exon 4, Chinese Crested Variant)  | Clear |
| Canine Multiple System Degeneration (SERAC1 Exon 15, Kerry Blue Terrier Variant)  | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| ○ Cardiomyopathy and Juvenile Mortality (YARS2)                                      | Clear |
|--|-------|
| Centronuclear Myopathy, CNM (PTPLA)  | Clear |
| Cerebellar Hypoplasia (VLDLR, Eurasier Variant)                                      | Clear |
| Chondrodystrophy (ITGA10, Norwegian Elkhound and Karelian Bear Dog Variant)          | Clear |
| Cleft Lip and/or Cleft Palate (ADAMTS20, Nova Scotia Duck Tolling Retriever Variant) | Clear |
| Cleft Palate, CP1 (DLX6 intron 2, Nova Scotia Duck Tolling Retriever Variant)        | Clear |
| Cobalamin Malabsorption (CUBN Exon 8, Beagle Variant)                                | Clear |
| Cobalamin Malabsorption (CUBN Exon 53, Border Collie Variant)                        | Clear |
| ○ Collie Eye Anomaly (NHEJ1)   | Clear |
| Omplement 3 Deficiency, C3 Deficiency (C3)   | Clear |
| Congenital Cornification Disorder (NSDHL, Chihuahua Variant)                         | Clear |
| Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant)                  | Clear |
| Congenital Hypothyroidism (TPO, Tenterfield Terrier Variant)                         | Clear |
| Congenital Hypothyroidism with Goiter (TPO Intron 13, French Bulldog Variant)        | Clear |
| Congenital Hypothyroidism with Goiter (SLC5A5, Shih Tzu Variant)                     | Clear |
| Congenital Macrothrombocytopenia (TUBB1 Exon 1, Cairn and Norfolk Terrier Variant)   | Clear |
| Congenital Myasthenic Syndrome, CMS (COLQ, Labrador Retriever Variant)               | Clear |
| Congenital Myasthenic Syndrome, CMS (COLQ, Golden Retriever Variant)                 | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Ongenital Myasthenic Syndrome, CMS (CHAT, Old Danish Pointing Dog Variant)                     | Clear |
|--|-------|
| Ongenital Myasthenic Syndrome, CMS (CHRNE, Jack Russell Terrier Variant)                       | Clear |
| Congenital Stationary Night Blindness (LRIT3, Beagle Variant)                                  | Clear |
| Congenital Stationary Night Blindness (RPE65, Briard Variant)                                  | Clear |
| Craniomandibular Osteopathy, CMO (SLC37A2)   | Clear |
| Craniomandibular Osteopathy, CMO (SLC37A2 Intron 16, Basset Hound Variant)                     | Clear |
| Cystinuria Type I-A (SLC3A1, Newfoundland Variant)   | Clear |
| Oystinuria Type II-A (SLC3A1, Australian Cattle Dog Variant)                                   | Clear |
| Cystinuria Type II-B (SLC7A9, Miniature Pinscher Variant)                                      | Clear |
| Oay Blindness (CNGB3 Deletion, Alaskan Malamute Variant)                                       | Clear |
| Oay Blindness (CNGA3 Exon 7, Labrador Retriever Variant)                                       | Clear |
| Oay Blindness (CNGB3 Exon 6, German Shorthaired Pointer Variant)                               | Clear |
| Opening Syndrome of Dobermans, DVDob, DINGS (MYO7A)  | Clear |
| Demyelinating Polyneuropathy (SBF2/MTRM13)   | Clear |
| Oental-Skeletal-Retinal Anomaly (MIA3, Cane Corso Variant)                                     | Clear |
| Oiffuse Cystic Renal Dysplasia and Hepatic Fibrosis (INPP5E Intron 9, Norwich Terrier Variant) | Clear |
| Oilated Cardiomyopathy, DCM (RBM20, Schnauzer Variant)   | Clear |
| Oilated Cardiomyopathy, DCM1 (PDK4, Doberman Pinscher Variant 1)                               | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| O Dilated Cardiomyopathy, DCM2 (TTN, Doberman Pinscher Variant 2)               | Clear |
|---|-------|
| Oisproportionate Dwarfism (PRKG2, Dogo Argentino Variant)                       | Clear |
| Ory Eye Curly Coat Syndrome (FAM83H Exon 5)                                     | Clear |
| Oystrophic Epidermolysis Bullosa (COL7A1, Central Asian Shepherd Dog Variant)   | Clear |
| Oystrophic Epidermolysis Bullosa (COL7A1, Golden Retriever Variant)             | Clear |
| Early Bilateral Deafness (LOXHD1 Exon 38, Rottweiler Variant)                   | Clear |
| Early Onset Adult Deafness, EOAD (EPS8L2 Deletion, Rhodesian Ridgeback Variant) | Clear |
| Early Onset Cerebellar Ataxia (SEL1L, Finnish Hound Variant)                    | Clear |
| Ehlers Danlos (ADAMTS2, Doberman Pinscher Variant)                              | Clear |
| Enamel Hypoplasia (ENAM Deletion, Italian Greyhound Variant)                    | Clear |
| Enamel Hypoplasia (ENAM SNP, Parson Russell Terrier Variant)                    | Clear |
| Episodic Falling Syndrome (BCAN)  | Clear |
| Exercise-Induced Collapse, EIC (DNM1)   | Clear |
| Factor VII Deficiency (F7 Exon 5)   | Clear |
| Factor XI Deficiency (F11 Exon 7, Kerry Blue Terrier Variant)                   | Clear |
| Familial Nephropathy (COL4A4 Exon 3, Cocker Spaniel Variant)                    | Clear |
| Familial Nephropathy (COL4A4 Exon 30, English Springer Spaniel Variant)         | Clear |
| Fanconi Syndrome (FAN1, Basenji Variant)  | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Fetal-Onset Neonatal Neuroaxonal Dystrophy (MFN2, Giant Schnauzer Variant)  | Clear |
|---|-------|
| Glanzmann's Thrombasthenia Type I (ITGA2B Exon 13, Great Pyrenees Variant)  | Clear |
| Glanzmann's Thrombasthenia Type I (ITGA2B Exon 12, Otterhound Variant)  | Clear |
| Globoid Cell Leukodystrophy, Krabbe disease (GALC Exon 5, Terrier Variant)  | Clear |
| Glycogen Storage Disease Type IA, Von Gierke Disease, GSD IA (G6PC, Maltese Variant)  | Clear |
| Glycogen Storage Disease Type IIIA, GSD IIIA (AGL, Curly Coated Retriever Variant)  | Clear |
| Glycogen storage disease Type VII, Phosphofructokinase Deficiency, PFK Deficiency (PFKM, Whippet<br>and English Springer Spaniel Variant) | Clear |
| Glycogen storage disease Type VII, Phosphofructokinase Deficiency, PFK Deficiency (PFKM, Wachtelhund Variant)                             | Clear |
|   | Clear |
| Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2 (TTC8)  | Clear |
| Goniodysgenesis and Glaucoma, Pectinate Ligament Dysplasia, PLD (OLFM3)   | Clear |
| Hemophilia A (F8 Exon 10, Boxer Variant)  | Clear |
|   | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Hemophilia B (F9 Exon 7, Rhodesian Ridgeback Variant)  | Clear |
|--|-------|
| Hereditary Ataxia, Cerebellar Degeneration (RAB24, Old English Sheepdog and Gordon Setter Variant)   | Clear |
| Hereditary Cataracts (HSF4 Exon 9, Australian Shepherd Variant)  | Clear |
| Hereditary Footpad Hyperkeratosis (FAM83G, Terrier and Kromfohrlander Variant)   | Clear |
| Hereditary Footpad Hyperkeratosis (DSG1, Rottweiler Variant)   | Clear |
| Hereditary Nasal Parakeratosis (SUV39H2 Intron 4, Greyhound Variant)   | Clear |
|  | Clear |
| Hereditary Vitamin D-Resistant Rickets (VDR)   | Clear |
| Hypocatalasia, Acatalasemia (CAT)  | Clear |
| Hypomyelination and Tremors (FNIP2, Weimaraner Variant)  | Clear |
| Hypophosphatasia (ALPL Exon 9, Karelian Bear Dog Variant)  | Clear |
| O Ichthyosis (NIPAL4, American Bulldog Variant)  | Clear |
| O Ichthyosis (SLC27A4, Great Dane Variant)   | Clear |
| Orange Content of the | Clear |
| O Ichthyosis, ICH1 (PNPLA1, Golden Retriever Variant)  | Clear |
| ✓ Inflammatory Myopathy (SLC25A12)   | Clear |
|  | Clear |
| Inherited Selected Cobalamin Malabsorption with Proteinuria (CUBN, Komondor Variant)   | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Intervertebral Disc Disease (Type I) (FGF4 retrogene - CFA12)                              | Clear |
|--|-------|
| Intestinal Lipid Malabsorption (ACSL5, Australian Kelpie)                                  | Clear |
| Junctional Epidermolysis Bullosa (LAMA3 Exon 66, Australian Cattle Dog Variant)            | Clear |
| Junctional Epidermolysis Bullosa (LAMB3 Exon 11, Australian Shepherd Variant)              | Clear |
| Juvenile Epilepsy (LGI2)   | Clear |
| Juvenile Laryngeal Paralysis and Polyneuropathy (RAB3GAP1, Rottweiler Variant)             | Clear |
| Juvenile Myoclonic Epilepsy (DIRAS1)   | Clear |
|  | Clear |
| Lagotto Storage Disease (ATG4D)  | Clear |
| Laryngeal Paralysis (RAPGEF6, Miniature Bull Terrier Variant)                              | Clear |
| Late Onset Spinocerebellar Ataxia (CAPN1)  | Clear |
| Late-Onset Neuronal Ceroid Lipofuscinosis, NCL 12 (ATP13A2, Australian Cattle Dog Variant) | Clear |
| Leonberger Polyneuropathy 1 (LPN1, ARHGEF10)   | Clear |
| O Leonberger Polyneuropathy 2 (GJA9)   | Clear |
| Lethal Acrodermatitis, LAD (MKLN1)   | Clear |
| Leukodystrophy (TSEN54 Exon 5, Standard Schnauzer Variant)                                 | Clear |
|  | Clear |
| Limb Girdle Muscular Dystrophy (SGCD, Boston Terrier Variant)                              | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| <ul> <li>Limb-Girdle Muscular Dystrophy 2D (SGCA Exon 3, Miniature Dachshund Variant)</li> </ul>                     | Clear |
|--|-------|
| O Long QT Syndrome (KCNQ1)   | Clear |
| Lundehund Syndrome (LEPREL1)   | Clear |
| Macular Corneal Dystrophy, MCD (CHST6)   | Clear |
| Malignant Hyperthermia (RYR1)  | Clear |
| May-Hegglin Anomaly (MYH9)   | Clear |
| Methemoglobinemia (CYB5R3, Pit Bull Terrier Variant)   | Clear |
| Methemoglobinemia (CYB5R3)   | Clear |
| Microphthalmia (RBP4 Exon 2, Soft Coated Wheaten Terrier Variant)  | Clear |
| Mucopolysaccharidosis IIIB, Sanfilippo Syndrome Type B, MPS IIIB (NAGLU, Schipperke Variant)                         | Clear |
| Mucopolysaccharidosis Type IIIA, Sanfilippo Syndrome Type A, MPS IIIA (SGSH Exon 6, Dachshund Variant)               | Clear |
| Mucopolysaccharidosis Type IIIA, Sanfilippo Syndrome Type A, MPS IIIA (SGSH Exon 6, New Zealand<br>Huntaway Variant) | Clear |
| Mucopolysaccharidosis Type VI, Maroteaux-Lamy Syndrome, MPS VI (ARSB Exon 5, Miniature Pinscher Variant)             | Clear |
| Mucopolysaccharidosis Type VII, Sly Syndrome, MPS VII (GUSB Exon 5, Terrier Brasileiro Variant)                      | Clear |
| Muscular Dystrophy (DMD, Cavalier King Charles Spaniel Variant 1)  | Clear |
| Muscular Dystrophy (DMD, Golden Retriever Variant)   | Clear |
| Musladin-Lueke Syndrome, MLS (ADAMTSL2)  | Clear |
| Myasthenia Gravis-Like Syndrome (CHRNE, Heideterrier Variant)  | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Myotonia Congenita (CLCN1 Exon 23, Australian Cattle Dog Variant)                        | Clear |
|--|-------|
| Myotonia Congenita (CLCN1 Exon 7, Miniature Schnauzer Variant)                           | Clear |
| Narcolepsy (HCRTR2 Exon 1, Dachshund Variant)  | Clear |
| Narcolepsy (HCRTR2 Intron 4, Doberman Pinscher Variant)                                  | Clear |
| Narcolepsy (HCRTR2 Intron 6, Labrador Retriever Variant)                                 | Clear |
| Nemaline Myopathy (NEB, American Bulldog Variant)  | Clear |
| Neonatal Cerebellar Cortical Degeneration (SPTBN2, Beagle Variant)                       | Clear |
| Neonatal Encephalopathy with Seizures, NEWS (ATF2)                                       | Clear |
| Neonatal Interstitial Lung Disease (LAMP3)   | Clear |
| Neuroaxonal Dystrophy, NAD (VPS11, Rottweiler Variant)                                   | Clear |
| Neuroaxonal Dystrophy, NAD (TECPR2, Spanish Water Dog Variant)                           | Clear |
| Neuronal Ceroid Lipofuscinosis 1, NCL 1 (PPT1 Exon 8, Dachshund Variant 1)               | Clear |
| Neuronal Ceroid Lipofuscinosis 10, NCL 10 (CTSD Exon 5, American Bulldog Variant)        | Clear |
| Neuronal Ceroid Lipofuscinosis 2, NCL 2 (TPP1 Exon 4, Dachshund Variant 2)               | Clear |
| Neuronal Ceroid Lipofuscinosis 5, NCL 5 (CLN5 Exon 4 SNP, Border Collie Variant)         | Clear |
| Neuronal Ceroid Lipofuscinosis 5, NCL 5 (CLN5 Exon 4 Deletion, Golden Retriever Variant) | Clear |
| Neuronal Ceroid Lipofuscinosis 6, NCL 6 (CLN6 Exon 7, Australian Shepherd Variant)       | Clear |
| Neuronal Ceroid Lipofuscinosis 7, NCL7 (MFSD8, Chihuahua and Chinese Crested Variant)    | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8, Australian Shepherd Variant)                                    | Clear |
|--|-------|
| Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8 Exon 2, English Setter Variant)                                  | Clear |
| Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8 Insertion, Saluki Variant)                                       | Clear |
| Neuronal Ceroid Lipofuscinosis, Cerebellar Ataxia, NCL4A (ARSG Exon 2, American Staffordshire Terrier Variant) | Clear |
| Oculocutaneous Albinism, OCA (SLC45A2 Exon 6, Bullmastiff Variant)   | Clear |
| Oculocutaneous Albinism, OCA (SLC45A2, Small Breed Variant)  | Clear |
| Oculoskeletal Dysplasia 2 (COL9A2, Samoyed Variant)  | Clear |
| Osteochondrodysplasia (SLC13A1, Poodle Variant)  | Clear |
| Osteogenesis Imperfecta (COL1A2, Beagle Variant)   | Clear |
| Osteogenesis Imperfecta (SERPINH1, Dachshund Variant)  | Clear |
| Osteogenesis Imperfecta (COL1A1, Golden Retriever Variant)   | Clear |
| P2Y12 Receptor Platelet Disorder (P2Y12)   | Clear |
| Pachyonychia Congenita (KRT16, Dogue de Bordeaux Variant)  | Clear |
| Paroxysmal Dyskinesia, PxD (PIGN)  | Clear |
| Persistent Mullerian Duct Syndrome, PMDS (AMHR2)   | Clear |
| Pituitary Dwarfism (POU1F1 Intron 4, Karelian Bear Dog Variant)  | Clear |
| Polycystic Kidney Disease, PKD (PKD1)  | Clear |
| Pompe's Disease (GAA, Finnish and Swedish Lapphund, Lapponian Herder Variant)                                  | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

|  | Clear                   |
|--|-------------------------|
| Primary Ciliary Dyskinesia, PCD (NME5, Alaskan Malamute Variant)   | Clear                   |
| Primary Ciliary Dyskinesia, PCD (CCDC39 Exon 3, Old English Sheepdog Variant)  | Clear                   |
| Primary Hyperoxaluria (AGXT)   | Clear                   |
| Primary Lens Luxation (ADAMTS17)   | Clear                   |
| Primary Open Angle Glaucoma (ADAMTS17 Exon 11, Basset Fauve de Bretagne Variant)   | Clear                   |
| Primary Open Angle Glaucoma (ADAMTS10 Exon 17, Beagle Variant)   | Clear                   |
| Primary Open Angle Glaucoma (ADAMTS10 Exon 9, Norwegian Elkhound Variant)  | Clear                   |
| Primary Open Angle Glaucoma and Primary Lens Luxation (ADAMTS17 Exon 2, Chinese Shar-Pei Variant)  | Clear                   |
|  |                         |
| Progressive Retinal Atrophy (SAG)  | Clear                   |
| <ul> <li>Progressive Retinal Atrophy (SAG)</li> <li>Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)</li> </ul>  | Clear<br>Clear          |
|  |                         |
| Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)   | Clear                   |
| <ul> <li>Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)</li> <li>Progressive Retinal Atrophy, Bardet-Biedl Syndrome (BBS2 Exon 11, Shetland Sheepdog Variant)</li> </ul>   | Clear                   |
| <ul> <li>Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)</li> <li>Progressive Retinal Atrophy, Bardet-Biedl Syndrome (BBS2 Exon 11, Shetland Sheepdog Variant)</li> <li>Progressive Retinal Atrophy, CNGA (CNGA1 Exon 9)</li> </ul>   | Clear<br>Clear<br>Clear |
| <ul> <li>Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)</li> <li>Progressive Retinal Atrophy, Bardet-Biedl Syndrome (BBS2 Exon 11, Shetland Sheepdog Variant)</li> <li>Progressive Retinal Atrophy, CNGA (CNGA1 Exon 9)</li> <li>Progressive Retinal Atrophy, crd1 (PDE6B, American Staffordshire Terrier Variant)</li> </ul>  | Clear<br>Clear<br>Clear |
| <ul> <li>✓ Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)</li> <li>✓ Progressive Retinal Atrophy, Bardet-Biedl Syndrome (BBS2 Exon 11, Shetland Sheepdog Variant)</li> <li>✓ Progressive Retinal Atrophy, CNGA (CNGA1 Exon 9)</li> <li>✓ Progressive Retinal Atrophy, crd1 (PDE6B, American Staffordshire Terrier Variant)</li> <li>✓ Progressive Retinal Atrophy, crd4/cord1 (RPGRIP1)</li> </ul> | Clear Clear Clear Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Progressive Retinal Atrophy, rcd1 (PDE6B Exon 21, Irish Setter Variant)                | Clear |
|--|-------|
| Progressive Retinal Atrophy, rcd3 (PDE6A)  | Clear |
| Proportionate Dwarfism (GH1 Exon 5, Chihuahua Variant)                                 | Clear |
| Protein Losing Nephropathy, PLN (NPHS1)  | Clear |
| Pyruvate Dehydrogenase Deficiency (PDP1, Spaniel Variant)                              | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 5, Basenji Variant)                              | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 7, Beagle Variant)                               | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 10, Terrier Variant)                             | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 7, Labrador Retriever Variant)                   | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 7, Pug Variant)                                  | Clear |
| Raine Syndrome (FAM20C)  | Clear |
| Recurrent Inflammatory Pulmonary Disease, RIPD (AKNA, Rough Collie Variant)            | Clear |
| Retina Dysplasia and/or Optic Nerve Hypoplasia (SIX6 Exon 1, Golden Retriever Variant) | Clear |
| Sensory Neuropathy (FAM134B, Border Collie Variant)                                    | Clear |
| Severe Combined Immunodeficiency, SCID (PRKDC, Terrier Variant)                        | Clear |
| Severe Combined Immunodeficiency, SCID (RAG1, Wetterhoun Variant)                      | Clear |
| Shaking Puppy Syndrome (PLP1, English Springer Spaniel Variant)                        | Clear |
| Shar-Pei Autoinflammatory Disease, SPAID, Shar-Pei Fever (MTBP)                        | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

| Skeletal Dysplasia 2, SD2 (COL11A2, Labrador Retriever Variant)                         | Clear |
|---|-------|
| Skin Fragility Syndrome (PKP1, Chesapeake Bay Retriever Variant)                        | Clear |
| Spinocerebellar Ataxia (SCN8A, Alpine Dachsbracke Variant)                              | Clear |
| Spinocerebellar Ataxia with Myokymia and/or Seizures (KCNJ10)                           | Clear |
| Spongy Degeneration with Cerebellar Ataxia 1 (KCNJ10)                                   | Clear |
| Spongy Degeneration with Cerebellar Ataxia 2 (ATP1B2)                                   | Clear |
| Stargardt Disease (ABCA4 Exon 28, Labrador Retriever Variant)                           | Clear |
| Succinic Semialdehyde Dehydrogenase Deficiency (ALDH5A1 Exon 7, Saluki Variant)         | Clear |
| ⊘ Thrombopathia (RASGRP1 Exon 5, American Eskimo Dog Variant)                           | Clear |
| Thrombopathia (RASGRP1 Exon 5, Basset Hound Variant)                                    | Clear |
| Thrombopathia (RASGRP1 Exon 8, Landseer Variant)  | Clear |
| Trapped Neutrophil Syndrome, TNS (VPS13B)   | Clear |
| Ullrich-like Congenital Muscular Dystrophy (COL6A3 Exon 10, Labrador Retriever Variant) | Clear |
| Ullrich-like Congenital Muscular Dystrophy (COL6A1 Exon 3, Landseer Variant)            | Clear |
| Unilateral Deafness and Vestibular Syndrome (PTPRQ Exon 39, Doberman Pinscher)          | Clear |
|   | Clear |
|   | Clear |
| ✓ Von Willebrand Disease Type III, Type III vWD (VWF Exon 4, Terrier Variant)           | Clear |



DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

### **OTHER RESULTS**

| On Willebrand Disease Type III, Type III vWD (VWF Intron 16, Nederlandse Kooikerhondje Variant) | Clear     |
|---|-----------|
| On Willebrand Disease Type III, Type III vWD (VWF Exon 7, Shetland Sheepdog Variant)            | Clear     |
| X-Linked Hereditary Nephropathy, XLHN (COL4A5 Exon 35, Samoyed Variant 2)                       | Clear     |
| X-Linked Myotubular Myopathy (MTM1, Labrador Retriever Variant)                                 | Clear     |
| X-Linked Progressive Retinal Atrophy 1, XL-PRA1 (RPGR)  | Clear     |
| X-linked Severe Combined Immunodeficiency, X-SCID (IL2RG Exon 1, Basset Hound Variant)          | Clear     |
| X-linked Severe Combined Immunodeficiency, X-SCID (IL2RG, Corgi Variant)                        | Clear     |
| Xanthine Urolithiasis (XDH, Mixed Breed Variant)  | Clear     |
| β-Mannosidosis (MANBA Exon 16, Mixed-Breed Variant)   | Clear     |
| Mast Cell Tumor   | No result |

Registration: N/A IHR 2231041 \rightarrow\text{embark} Microchip: 250268780939544



23%

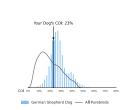
DNA Test Report Test Date: January 17th, 2024 embk.me/uandmeuriolesgriffesdesterressauva

#### INBREEDING AND DIVERSITY

CATEGORY RESULT

Inbreeding | Gene: n/a | Genetic Result: 23%

Inbreeding is a measure of how closely related this dog's parents were. The higher the number, the more closely related the parents. In general, greater inbreeding is associated with increased incidence of genetically inherited conditions.

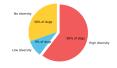


#### Immune Response 1 | Gene: DRB1 | Genetic Result: High Diversity

Diversity in the Major Histocompatibility Complex (MHC) region of the genome has been found in some studies to be associated with the incidence of certain autoimmune diseases. Dogs that have less diversity in the MHC region—i.e. the Dog Leukocyte Antigen (DLA) inherited from the mother is similar to the DLA inherited from the father—are considered less immunologically diverse. A High Diversity result means the dog has two highly dissimilar haplotypes. A Low Diversity result means the dog has two similar but not identical haplotypes. A No Diversity result means the dog has inherited identical haplotypes from both parents. Some studies have shown associations between certain DRB1 haplotypes and autoimmune diseases such as Cushing's disease, but these findings have yet to be scientifically validated.

#### **High Diversity**

How common is this amount of diversity in purebreds:

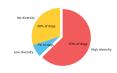


#### Immune Response 2 | Gene: DQA1 and DQB1 | Genetic Result: High Diversity

Diversity in the Major Histocompatibility Complex (MHC) region of the genome has been found in some studies to be associated with the incidence of certain autoimmune diseases. Dogs that have less diversity in the MHC region—i.e. the Dog Leukocyte Antigen (DLA) inherited from the mother is similar to the DLA inherited from the father—are considered less immunologically diverse. A High Diversity result means the dog has two highly dissimilar haplotypes. A Low Diversity result means the dog has two similar but not identical haplotypes. A No Diversity result means the dog has inherited identical haplotypes from both parents. A number of studies have shown correlations of DQA-DQB1 haplotypes and certain autoimmune diseases; however, these have not yet been scientifically validated.

#### **High Diversity**

How common is this amount of diversity in purebreds:



Registration: N/A IHR 2231041 \rightarrow\text{embark} Microchip: 250268780939543