

Horse: All About The Gifts Owner: Justin Stohlman

Horse and Owner Information

Horse	All About The Gifts	DOB	2014-05-03
Breed	Paint	Age	3 years, 6 months
Color	Bay Overo	Sex	Mare
Discipline	Hunter Under Saddle	Height	16.1 hands
Registry	APHA	Reg Number	1050225
Sire	The Gift Of Midas	Dam	Alloutplacestolook
Sire Reg & No.	APHA 836785	Dam Reg & No.	APHA 793497
Comments			

Owner

Justin Stohlman

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Results Summary

Coat Color:

All About The Gifts has one Red allele and one Black, indicating her base coat color appears Black. Two copies of the Dominant Agouti allele were detected; invisible on a Red base, it pushes/restricts Black out to points; legs, ear tips, etc. appearing Bay. One Dominant White 20 allele and one Frame/Lethal White Overo allele each was detected which may result in White markings. As a result of the allele count in each of the following, she has a minimum 50% chance of passing Red or Black, and 100% Dominant Agouti, and 50% Dominant White 20 and/or Frame/Lethal White Overo to any offspring.

Allele Summary: AA, Ee, W20/n, LWO/n, CT (Mid-distance Type), Gaited/n

Traits:

All About The Gifts' testing indicated the presence of one Frame/Lethal White Overo (LWO) allele resulting in "Carrier" status. Caution is recommended when breeding to avoid another carrier and thus, 25% chance of foal death. Her testing has indicated the presence of one DMRT3 (Gaited) allele, and she may, therefore, pass it to 50% of any offspring.

Please note:

Your analysis is ongoing and may include some regions marked with an asterisk denoting the following.

* Discovery - This gene detection is in the early stages of discovery and will have varying reliability results.

** Inconclusive - Not a bad omen! Simply put, the gene of interest did not reveal itself (neither a positive nor a negative; no result, therefore unknown).

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Coat Color Results

lase				
Agouti	+/+	ASIP	AA - Two dominant Agouti elleles detected; restricts any Black base to appear Bay.	More about A
Black/Red	+/-	MC1R	Ee - One Black allele detected and one Red.	More about E
odifier				
Brindle/IP	4-	IKBKG	No Brindle/IP alleles detected.	More about IP
Grey	4-	STX17A	No Grey alleles detected	More about G
ilution				
Champagne	4-	SLC36A1	No Champagne alleles detected	More about CH
Cream	4-	SLC45A2	No Cream alleles detected	More about CR
Dun	4-4-	твхз	nd2/nd2 (non-dun). Two non-dun2 alleles detected. No Dun or non-Dun Primitive Marking alleles detected.	More about Dun
Pearl	afe.	SLC45A2	No Pearl alleles detected.	More about pri
Silver	4-	PMEL17	No Silver aliales detected.	More about Z

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Coat Color Results, continued

White Patterns Results				
Dominant White	+/-	KIT	W20/n - One Dominant White 20 allele detected (DW1-21).	More about DW
Frame Overo (LWO)	+/-	EDNRB	LWO/n - One Frame Overo (LWO) allele detected.	More about LWO
Leopard Complex Spotting (LP)	4-	TRPM1	No Leopard Complex Spotting (LP) alleles detected.	More about LP
Pattern 1 (LP modification)	4-	RFWD3	No Pattern 1 (LP modification) alleles detected.	More about PATN1
Splashed White (MITF)	$d_{-1}d_{-1}$	MITF	No Splashed White 1 nor Splashed White 3 alleles detected.	More about SW (MITF)
Splashed White (PAX3)	-1-,-1-	PAX3	No Splashed White 2 nor Splashed White 4 alleles detected.	More about SW (PAX3)
Sabino 1	4-	KIT	No Sabino 1 alleles detected	More about SB1
Tobiano	-1-	ECA3	No Tobiano alleles detected,	More about TO



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Health Genetics 1

mmuna Suatam				
mmune System Foal Immunodeficiency Syndrome	4-	SLC5A3	No Foal immunodeficiency Syndrome alleles detected.	More about fis
Severe Combined Immunodeficiency	4-	DNAPK	No Severe Combined Immunodeficiency alleles detected.	More about soid
West Nile*	4.	OAS1	Normal susceptibility to West Nile Virus,	More about WNVR*
Muscle Disorders				
Glycogen Branching Enzyme Deficiency	-/-	GBE1	No Glycogen Branching Enzyme Deficiency alleles detected.	More about gbed
Hyperkalemic Periodic Paralysis	4-	SCN4A	No Hyperkalemic Periodic Paralysis alleles detected.	More about HYPP
Malignant Hyperthermia	4-	RYR1	No Mailgnant Hyperthermia alleles detected.	More about MH
Myotonia	-/-	CLCN4	No Myotonia alleles detected.	More about myt
Polysaccharide Storage Myopathy (type 1)	J.	GYS1	No Polysaccharida Storage Myopathy (type 1) alleles detected.	More about PSSM1



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Health Genetics 2

Neurologic Disorders				
Cerebellar Abiotrophy	4-	MUTYH	No Cerebellar Abiotrophy alleles detected.	More about ca
Lavender Foal Syndrome	4-	MYOSA	No Lavender Foal Syndrome alleles detected.	More about its
Reproductive Disorders				
Androgen Insensitivity	-/-	AR	No Androgen Insensitivity alleles detected.	More about as
IAR - Subfertility*	+/-,+/-	FKBP6	Two IAR Subfertility* alleles detected; likely no effect,	More about iar*
Skin Disorders				
Hereditary Equine Regional Dermal Asthenia	4-	PPIB	No Hereditary Equine Regional Dermal Asthenia alleles detected.	More about herda
Junctional Epidermolysa Bullosis (type 1)	4-	LAMCZ	No Junctional Epidermolysa Bullosis (type 1) alleles detected.	More about jeb1
Junctional Epidermolysa Bullosis (type 2")	4-	LAMA3	No Junctional Epidermolysa Bullosis (type 2*) alleles detected.	More about jeb2*
Warmblood Fragile Foal Syndrome	4-	PLOD1	No Warmblood Fragile Foal Syndrome alleles detected.	More about WFFS



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Other Genetics

Trait Genetics				
Lordosis*	+/-,-/-,-/-	ECA20	No pattern of Lordosis* alleles detected.	More about L*
Curiosity/Vigilance*	+/+	DRD4	Cur - GG - Two Curiosity alleles detected; likely more curious than vigilant.	More about Cur/Vig
Myostatin/Speed	*/-	MSTN	CT (Mid-distance Type) - One Sprint and one Endurance allele detected; likely Mid-distance ability.	More about MSTN
Gait	*/-	DMRT3	Gaitedin - One Gait allele detected.	More about Gaited



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Inheritance Probabilities



Coat Color Inheritance Probabilities: The bar graph above depicts the number of alleles for specific coat color phenotypes based upon your horse's genetic testing results. Completely filled red bar represents two such alleles (homozygous) and a half-filled yellow bar represents one such allele (heterozygous).

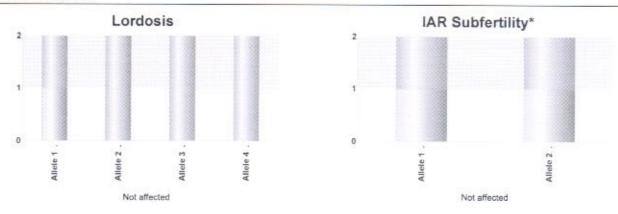
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Inheritance Probabilities



Multi-allele Risk Charts: Each chart represents a trait, and each bar indicates a distinct risk or allele presence. These act in combination to produce the trait. A red bar indicates the horse carries 2 risk alleles at the site; a partly-yellow bar indicates 1 risk allele; and a fully-grey bar indicates 0 risk alleles. If all bars are red, then the horse carries two risk alleles at each risk site and is likely affected. If all bars contain yellow or red, but are not all red, then the horse is likely a carrier. Otherwise, the horse is not a likely a carrier of the tested trait.



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Defining Genetics & More Info

Allele: One of two or more alternative forms of a gene that arise by mutation and are found at the same place on a chromosome.

Alleles: Heterozygous vs. Homozygous? Allele calls are written in a way that denotes their origin and whether they are DOMINANT (uppercase) or recessive (lowercase). For example, at MC1R (also known as extension), Black is dominant and thus written as "E" whereas Red is recessive and thus denoted as "e". Therefore, an

EE horse is homozygous for Black (and thus appears black), an ee horse is homozygous for Red (appears Red), and an Ee horse is

heterozygous (shows the dominant allele, thus is Black).

Gene: A unit of heredity that is transferred from a parent to offspring and is thought to determine some characteristic of the offspring.

Genotype: The genetic constitution or make up of an individual organism.

Heterozygous: A pair of genes which are different (not the same). One is typically dominant and one recessive.

Homozygous: A pair of genes that are identical (of one type).

Phenotype: The observable or visible characteristics of an individual resulting from their genotype or the interaction of their various genes and environment.

The results depicted in this report do not constitute veterinary or medical advice. Any medical of veterinary advice should be sought from your veterinarian regarding these results or any health issues or questions you may have about your animal. Breed, sex, gene interaction, unknown genes and individual variances may impact the results, phenotypes, and behaviors in any animal in unknown and unpredictable ways. Please be advised that your animals' health is important to us and you should feel free to contact us should you have any further questions or feedback on our diagnostic platform, results reporting, or general questions. We value your input and thank you!

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