



**EQUINE JUVENILE SPINOCEREBELLAR ATAXIA
 TEST REPORT**

<i>Provided Information:</i>		<i>Case:</i>	NQ113484
<i>Name:</i>	LONE WOOLF	<i>Date Received:</i>	12-Aug-2024
<i>Registration:</i>	6094246	<i>Report Issue Date:</i>	14-Aug-2024
		<i>Report ID:</i>	9751-2127-4094-7091
Verify report at vgl.ucdavis.edu/verify			
<i>DOB:</i> 04/03/2020 <i>Sex:</i> Stallion <i>Breed:</i> Quarter Horse			
<i>Sire:</i>	METALLIC REBEL	<i>Dam:</i>	A LITTLE REYLENA
<i>Reg:</i>	5506591	<i>Reg:</i>	4522460
<i>Microchip:</i>		<i>Microchip:</i>	

RESULT

INTERPRETATION

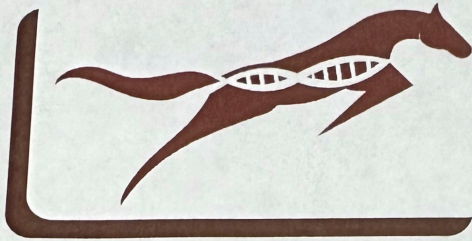
Equine Juvenile Spinocerebellar Ataxia	N/N
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Normal. No copies of the allele associated with equine juvenile spinocerebellar ataxia (EJSCA) detected.

One Page Diagnostics Report

EtalonDX.com

LabID: 042924_047



Horse: Lone Woolf
Breed: American Quarter Horse
Date of Birth: N/A
Sex: Stallion

Owner: Samantha Gillis
Phone:
Email: sammy.gillis@icloud.com

Upgrade available for Performance Variants (Temperament, Speed, Gait)

UPGRADE NOW

Variant Summary



MYHM: n/n

Performance Variants

Trait	Genotype	Notes
Upgrade now to access Performance results (Temperament, Speed, Gait)		

Health Results

Trait	Genotype	Notes
Myosin-Heavy Chain Myopathy (MYHM)	n/n	No variant detected - Negative

Coat Color Results

Trait	Genotype	Notes
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AMERICAN QUARTER HORSE GENETIC HEALTH PANEL TEST REPORT

<p><i>Client/Owner/Agent Information:</i> AMERICAN QUARTER HORSE ASSOCIATION</p> <p><i>Provided Information:</i> <i>Name:</i> LONE WOLF <i>Registration:</i> 6094246</p>	<p><i>Date Received:</i> 07-Jan-2021 <i>Report Issue Date:</i> 04-Sep-2024 <i>Report ID:</i> 9597-1567-4039-1171 <i>Reissue of:</i> 4760-5394-2252-5108</p>
<p><i>DOB:</i> 04/03/2020 <i>Sex:</i> Stallion <i>Breed:</i> Quarter Horse <i>Alt. ID:</i> 7364062</p>	
<p><i>Sire:</i> METALLIC REBEL <i>Dam:</i> A LITTLE REYLENA <i>Reg:</i> 5506591 <i>Reg:</i> 4522460 <i>Microchip:</i> <i>Microchip:</i></p>	

RESULT

INTERPRETATION

Test Name	Result	Interpretation
Glycogen Branching Enzyme Deficiency (GBED)	N/G	Carrier. One copy of the GBED allele detected.
Hereditary Equine Regional Dermal Asthenia (HERDA)	N/N	Normal. No copies of the HERDA allele detected.
Hyperkalemic Periodic Paralysis (HYPP)	N/N	Normal. No copies of the HYPP allele detected.
Malignant Hyperthermia (MH)	N/N	Normal. No copies of the MH allele detected.
Polysaccharide Storage Myopathy Type 1 (PSSM1)	N/N	Normal. No copies of the PSSM1 allele detected.

Additional Information

If testing for a disease or a disorder was performed and results indicate the animal is affected or at risk, we recommend contacting your veterinarian for further clinical evaluation and for additional information on disease and management.

For more detailed information on American Quarter Horse Genetic Health Panel test results, please visit our website at: vgl.ucdavis.edu/panel/quarter-horse-disease-panel

License Information

The GBED test is performed under a license agreement with the University of Minnesota.

Results are determined using PCR-based methods. The results relate only to the sample tested as identified by the submitter (for example, identity and/or breed).

Report authorized by Dr. Rebecca Bellone, VGL Director

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