AMERICAN KENNEL CLUB

GOLDEN STAR SUPREME SANDY

NUMBER SS39021408

SEX FEMALE DATE OF BIRTH DECEMBER 20, 2022



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CERTIFICATE ISSUED AUGUST 11, 2023 This certificate invalidates all previous certificates issued. If a date appears after the name and number of the sire and dam, it indicates the issue of the Stud Book Register in which the sire or dam is published.

For Transfer Instructions, see back of Certificate.

This Certificate issued with the right to correct or revoke by the American Kennel Club.

NAME GOLDEN STAR SUPREME SAND BREED GOLDEN RETRIEVER COLOR GOLDEN SIRE ALEX SUPREME SS08532402 04-20 (AKC DNA #V DAM ANNABELLE JOY II SS07105911 04-23 BREEDER ELI A YODER OWNER OWEN YODER 2349 OLD BEN BOW RD UNION GROVE NC 28689-9072

REGISTRATION CERTIFICATE

SS08532402 04-20 (AKC DNA #V927163)

	AMERICAN Gine		MERICAN	SS07105911 (04-23) DK GLDN				GOLDEN RETRIEVER FEMALE GLDN Date Whelped: 12/20/2022 Breeder: ELI A YODER	SS39021408			Sire SS08532402 (04-20) DK GLDN AKC DNA #V927163			ł	Certified Di	L CLUB
	secretary and	CRYSTAL OF HAZEL BY POPE SR92454604 (11-18) LT GLDN				SS00279503 (11-18) LT GLDN AKC DNA #V911188	HOGAN BY ULRICH'S DAKOTA OF			SS03400809 (02-19) GLDN				#V825885	MARTINS CREEK BRUTUS	Edigree	FOUNDED 1884
at this pedigree was compiled from official St	SR86689704 (07-16) LT GLDN	- HAZEL BY AXLE OF DUMBLEDORE LL	SR87271207 (07-16) LT GLDN	POPE BY DEACON OF GOLDEN DUCK	SR90930806 (10-17) LI GLDN	YOLIS ENGLISH CUPCAKE	SR90262503 (04-17) LT GLDN	ULRICH'S DAKOTA	SR96438303 (04-18) GLDN	- STARS-SWEET-SALLY	#V834783	SHETLERS BUTCH SR79234903 (02-16) LT GLDN AKC DNA	SR87188603 (04-16) LT GLDN		SR77367808 (09-14) GLDN AKC DNA #V744607	VALLEY OF THE FRANK	
ud Book records on March 4, 2024.	RONDY'S SONDRA SR74030301 (06-15) LT GLDN	RONDY'S AXLE SR80833712 (06-15) LT GLDN AKC DNA #V750511	NESA'S WHITE GOAST SR75067605 (07-15) LT GLDN	DEACON OF GOLDEN DUCK ROMEO SR75768304 (07-15) LT GLDN	ROXY LOU PRINCESA SR76051302 (08-14) LT GLDN	RUS PEKOS MISTER TVISTER SR59175101 (06-10) OFA24F OFEL24 CHIC68962 LT GLDN (RUS) AKC DNA #V586327	ZAMPANZAR PURE WISH CGC SR73347101 (02-14) LT GLDN (SPA) AKC DNA #V709153	NENUORAMOS WINGS OF WIND SR83729701 (01-15) LT GLDN (LIT) AKC DNA #V737733	DAISY STAR III SR75713003 (11-14) LT GLDN	NISLEYS THOMAS SR76478902 (11-14) LT GLDN AKC DNA #V799223	SHETLERS TRIXIE SR73539302 (12-13) GLDN	SHETLERS BO SR73485009 (12-13) LT GLDN AKC DNA #V720614	GOLDEN WINE ROXIE SR51380806 (10-09) LT GLDN	PURRFECT BISHOP SR38122307 (10-08) LT GLDN	SADIE MAHLADY SR55420603 (06-11) LT GLDN	XERXES XEBEC SR73539306 (08-13) GLDN AKC DNA #V785083	

THE AMERICAN KENNEL CLUB

Research Pedigree - 5 Generation Golden Star Supreme Sandy

Name: Golden Star Supreme Sandy AKC #: SS390214/08 Birth Date: 12/20/2022 Colors/Markings: Golden Breeder(s): Eli A Yoder

Breed/Variety: Golden Retriever Sex: Female

Golden Star Supreme Sandy SS390214/08 Golden			Valley Of The Frank SR773678/08 09-14	Xerxes Xebec SR735393/06 08-13 Golden AKC DNA #V785083	Jacobs Golden Ruger Pooh SR5550006/04 02-11 Golden AKC DNA #V720612 Wilsons Daisy South SR551567/08 02-11 Golden
			Golden AKC DNA #V744607	Sadie Mahlady SR554206/03 06-11	Trigger Happy Day SR471603/06 07-09 Golden AKC DNA #V886678
		Martins Creek Brutus SR912797/01 05-17		Light Golden	Tootsie Golden Day SR476378/04 07-09 Light Golden
		Light Golden AKC DNA #V825885		Purrfect Bishop SR381223/07 10-08	Raber's Gingo SR229437/05 04-06 Light Golden
			Sbf Trixie	Light Golden	Rabers Goldie SR190848/02 04-06 Golden
			SB871886/03 04-16 Light Golden	Golden Wine Roxie SR513808/06 10-09	Sir Moose A Lot Miller SR190686/02 08-06 Golden AKC DNA #V439753
	Alex Supreme SS085324/02 04-20			Light Golden	Bennets Ranger Shaxira SR387811/02 11-08 Golden
	Dark Golden AKC DNA #V927163			Shetlers Bo SR734850/09 12-13	Hillsides Sir Milton SR650206/10 03-12 Light Golden AKC DNA #V662146
			Shetlers Butch SR792349/03 02-16 Light Golden AKC DNA #V834783	Light Golden AKC DNA #V720614	Sassarina Lane Wilson SR623250/05 03-12 Light Golden
				Shetlers Trixie SR735393/02 12-13	Jacobs Golden Ruger Pooh SR550006/04 02-11 Golden AKC DNA #V720612
		Cozy Lane's Lexi		Golden	Wilsons Daisy South SR551567/08 02-11 Golden
		SS034008/09 02-19 Golden		Nisleys Thomas SR764789/02 11-14 Light Golden	Windsong's Ringo Star SR447351/04 03-09 Light Golden OFA24G OFEL24 AKC DNA #V598680
			Stars-Sweet-Sally	AKC DNA #V799223	Misty's Star SR615507/08 01-12 Light Golden
			SR964383/03 04-18 Golden	Daisy Star III SR757130/03 11-14	Star Light's Pudgey SR694534/07 02-13 Light Golden OFA55G AKC DNA #V703519
				Light Golden	Lisa's Biscuit SR534894/07 04-11 Golden

				Nenuoramos Wings Of Wind SR837297/01 01-15 (Lithuania)	Kephles Keep In Touch PKR NO5630/05
			<u>Ulrich's Dakota</u> SR902625/03 04-17	Light Golden AKC DNA #V737733	Nenuoramos Tekera Twist LSVK GR0755/06
			SK902625/03/04-17 Light Golden	Zampanzar Pure Wish CGC SR733471/01 02-14 (Spain)	Dewmist Silver Zampanzar LOE 1411717
		<u>Hogan By Ulrich's Dakota Of</u>		Light Golden AKC DNA #V709153	Zampanzar Vanilla Spice LOE 1830731
		Yolis SS002795/03 11-18 Light Golden		Rus Pekos Mister Tvister SR591751/01 06-10 (Russia)	Derby For Ural Evidog RKF 2073618 09-09
		AŘC DNA #V911188	Yolis English Cupcake SR909308/06 10-17 Light Golden	Light Golden OFA24F OFEL24 AKC DNA #V586327	Rus Pekos Doris RKF 1889966
	Annabelle Joy II SS071059/11 04-23 Dark Golden	9/11 04-23		Roxy Lou Princesa SR760513/02 08-14 Light Golden	Skippy Of Sanitacteam Day SR462284/01 04-09 (Yugoslavia) Light Golden AKC DNA #V502271
					Luna Beatrice Of Norwood SR675711/05 04-13 Light Golden
			Pope By Deacon Of Golden Duck Romeo SR872712/07 07-16 Light Golden	<mark>Deacon Of Golden Duck Romeo</mark> SR757693/04 07-15 Light Golden	Charles Henry Of Golden Duck Romeo SR582668/01 06-11 (Yugoslavia) Light Golden OFA25G OFEL27 AKC DNA #V593212
					Whitegold's Milly SR625181/04 03-12 Light Golden OFA39G
				Nesa's White Goast SR750676/05 07-15	Indy's White Ghost SR473164/01 06-09 Light Golden AKC DNA #V581546
				Light Golden	Juzzies White Ghost SR613979/08 06-11 Light Golden
				Rondy's Axle SR808337/12 06-15	Dumbledore II CGC TKN SR771799/01 08-13 (Yugoslavia) Light Golden OFA29F OFEL24 AKC DNA #V696747
			Hazel By Axle Of Dumbledore Ll SR866897/04 07-16 Light Golden	Light Golden AKC DNA #V750511	Clumbret Near To Heart CA THD CGC TKN SR786046/02 04-14 (Yugoslavia) Light Golden OFEL24 AKC DNA #V705649
				<u>Rondy's Sondra</u> SR740303/01 06-15	Lil Bit-O-Golden's Matia SR660049/10 03-12 Light Golden AKC DNA #V659121
	-			Light Golden	Oakbrook's Classy Chloe SR578315/02 11-12 Light Golden

2024 American Kennel Club

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BREED ANCESTRY

Golden Retriever : 73.6%
Firsh Setter : 19.8%
English Cocker Spaniel : 6.6%

GENETIC STATS

Predicted adult weight: 53 lbs

TEST DETAILS

Kit number: EM-19766660 Swab number: 31220412303489

BREED ANCESTRY BY CHROMOSOME

Our advanced test identifies from where Sandy inherited every part of the chromosome pairs in her genome.

			Breed co	lors:			
	Gol	den Retrieve	er Irish Setter	Englis	h Cocker Spanie	el -	
1	_	2		3		4	
5		6		7	-	8	
9		10		11		12	
13		14		15		16	
17		18		19		20	
21		22		23		24	
25		26		27		28	
29		30		31		32	
33		34		35		36	
37	-	38	-				







Fun Fact

A Golden Retriever is also pictured in the Guinness Book of World's Records for "Most tennis balls held in mouth" (with 6). Test Date: September 1st, 2023



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GOLDEN RETRIEVER

The Golden Retriever was developed in the early 19th century as an ideal hunting companion, able to retrieve birds on both land and water in the marshy Scottish countryside. Their friendliness and intelligence makes the both a popular family pet and an excellent working dog, well suited for being a service dog, therapy dog or for search and rescue. The third most popular breed in the US, the American and Canadian Goldens are generally lankier and darker than their British counterparts. Their wavy, feathered topcoat is water resistant, their undercoat helps them with thermoregulation and both coats have a tendency for heavy seasonal shedding. Goldens need lots of exercise (especially when younger), and their love of play and water means their owners usually get a lot of exercise too! In 2013, the 100th anniversary of Britain's Golden Retriever Club, Goldens from around the world came made the pilgrimage to the breed's birthplace in Scotland, where 222 of them posed in a single record-breaking photo. At the same time, the Golden Retriever Lifetime Study was getting started in the United States, recruiting 3,000 Golden Retrievers for a lifetime study aimed at understanding how genetics, lifestyle and environment influences healthy aging and cancer risk in Goldens.





Test Date: September 1st, 2023



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IRISH SETTER

The Irish Setter is a very regal, yet athletic pup that turn everyone's head at the dog park. They originated in Ireland during the 1700's as a hunting companion. The breed later took America by storm in the late 1800's and again during the 1960's and 70's. With their origins rooted in hunting, Irish Setters come packed to the brim with energy. Expect endless amounts of play dates with a tennis ball. This is also a sensitive breed, which can turn into behavioral problems without good training and socialization from an early age. Irish Setters have a gorgeous coat, which requires a lot of work. These dogs need groomed more frequently along with occasional baths. With great beauty comes great responsibility.

Fun Fact

Irish Setters used to have white in their coat, but it was bred out during the 1800's in Ireland.







Test Date: September 1st, 2023

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ENGLISH COCKER SPANIEL

The English Cocker Spaniel is a breed of gun dog. There are "field" or "working" cockers and "show" cockers. An active sporting dog, the English Cocker Spaniel's compact, solid body practically vibrates with energy and enthusiasm, particularly when at work in the field. Although known for its soft, melting spaniel expression, the breed is a tough worker, capable of covering ground effortlessly and penetrating the densest of cover. His coat can be solid-colored (black, liver or shades of red) or particolored, including ticking or roaning. Prone to ear infections. During the summer, the ears should be checked often. Hanging close to the ground as they do, they can become host to ticks or burrs, often the cause of deafness. The Cocker can gain weight easily; do not overfeed.

Fun Fact

The Cocker is part of the royal family. The Duke and Duchess of Cambridge, also known as Prince William and Kate Middleton, adopted a cocker spaniel puppy in 2012. The puppy, named Lupo, is the son of a cocker spaniel owned by the duchess' mother. Lupo is the latest in a long line of dogs in the royal family.

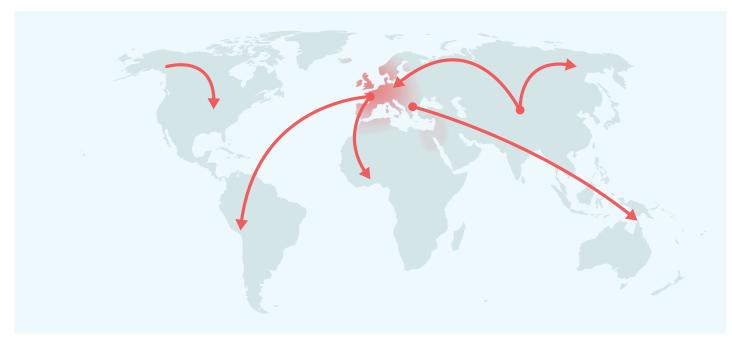




Test Date: September 1st, 2023

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MATERNAL LINE



Through Sandy's mitochondrial DNA we can trace her mother's ancestry back to where dogs and people first became friends. This map helps you visualize the routes that her ancestors took to your home. Their story is described below the map.

HAPLOGROUP: A1e

This female lineage likely stems from some of the original Central Asian wolves that were domesticated into modern dogs starting about 15,000 years ago. It seemed to be a fairly rare dog line for most of dog history until the past 300 years, when the lineage seemed to "explode" out and spread quickly. What really separates this group from the pack is its presence in Alaskan village dogs and Samoyeds. It is possible that this was an indigenous lineage brought to the Americas from Siberia when people were first starting to make that trip themselves! We see this lineage pop up in overwhelming numbers of Irish Wolfhounds, and it also occurs frequently in popular large breeds like Bernese Mountain Dogs, Saint Bernards and Great Danes. Shetland Sheepdogs are also common members of this maternal line, and we see it a lot in Boxers, too. Though it may be all mixed up with European dogs thanks to recent breeding events, its origins in the Americas makes it a very exciting lineage for sure!

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HAPLOTYPE: A2a

Part of the large A1e haplogroup, we see this haplotype in village dogs up and down the Americas as well as French Polynesia. Among the breed dogs we have detected it in, we see it most frequently in English Springer Spaniels, Papillons, and Collies.



Test Date: September 1st, 2023



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RESULT

TRAITS: COAT COLOR

TRAIT

E Locus (MC1R)

The E Locus determines if and where a dog can produce dark (black or brown) hair. Dogs with two copies of the recessive **e** allele do not produce dark hairs at all, and will be "red" over their entire body. The shade of red, which can range from a deep copper to yellow/gold to cream, is dependent on other genetic factors including the Intensity loci. In addition to determining if a dog can develop dark hairs at all, the E Locus 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 the **Em** allele usually have a melanistic mask (dark facial hair as commonly seen in the German Shepherd and Pug). Dogs with no copies of **Em** but one or two copies of the **Eg** allele usually have a melanistic "widow's peak" (dark forehead hair as commonly seen in the Afghan Hound and Borzoi, where it is called either "grizzle" or "domino").

No dark hairs anywhere (ee)

K Locus (CBD103)

The K Locus K^B allele "overrides" the A Locus, meaning that it prevents the A Locus genotype from affecting coat color. For this reason, the K^B allele is referred to as the "dominant black" allele. As a result, dogs with at least one K^B allele will usually have solid black or brown coats (or red/cream coats if they are **ee** at the E Locus) regardless of their genotype at the A Locus, although several other genes could impact the dog's coat and cause other patterns, such as white spotting. Dogs with the **k**^y**k**^y genotype will show a coat color pattern based on the genotype they have at the A Locus. Dogs who test as **K**^B**k**^y may be brindle rather than black or brown.

Not expressed (K^Bk^y)





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RESULT

TRAITS: COAT COLOR (CONTINUED)

TRAIT

Intensity Loci LINKAGE

Areas of a dog's coat where dark (black or brown) pigment is not expressed either contain red/yellow pigment, or no pigment at all. Five locations across five chromosomes explain approximately 70% of red pigmentation "intensity" variation across all dogs. Dogs with a result of **Intense Red Pigmentation** will likely have deep red hair like an Irish Setter or "apricot" hair like some Poodles, dogs with a result of **Intermediate Red Pigmentation** will likely have tan or yellow hair like a Soft-Coated Wheaten Terrier, and dogs with **Dilute Red Pigmentation** will likely have cream or white hair like a Samoyed. Because the mutations we test may not directly cause differences in red pigmentation intensity, we consider this to be a linkage test.

Any pigmented hair likely yellow or tan (Intermediate Red Pigmentation)

A Locus (ASIP)

The A Locus controls switching between black and red pigment in hair cells, but it will only be expressed in dogs that are not **ee** at the E Locus and are **k**^y**k**^y at the K Locus. Sable (also called "Fawn") dogs have a mostly or entirely red coat with some interspersed black hairs. Agouti (also called "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.

D Locus (MLPH)

The D locus result that we report is determined by two different genetic variants that can work together to cause diluted pigmentation. These are the common **d** allele, also known as "**d1**", and a less common allele known as "**d2**". Dogs with two **d** alleles, regardless of which variant, will have all black pigment lightened ("diluted") to gray, or brown pigment lightened to lighter brown in their hair, skin, and sometimes eyes. There are many breed-specific names for these dilute colors, such as "blue", "charcoal", "fawn", "silver", and "Isabella". Note that in certain breeds, dilute dogs have a higher incidence of Color Dilution Alopecia. Dogs with one **d** allele will not be dilute, but can pass the **d** allele on to their puppies. To view your dog's **d1** and **d2** test results, click the "SEE DETAILS" link in the upper right hand corner of the "Base Coat Color" section of the Traits page, and then click the "VIEW SUBLOCUS RESULTS" link at the bottom of the page.

Not expressed (ata)

Not expressed (DD)

Registration:







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RESULT

TRAITS: COAT COLOR (CONTINUED)

TRAIT

Cocoa (HPS3)

Dogs with the coco genotype will produce dark brown pigment instead of black in both their hair and skin.No co alleles, notDogs with the Nco genotype will produce black pigment, but can pass the co allele on to their puppies.expressed (NN)Dogs that have the coco genotype as well as the bb genotype at the B locus are generally a lighter brownthan dogs that have the Bb or BB genotypes at the B locus.

B Locus (TYRP1)

Dogs with two copies of the **b** allele produce brown pigment instead of black in both their hair and skin. Dogs with one copy of the **b** allele will produce black pigment, but can pass the **b** allele on to their puppies. E Locus **ee** dogs that carry two **b** alleles will have red or cream coats, but have brown noses, eye rims, and footpads (sometimes referred to as "Dudley Nose" in Labrador Retrievers). "Liver" or "chocolate" is the preferred color term for brown in most breeds; in the Doberman Pinscher it is referred to as "red".

Likely black colored nose/feet (BB)

Not expressed (NI)

Saddle Tan (RALY)

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 **a**^t allele, so dogs that do not express **a**^t are not influenced by this gene.

S Locus (MITF)

The S Locus determines white spotting and pigment distribution. MITF controls where pigment is produced, and an insertion in the MITF gene causes a loss of pigment in the coat and skin, resulting in white hair and/or pink skin. Dogs with two copies of this variant will likely have breed-dependent white patterning, with a nearly white, parti, or piebald coat. Dogs with one copy of this variant will have more limited white spotting and may be considered flash, parti or piebald. This MITF variant does not explain all white spotting patterns in dogs and other variants are currently being researched. Some dogs may have small amounts of white on the paws, chest, face, or tail regardless of their S Locus genotype.

Likely to have little to no white in coat (SS)

Registration:







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No merle alleles (mm)

RESULT

TRAITS: COAT COLOR (CONTINUED)

TRAIT

M Locus (PMEL)

Merle coat patterning is common to several dog breeds including the Australian Shepherd, Catahoula Leopard Dog, and Shetland Sheepdog, among many others. Merle arises from an unstable SINE insertion (which we term the "M*" allele) that disrupts activity of the pigmentary gene PMEL, leading to mottled or patchy coat color. Dogs with an **M*m** result are likely to be phenotypically 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 be phenotypically merle. Dogs with an **mm** result have no merle alleles and are unlikely to have a merle coat pattern.

Note that Embark does not currently distinguish between the recently described cryptic, atypical, atypical+, classic, and harlequin merle alleles. Our merle test only detects the presence, but not the length of the SINE insertion. We do not recommend making breeding decisions on this result alone. Please pursue further testing for allelic distinction prior to breeding decisions.

R Locus (USH2A) LINKAGE

The R Locus regulates the presence or absence of the roan coat color pattern. Partial duplication of the USH2A gene is strongly associated with this coat pattern. Dogs with at least one **R** allele will likely have roaning on otherwise uniformly unpigmented white areas. Roan appears in white areas controlled by the S Locus but not in other white or cream areas created by other loci, such as the E Locus with **ee** along with Dilute Red Pigmentation by I Locus (for example, in Samoyeds). Mechanisms for controlling the extent of roaning are currently unknown, and roaning can appear in a uniform or non-uniform pattern. Further, non-uniform roaning may appear as ticked, and not obviously roan. The roan pattern can appear with or without ticking.

Likely no impact on coat pattern (rr)

H Locus (Harlequin)

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. This trait is thought to be homozygous lethal; a living dog with an **HH** genotype has never been found.

No harlequin alleles (hh)







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RESULT

TRAITS: OTHER COAT TRAITS

TRAIT

Furnishings (RSP02) LINKAGE

Dogs with one or two copies of the F allele have "furnishings": the mustache, beard, and eyebrows characteristic of breeds like the Schnauzer, Scottish Terrier, and Wire Haired Dachshund. A dog with two I alleles will not have furnishings, which is sometimes called an "improper coat" in breeds where and/or eyebrows) (II) furnishings are part of the breed standard. The mutation is a genetic insertion which we measure indirectly using a linkage test highly correlated with the insertion.

Likely unfurnished (no mustache, beard,

Coat Length (FGF5)

The FGF5 gene is known to affect hair length in many different species, including cats, dogs, mice, and humans. In dogs, the T allele confers a long, silky haircoat as observed in the Yorkshire Terrier and the Long Haired Whippet. The ancestral G allele causes a shorter coat as seen in the Boxer or the American Staffordshire Terrier. In certain breeds (such as Corgi), the long haircoat is described as "fluff."

Likely long coat (TT)

Shedding (MC5R)

Dogs with at least one copy of the ancestral C allele, like many Labradors and German Shepherd Dogs, are heavy or seasonal shedders, while those with two copies of the **T** allele, including many Boxers, Shih Tzus and Chihuahuas, tend to be lighter shedders. Dogs with furnished/wire-haired coats caused by RSPO2 (the furnishings gene) tend to be low shedders regardless of their genotype at this gene.

Hairlessness (FOXI3) LINKAGE

A duplication in the FOXI3 gene causes hairlessness over most of the body as well as changes in tooth shape and number. This mutation occurs in Peruvian Inca Orchid, Xoloitzcuintli (Mexican Hairless), and Chinese Crested (other hairless breeds have different mutations). Dogs with the NDup genotype are likely to be hairless while dogs with the NN genotype are likely to have a normal coat. The DupDup genotype has never been observed, suggesting that dogs with that genotype cannot survive to birth. Please note that this is a linkage test, so it may not be as predictive as direct tests of the mutation in some lines.

Likely heavy/seasonal shedding (CT)

Very unlikely to be hairless (NN)

Hairlessness (SGK3)

Hairlessness in the American Hairless Terrier arises from a mutation in the SGK3 gene. 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

Very unlikely to be hairless (NN)

Registration:







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RESULT

TRAITS: OTHER COAT TRAITS (CONTINUED)

TRAIT

Oculocutaneous Albinism Type 2 (SLC45A2) LINKAGE

Dogs with two copies **DD** of this deletion in the SLC45A2 gene have oculocutaneous albinism (OCA), also known as Doberman Z Factor Albinism, a recessive condition characterized by severely reduced or absent pigment in the eyes, skin, and hair. Affected dogs sometimes suffer from vision problems due to lack of eye pigment (which helps direct and absorb ambient light) and are prone to sunburn. Dogs with a single copy of the deletion **ND** will not be affected but can pass the mutation on to their offspring. 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. Please note that this is a linkage test, so it may not be as predictive as direct tests of the mutation in some lines.

Coat Texture (KRT71)

Dogs with a long coat and at least one copy of the **T** allele have a wavy or curly coat characteristic of Poodles and Bichon Frises. Dogs with two copies of the ancestral **C** allele are likely to have a straight coat, but there are other factors that can cause a curly coat, for example if they at least one **F** allele for the Furnishings (RSPO2) gene then they are likely to have a curly coat. Dogs with short coats may carry one or two copies of the **T** allele but still have straight coats.

Likely straight coat (CC)





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RESULT

TRAITS: OTHER BODY FEATURES

TRAIT

Muzzle Length (BMP3)

Dogs in medium-length muzzle (mesocephalic) breeds like Staffordshire Terriers and Labradors, and long muzzle (dolichocephalic) breeds like Whippet and Collie have one, or more commonly two, copies of the ancestral **C** allele. Dogs in many short-length muzzle (brachycephalic) breeds such as the English Bulldog, Pug, and Pekingese have two copies of the derived **A** allele. At least five different genes affect muzzle length in dogs, with BMP3 being the only one with a known causal mutation. For example, the skull shape of some breeds, including the dolichocephalic Scottish Terrier or the brachycephalic Japanese Chin, appear to be caused by other genes. Thus, dogs may have short or long muzzles due to other genetic factors that are not yet known to science.

Likely medium or long muzzle (CC)

Tail Length (T)

Whereas most dogs have two **C** alleles and a long tail, dogs with one **G** allele are likely to have a bobtail, which is an unusually short or absent tail. This mutation causes natural bobtail in many 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 the **GG** genotype do not survive to birth. Please note that this mutation does not explain every natural bobtail! 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, these breeds do not have this mutation. This suggests that other unknown genetic mutations can also lead to a natural bobtail.

Hind Dewclaws (LMBR1)

Common in certain breeds such as the Saint Bernard, hind dewclaws are extra, nonfunctional digits located midway between a dog's paw and hock. Dogs with at least one copy of the **T** allele have about a 50% chance of having hind dewclaws. Note that other (currently unknown to science) mutations can also cause hind dewclaws, so some **CC** or **TC** dogs will have hind dewclaws.

Likely normal-length tail (CC)

Unlikely to have hind dew claws (CC)

Registration:



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TRAITS: OTHER BODY FEATURES (CONTINUED)

Blue Eye Color (ALX4) LINKAGE

Back Muscling & Bulk, Large Breed (ACSL4)

Embark researchers discovered this large duplication associated with blue eyes in Arctic breeds like Siberian Husky as well as tri-colored (non-merle) Australian Shepherds. Dogs with at least one copy of the duplication (Dup) are more likely to have at least one blue eye. Some dogs with the duplication may have only one blue eye (complete heterochromia) or may not have blue eyes at all; nevertheless, they can still pass the duplication and the trait to their offspring. NN dogs do not carry this duplication, but may have blue eyes due to other factors, such as merle. Please note that this is a linkage test, so it may not be as predictive as direct tests of the mutation in some lines.

The T allele is associated with heavy muscling along the back and trunk in characteristically "bulky" large-

Rottweiler. The "bulky" T allele is absent from leaner shaped large breed dogs like the Great Dane, Irish

Wolfhound, and Scottish Deerhound, which are fixed for the ancestral C allele. Note that this mutation does not seem to affect muscling in small or even mid-sized dog breeds with notable back muscling, including

breed dogs including the Saint Bernard, Bernese Mountain Dog, Greater Swiss Mountain Dog, and

the American Staffordshire Terrier, Boston Terrier, and the English Bulldog.

Less likely to have blue eyes (NN)

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Likely normal muscling (CC)



SANDY

DNA Test Report

TRAIT



RESULT





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TRAITS: BODY SIZE		
TRAIT		RESULT
Body Size (IGF1)		
The I allele is associated with smaller body siz	ze.	Larger (NN)
Body Size (IGFR1)		Larger (GG)
The A allele is associated with smaller body si	ze.	
Body Size (STC2)		Larger (TT)
The A allele is associated with smaller body si	ze.	
Body Size (GHR - E191K)		Intermediate (GA)
The A allele is associated with smaller body si	ze.	interneulate (GA)
Body Size (GHR - P177L)		Larger (CC)
The ${\bf T}$ allele is associated with smaller body si	ze.	





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RESULT

TRAITS: PERFORMANCE

TRAIT

Altitude Adaptation (EPAS1)

This mutation causes dogs to be especially tolerant of low oxygen environments (hypoxia), such as those found at high elevations. Dogs with at least one **A** allele are less susceptible to "altitude sickness." This mutation was originally identified in breeds from high altitude areas such as the Tibetan Mastiff.

Appetite (POMC) LINKAGE

This mutation in the POMC gene is found primarily in Labrador and Flat Coated Retrievers. Compared to
dogs with no copies of the mutation (NN), dogs with one (ND) or two (DD) copies of the mutation are more
likely to have high food motivation, which can cause them to eat excessively, have higher body fat
motiva
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.Norma

Normal food motivation (NN)



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DNA Test Report

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HEALTH REPORT

How to interpret Sandy's genetic health results:

If Sandy 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 Sandy 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

Of the 255 genetic health risks we analyzed, we found 2 results that you should learn about.

Notable results (2)

ALT Activity

Ichthyosis, ICH1

Clear results

Breed-relevant (20)

Other (233)







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BREED-RELEVANT RESULTS

Research studies indicate that these results are more relevant to dogs like Sandy, and may influence her chances of developing certain health conditions.

Ichthyosis, ICH1 (PNPLA1, Golden Retriever Variant)	Notable
Acral Mutilation Syndrome (GDNF-AS, Spaniel and Pointer Variant)	Clear
Bernard-Soulier Syndrome, BSS (GP9, Cocker Spaniel Variant)	Clear
Canine Leukocyte Adhesion Deficiency Type I, CLAD I (ITGB2, Setter Variant)	Clear
Congenital Myasthenic Syndrome, CMS (COLQ, Golden Retriever Variant)	Clear
O Degenerative Myelopathy, DM (SOD1A)	Clear
O Dystrophic Epidermolysis Bullosa (COL7A1, Golden Retriever Variant)	Clear
Exercise-Induced Collapse, EIC (DNM1)	Clear
Samilial Nephropathy (COL4A4 Exon 3, Cocker Spaniel Variant)	Clear
Glycogen storage disease Type VII, Phosphofructokinase Deficiency, PFK Deficiency (PFKM, Whippet and English Springer Spaniel Variant)	Clear
Golden Retriever Progressive Retinal Atrophy 1, GR-PRA1 (SLC4A3)	Clear
Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2 (TTC8)	Clear
Intervertebral Disc Disease (Type I) (FGF4 retrogene - CFA12)	Clear
Muscular Dystrophy (DMD, Golden Retriever Variant)	Clear
Neuronal Ceroid Lipofuscinosis 5, NCL 5 (CLN5 Exon 4 Deletion, Golden Retriever Variant)	Clear
Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8 Exon 2, English Setter Variant)	Clear
Osteogenesis Imperfecta (COL1A1, Golden Retriever Variant)	Clear
Progressive Retinal Atrophy, prcd (PRCD Exon 1)	Clear





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BREED-RELEVANT RE	SULTS		
Progressive Retinal Atrophy,	rcd1 (PDE6B Exon 21, Irish Setter Variant)	Clear	
Retina Dysplasia and/or Optio	c Nerve Hypoplasia (SIX6 Exon 1, Golden Retriever Variant)	Clear	
Von Willebrand Disease Type	I, Type I vWD (VWF)	Clear	

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OTHER RESULTS

Research has not yet linked these conditions to dogs with similar breeds to Sandy. Review any increased risk or notable results to understand her potential risk and recommendations.

O ALT Activity (GPT)	Notable
2-DHA Kidney & Bladder Stones (APRT)	Clear
Alaskan Husky Encephalopathy (SLC19A3)	Clear
Alaskan Malamute Polyneuropathy, AMPN (NDRG1 SNP)	Clear
Alexander Disease (GFAP)	Clear
Anhidrotic Ectodermal Dysplasia (EDA Intron 8)	Clear
Autosomal Dominant Progressive Retinal Atrophy (RHO)	Clear
Bald Thigh Syndrome (IGFBP5)	Clear
Bully Whippet Syndrome (MSTN)	Clear
Canine Elliptocytosis (SPTB Exon 30)	Clear
Canine Fucosidosis (FUCA1)	Clear
Canine Leukocyte Adhesion Deficiency Type III, CLAD III (FERMT3, German Shepherd Variant)	Clear
Canine Multifocal Retinopathy, cmr1 (BEST1 Exon 2)	Clear
Canine Multifocal Retinopathy, cmr2 (BEST1 Exon 5, Coton de Tulear Variant)	Clear
Canine Multifocal Retinopathy, cmr3 (BEST1 Exon 10 Deletion, Finnish and Swedish Lapphund, Lapponian Herder Variant)	Clear
Canine Multiple System Degeneration (SERAC1 Exon 4, Chinese Crested Variant)	Clear
Canine Multiple System Degeneration (SERAC1 Exon 15, Kerry Blue Terrier Variant)	Clear
Cardiomyopathy and Juvenile Mortality (YARS2)	Clear



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OTHER RESULTS

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
Complement 3 Deficiency, C3 Deficiency (C3)	Clear
Congenital Cornification Disorder (NSDHL, Chihuahua Variant)	Clear
 Congenital Cornification Disorder (NSDHL, Chihuahua Variant) Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant) 	Clear Clear
Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant)	Clear
 Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant) Congenital Hypothyroidism (TPO, Tenterfield Terrier Variant) 	Clear Clear
 Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant) Congenital Hypothyroidism (TPO, Tenterfield Terrier Variant) Congenital Hypothyroidism with Goiter (TPO Intron 13, French Bulldog Variant) 	Clear Clear Clear
 Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant) Congenital Hypothyroidism (TPO, Tenterfield Terrier Variant) Congenital Hypothyroidism with Goiter (TPO Intron 13, French Bulldog Variant) Congenital Hypothyroidism with Goiter (SLC5A5, Shih Tzu Variant) 	Clear Clear Clear Clear
 Congenital Hypothyroidism (TPO, Rat, Toy, Hairless Terrier Variant) Congenital Hypothyroidism (TPO, Tenterfield Terrier Variant) Congenital Hypothyroidism with Goiter (TPO Intron 13, French Bulldog Variant) Congenital Hypothyroidism with Goiter (SLC5A5, Shih Tzu Variant) Congenital Macrothrombocytopenia (TUBB1 Exon 1, Cairn and Norfolk Terrier Variant) 	Clear Clear Clear Clear Clear

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OTHER RESULTS		
⊘ Congenital Stationary Night Blindness (LF	lT3, Beagle Variant)	Clear
Ongenital Stationary Night Blindness (RF	PE65, Briard Variant)	Clear
Craniomandibular Osteopathy, CMO (SLC3	37A2)	Clear
Craniomandibular Osteopathy, CMO (SLC3	7A2 Intron 16, Basset Hound Variant)	Clear
🔗 Cystinuria Type I-A (SLC3A1, Newfoundlar	d Variant)	Clear
Cystinuria Type II-A (SLC3A1, Australian C	attle Dog Variant)	Clear
🔗 Cystinuria Type II-B (SLC7A9, Miniature Pi	nscher Variant)	Clear
Day Blindness (CNGB3 Deletion, Alaskan N	Malamute Variant)	Clear
Day Blindness (CNGA3 Exon 7, German Sh	epherd Variant)	Clear
Day Blindness (CNGA3 Exon 7, Labrador Register)	etriever Variant)	Clear
Day Blindness (CNGB3 Exon 6, German Sh	orthaired Pointer Variant)	Clear
Deafness and Vestibular Syndrome of Dob	ermans, DVDob, DINGS (MYO7A)	Clear
Demyelinating Polyneuropathy (SBF2/MT	RM13)	Clear
Oental-Skeletal-Retinal Anomaly (MIA3, C	ane Corso Variant)	Clear
O Diffuse Cystic Renal Dysplasia and Hepati	c Fibrosis (INPP5E Intron 9, Norwich Terrier Variant)	Clear
Dilated Cardiomyopathy, DCM (RBM20, Sc	hnauzer Variant)	Clear
Dilated Cardiomyopathy, DCM1 (PDK4, Dot	berman Pinscher Variant 1)	Clear
Oilated Cardiomyopathy, DCM2 (TTN, Dob	erman Pinscher Variant 2)	Clear

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OTHER RESULTS

O Disproportionate Dwarfism (PRKG2, Dogo Argentino Variant)	Clear
Ory Eye Curly Coat Syndrome (FAM83H Exon 5)	Clear
O Dystrophic Epidermolysis Bullosa (COL7A1, Central Asian Shepherd Dog 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
Factor VII Deficiency (F7 Exon 5)	Clear
Sector XI Deficiency (F11 Exon 7, Kerry Blue Terrier Variant)	Clear
Familial Nephropathy (COL4A4 Exon 30, English Springer Spaniel Variant)	Clear
Fanconi Syndrome (FAN1, Basenji Variant)	Clear
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

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OTHER RESULTS

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, Wachtelhund Variant) 	Clear
GM1 Gangliosidosis (GLB1 Exon 2, Portuguese Water Dog Variant)	Clear
GM1 Gangliosidosis (GLB1 Exon 15, Shiba Inu Variant)	Clear
🧭 GM1 Gangliosidosis (GLB1 Exon 15, Alaskan Husky Variant)	Clear
GM2 Gangliosidosis (HEXA, Japanese Chin Variant)	Clear
GM2 Gangliosidosis (HEXB, Poodle Variant)	Clear
Goniodysgenesis and Glaucoma, Pectinate Ligament Dysplasia, PLD (OLFM3)	Clear
Hemophilia A (F8 Exon 11, German Shepherd Variant 1)	Clear
Hemophilia A (F8 Exon 1, German Shepherd Variant 2)	Clear
Hemophilia A (F8 Exon 10, Boxer Variant)	Clear
Hemophilia B (F9 Exon 7, Terrier Variant)	Clear
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

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OTHER RESULTS		
Hereditary Nasal Parakerato	osis (SUV39H2 Intron 4, Greyhound Variant)	Clear
Hereditary Nasal Parakerato	osis, HNPK (SUV39H2)	Clear
Hereditary Vitamin D-Resist	tant Rickets (VDR)	Clear
🔗 Hypocatalasia, Acatalasemi	a (CAT)	Clear
Hypomyelination and Tremo	ors (FNIP2, Weimaraner Variant)	Clear
🔗 Hypophosphatasia (ALPL E)	kon 9, Karelian Bear Dog Variant)	Clear
🔗 Ichthyosis (NIPAL4, America	an Bulldog Variant)	Clear
Ichthyosis (ASPRV1 Exon 2,	German Shepherd Variant)	Clear
🔗 Ichthyosis (SLC27A4, Great	Dane Variant)	Clear
🔗 Ichthyosis, Epidermolytic H	yperkeratosis (KRT10, Terrier Variant)	Clear
🔗 Inflammatory Myopathy (SL	C25A12)	Clear

Inherited Myopathy of Great Danes (BIN1) Clear \oslash Inherited Selected Cobalamin Malabsorption with Proteinuria (CUBN, Komondor Variant) \oslash Clear Intestinal Lipid Malabsorption (ACSL5, Australian Kelpie) Clear \oslash \oslash Junctional Epidermolysis Bullosa (LAMA3 Exon 66, Australian Cattle Dog Variant) Clear Junctional Epidermolysis Bullosa (LAMB3 Exon 11, Australian Shepherd Variant) Clear \oslash Juvenile Epilepsy (LGI2) Clear \oslash Juvenile Laryngeal Paralysis and Polyneuropathy (RAB3GAP1, Rottweiler Variant) Clear \oslash

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OTHER RESULTS

Juvenile Myoclonic Epilepsy (DIRAS1)	Clear
C L-2-Hydroxyglutaricaciduria, L2HGA (L2HGDH, Staffordshire Bull Terrier Variant)	Clear
Lagotto Storage Disease (ATG4D)	Clear
Laryngeal Paralysis (RAPGEF6, Miniature Bull Terrier Variant)	Clear
Conset Spinocerebellar Ataxia (CAPN1)	Clear
S Late-Onset Neuronal Ceroid Lipofuscinosis, NCL 12 (ATP13A2, Australian Cattle Dog Variant)	Clear
Contemporate Polyneuropathy 1 (LPN1, ARHGEF10)	Clear
Control Leonberger Polyneuropathy 2 (GJA9)	Clear
Control Lethal Acrodermatitis, LAD (MKLN1)	Clear
Leukodystrophy (TSEN54 Exon 5, Standard Schnauzer Variant)	Clear
Control Ligneous Membranitis, LM (PLG)	Clear
C Limb Girdle Muscular Dystrophy (SGCD, Boston Terrier Variant)	Clear
C Limb-Girdle Muscular Dystrophy 2D (SGCA Exon 3, Miniature Dachshund Variant)	Clear
Cong QT Syndrome (KCNQ1)	Clear
Lundehund Syndrome (LEPREL1)	Clear
Macular Corneal Dystrophy, MCD (CHST6)	Clear
Malignant Hyperthermia (RYR1)	Clear
May-Hegglin Anomaly (MYH9)	Clear

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OTHER RESULTS

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 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 3, German Shepherd Variant)	Clear
Mucopolysaccharidosis Type VII, Sly Syndrome, MPS VII (GUSB Exon 5, Terrier Brasileiro Variant)	Clear
Multiple Drug Sensitivity (ABCB1)	Clear
Muscular Dystrophy (DMD, Cavalier King Charles Spaniel Variant 1)	Clear
Musladin-Lueke Syndrome, MLS (ADAMTSL2)	Clear
Myasthenia Gravis-Like Syndrome (CHRNE, Heideterrier Variant)	Clear
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

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Clear

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OTHER RESULTS		
Nemaline Myopathy (NEB, American Bulld	og Variant)	Clear
Neonatal Cerebellar Cortical Degeneratio	n (SPTBN2, Beagle Variant)	Clear
Neonatal Encephalopathy with Seizures,	NEWS (ATF2)	Clear
Neonatal Interstitial Lung Disease (LAMPS)	3)	Clear
Neuroaxonal Dystrophy, NAD (VPS11, Rotte	weiler Variant)	Clear
Neuroaxonal Dystrophy, NAD (TECPR2, Sp	anish Water Dog Variant)	Clear
Neuronal Ceroid Lipofuscinosis 1, NCL 1 (F	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 6, NCL 6 (CLN6 Exon 7, Australian Shepherd Variant)	Clear
Neuronal Ceroid Lipofuscinosis 7, NCL 7 (N	MFSD8, Chihuahua and Chinese Crested Variant)	Clear
Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8, Australian Shepherd Variant)	Clear
Neuronal Ceroid Lipofuscinosis 8, NCL 8 (CLN8 Insertion, Saluki Variant)	Clear
 Neuronal Ceroid Lipofuscinosis, Cerebella Variant) 	ar Ataxia, NCL4A (ARSG Exon 2, American Staffordshire Terr	ier Clear
Oculocutaneous Albinism, OCA (SLC45A2	Exon 6, Bullmastiff Variant)	Clear
Oculocutaneous Albinism, OCA (SLC45A2	, Small Breed Variant)	Clear

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Oculoskeletal Dysplasia 2 (COL9A2, Samoyed Variant)



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OTHER RESULTS

Osteochondrodysplasia (SLC13A1, Poodle Variant)	Clear
Osteogenesis Imperfecta (COL1A2, Beagle Variant)	Clear
Osteogenesis Imperfecta (SERPINH1, Dachshund 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
Platelet Factor X Receptor Deficiency, Scott Syndrome (TMEM16F)	Clear
Polycystic Kidney Disease, PKD (PKD1)	Clear
Pompe's Disease (GAA, Finnish and Swedish Lapphund, Lapponian Herder Variant)	Clear
 Pompe's Disease (GAA, Finnish and Swedish Lapphund, Lapponian Herder Variant) Prekallikrein Deficiency (KLKB1 Exon 8) 	
	Clear
Prekallikrein Deficiency (KLKB1 Exon 8)	Clear Clear
 Prekallikrein Deficiency (KLKB1 Exon 8) Primary Ciliary Dyskinesia, PCD (NME5, Alaskan Malamute Variant) 	Clear Clear Clear
 Prekallikrein Deficiency (KLKB1 Exon 8) Primary Ciliary Dyskinesia, PCD (NME5, Alaskan Malamute Variant) Primary Ciliary Dyskinesia, PCD (CCDC39 Exon 3, Old English Sheepdog Variant) 	Clear Clear Clear Clear
 Prekallikrein Deficiency (KLKB1 Exon 8) Primary Ciliary Dyskinesia, PCD (NME5, Alaskan Malamute Variant) Primary Ciliary Dyskinesia, PCD (CCDC39 Exon 3, Old English Sheepdog Variant) Primary Hyperoxaluria (AGXT) 	Clear Clear Clear Clear Clear

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OTHER RESULTS

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
Progressive Retinal Atrophy (IFT122 Exon 26, Lapponian Herder Variant)	Clear
Progressive Retinal Atrophy, Bardet-Biedl Syndrome (BBS2 Exon 11, Shetland Sheepdog Variant)	Clear
Progressive Retinal Atrophy, CNGA (CNGA1 Exon 9)	Clear
Progressive Retinal Atrophy, crd1 (PDE6B, American Staffordshire Terrier Variant)	Clear
Progressive Retinal Atrophy, crd4/cord1 (RPGRIP1)	Clear
Progressive Retinal Atrophy, PRA1 (CNGB1)	Clear
Progressive Retinal Atrophy, PRA3 (FAM161A)	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
O Pyruvate Kinase Deficiency (PKLR Exon 7, Labrador Retriever Variant)	Clear

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OTHER RESULTS

Pyruvate Kinase Deficiency (PKLR Exon 7, Pug Variant)	Clear
Raine Syndrome (FAM20C)	Clear
Recurrent Inflammatory Pulmonary Disease, RIPD (AKNA, Rough Collie Variant)	Clear
Renal Cystadenocarcinoma and Nodular Dermatofibrosis (FLCN Exon 7)	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
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
O Thrombopathia (RASGRP1 Exon 5, American Eskimo Dog Variant)	Clear

Registration: American Kennel Club (AKC)



DNA Test Report

Test Date: September 1st, 2023

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OTHER RESULTS

O Thrombopathia (RASGRP1 Exon 5, Basset Hound Variant)	Clear
O Thrombopathia (RASGRP1 Exon 8, Landseer Variant)	Clear
Trapped Neutrophil Syndrome, TNS (VPS13B)	Clear
O Ullrich-like Congenital Muscular Dystrophy (COL6A3 Exon 10, Labrador Retriever Variant)	Clear
Ullrich-like Congenital Muscular Dystrophy (COL6A1 Exon 3, Landseer Variant)	Clear
O Unilateral Deafness and Vestibular Syndrome (PTPRQ Exon 39, Doberman Pinscher)	Clear
Urate Kidney & Bladder Stones (SLC2A9)	Clear
✓ Von Willebrand Disease Type II, Type II vWD (VWF, Pointer Variant)	Clear
Von Willebrand Disease Type III, Type III vWD (VWF Exon 4, Terrier Variant)	Clear
Von Willebrand Disease Type III, Type III vWD (VWF Intron 16, Nederlandse Kooikerhondje Variant)	Clear
Von 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

Registration: American Kennel Club (AKC)





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HEALTH REPORT

Notable result

ALT Activity

Sandy inherited both copies of the variant we tested for Alanine Aminotransferase Activity

Why is this important to your vet?

Sandy has two copies of a variant in the GPT gene and is likely to have a lower than average baseline ALT activity. ALT is a commonly used measure of liver health on routine veterinary blood chemistry panels. As such, your veterinarian may want to watch for changes in Sandy's ALT activity above their current, healthy, ALT activity. As an increase above Sandy's baseline ALT activity could be evidence of liver damage, even if it is within normal limits by standard ALT reference ranges.

What is Alanine Aminotransferase Activity?

Alanine aminotransferase (ALT) is a clinical tool that can be used by veterinarians to better monitor liver health. This result is not associated with liver disease. ALT is one of several values veterinarians measure on routine blood work to evaluate the liver. It is a naturally occurring enzyme located in liver cells that helps break down protein. When the liver is damaged or inflamed, ALT is released into the bloodstream.

How vets diagnose this condition

Genetic testing is the only way to provide your veterinarian with this clinical tool.

How this condition is treated

Veterinarians may recommend blood work to establish a baseline ALT value for healthy dogs with one or two copies of this variant.





Fembark

DNA Test Report

HEALTH REPORT

Notable result

Ichthyosis, ICH1

Sandy inherited one copy of the variant we tested for Ichthyosis, ICH1

What does this result mean?

This result should not impact Sandy's health but it could have consequences for siblings or other related dogs if they inherited two copies of the variant. We recommend discussing this result with their owners or breeders if you are in contact.

Impact on Breeding

Your dog carries this variant and will pass it on to ~50% of her offspring.

What is Ichthyosis, ICH1?

This skin disorder gets its name from the thick, darkly pigmented scales of skin ("ichthys" is Greek for "fish") that affected dogs display over most areas of the body, not including the head or extremities.

When signs & symptoms develop in affected dogs

As puppies, affected dogs can show signs of scaling. This disease tends to worsen with age.

How vets diagnose this condition

Examining the characteristic lesions is the first step in diagnosing lchthyosis. Confirmatory genetic testing and/or skin biopsies can also be performed.

How this condition is treated

There is no definitive treatment for ichthyosis: typically, ichthyotic dogs are maintained on a continuous treatment of mild antidandruff shampoos and moisturizing rinses. This is a chronic and frustrating condition to manage.

Actions to take if your dog is affected

• Following your veterinarian's advice on skin care and nutrition is the best way to manage ichthyosis.







5%

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INBREEDING AND DIVERSITY

CATEGORY

Coefficient Of Inbreeding

Our genetic COI measures the proportion of your dog's genome where the genes on the mother's side are identical by descent to those on the father's side.

MHC Class II - DLA DRB1

A Dog Leukocyte Antigen (DLA) gene, DRB1 encodes a major histocompatibility complex (MHC) protein involved in the immune response. Some studies have shown associations between certain DRB1 haplotypes and autoimmune diseases such as Addison's disease (hypoadrenocorticism) in certain dog breeds, but these findings have yet to be scientifically validated.

MHC Class II - DLA DQA1 and DQB1

DQA1 and DQB1 are two tightly linked DLA genes that code for MHC proteins involved in the immune response. A number of studies have shown correlations of DQA-DQB1 haplotypes and certain autoimmune diseases; however, these have not yet been scientifically validated.

RESULT

Your Doys COI : 5%

High Diversity

How common is this amount of diversity in mixed breed dogs:



High Diversity

How common is this amount of diversity in mixed breed dogs:



