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Check us out online at-----  
[www.WoronieckiRanchQuarterHorses.com](http://www.WoronieckiRanchQuarterHorses.com)  
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[woronieckiranch@westriv.com](mailto:woronieckiranch@westriv.com)

## 7 Identified Diseases Information as it Pertains to Woroniecki Ranch Quarter Horses

At Woroniecki Ranch Quarter Horses we take an ethical response to any genetic diseases as they are identified. AQHA previously had a 5-panel test requirement for breeding stallions since 2015. Two more diseases have been identified and AQHA has now required a 6-panel test. A 7<sup>th</sup> disease has been identified and could soon be added to the panel. We, as well as many other breeders, have decided to test for that (EJSCA). We also know that there could be many more diseases yet to be discovered. We order our tests through the VGL laboratory of the School of Veterinary Medicine at the University of California, Davis and provide those results to AQHA and buyers. VGL is internationally recognized as a pioneer and expert in DNA-based animal testing. The effects of these equine diseases are wide-ranging, from mild and manageable to severe and terminal. We have compiled a short description of each disorder tested. **In many instances we only test the necessary specific test based upon the parents' test results. If both parents are N/N on all or some diseases then the offspring is also N/N on those diseases by default. Please see ALL PAGES of this document link.**

**Glycogen Branching Enzyme Deficiency (GBED)** is a fatal genetic disorder that results from the inability to correctly store glycogen in several organs of the body. Most die within couple weeks of age, but none have been known to survive more than 2 months of age. These foals are often still born. **GBED is a recessive trait and only horses that inherit both recessive genes from each parent (G/G) will be afflicted. Carriers (N/G) and non-carriers (N/N) will have no problems in their lives as they will NOT be afflicted at all, and they will be able to perform all performance activities. If you decide to breed a carrier (N/G) it is highly advised to not breed to another carrier to avoid producing afflicted offspring.**

**Hereditary Equine Regional Dermal Asthenia (HERDA)** is an inherited skin condition primarily found in Quarter Horses that is characterized by hyperextensible skin, scarring, and severe lesions along the back of affected horses. **HERDA is a recessive trait and only horses that inherit both recessive genes from each parent (HRD/HRD) will be afflicted. Carriers (N/HRD) and non-carriers (N/N) will have no problems in their lives as they will NOT be afflicted at all, and they will be able to perform all performance activities. If you decide to breed a carrier (N/HRD) it is highly advised to not breed to another carrier to avoid producing afflicted offspring.**

**Hyperkalemic Periodic Paralysis (HYPP)** is an inherited disease of the muscles primarily found in Quarter Horses which is characterized by sporadic episodes of muscle tremors or paralysis. **HYPP is a dominant trait and carriers (N/H) will be afflicted but can be managed with careful nutritional care. It is highly recommended NOT to breed a carrier.**

**Formerly known as IMM, Myosin-heavy chain myopathy (MYHM)** is a muscle disease in Quarter Horses and related breeds that results in two distinct clinical disease presentations. The first presentation is called immune-mediated myositis or IMM and it is characterized by episodes of severe muscle atrophy following an autoimmune event. The second is severe muscle pain and damage termed non-exertional rhabdomyolysis or "tying-up" that is not associated with exercise and may or may not have muscle atrophy. **MYHM is a codominant trait and carriers (N/My) may develop a myosin-heavy chain myopathy. Horses with (My/My) may develop a more severe form of a myosin-heavy chain myopathy. It is highly recommended NOT to breed a carrier.** After consulting with veterinarians and experts in breeding who deem this disorder to not be as severe or common as HYPP or PSSM1, we have decided at this time to continue to breed certain individuals identified at WRQH. We will not breed carriers to carriers to minimize the potential. We have several aged horses that carry MYHM and have had no problems with them. If things prove differently, we will adjust at that time.

**Malignant Hyperthermia (MH)** is an inherited disease in which affected horses can be triggered by halogenated anesthetics, succinylcholine, stress, or excitement, which can induce a hyper-metabolic state characterized by symptoms including muscle contracture, elevated temperature, and an irregular heart rhythm. **MH is a dominant trait, and carriers (N/MH) will be afflicted if undergoing surgery or extreme stress. It is highly recommended NOT to breed a carrier.**

**Polysaccharide Storage Myopathy (PSSM1)** is a glycogen storage disease that results in the accumulation of abnormal complex sugars in muscle cells, which can lead to muscle pain, weakness, and reluctance to move. **PSSM1 is a dominant trait but carriers (N/PSSM1) can be managed with proper diet and exercise. It is highly recommended NOT to breed a carrier.**

**Equine Juvenile Spinocerebellar Ataxia (EJSCA)** is an inherited neurologic disease that causes ataxia. Affected foals develop ataxia, or incoordination, between 1 and 4 weeks of age. The disorder progresses within a few days until affected foals are unable to stand without assistance. **EJSCA is a recessive trait and only horses that inherit both recessive genes from each parent (JSA/JSA) will be afflicted. Carriers (N/JSA) and non-carriers (N/N) will have no problems in their lives as they will NOT be afflicted at all and they will be able to perform all performance activities. If deciding to breed a carrier (N/JSA) it is highly advised to not breed to another carrier to avoid producing afflicted offspring.**

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**Whiskey Up Gondor JW (AQHA)**

**2025 Buckskin Stallion**

GBED Status	N/N
HERDA Status	N/N
HYPP Status	N/N
MYHM Status	N/N
MH Status	N/N
PSSM1 Status	N/N
EJSCA Status	N/N

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# GLYCOGEN BRANCHING ENZYME DEFICIENCY (GBED) TEST REPORT

<b>Provided Information:</b> Name: <b>WHISKEY UP GONDOR JW</b> Registration: <b>AQHA PENDING</b>		<b>Case:</b> <b>NQ127370</b> <b>Date Received:</b> 28-Jul-2025 <b>Report Issue Date:</b> 30-Jul-2025 <b>Report ID:</b> 1466-8323-6227-2113 Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a>
DOB: <b>05/10/2025</b> Sex: <b>Stallion</b> Breed: <b>Quarter Horse</b>		
Sire: WHOOP UP WHISKEY Reg: 5359075 Microchip:		Dam: CABALLERO DRIFT JW Reg: 5741394 Microchip:

## RESULT

## INTERPRETATION

Glycogen Branching Enzyme Deficiency (GBED)	N/N
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Normal. No copies of the GBED allele detected.



VETERINARY GENETICS LABORATORY  
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## AQHA GENETIC DISEASE PANEL TEST RESULTS

<b>AMERICAN QUARTER HORSE ASSOCIATION</b> P.O. BOX 200 AMARILLO, TX 79168-0001		<b>Case:</b> <b>QHA192968</b> <b>Date Received:</b> 11-May-2015 <b>Print Date:</b> 15-May-2015 <b>Report ID:</b> 0187-9866-2321-9024 Verify report at <a href="http://www.vgl.ucdavis.edu/myvgl/verify.html">www.vgl.ucdavis.edu/myvgl/verify.html</a>
<b>Horse: WHOOP UP WHISKEY</b> <b>YOB: 2010 Sex: Stallion Breed: Quarter Horse Alt. ID: 6281311</b>		<b>Reg: 5359075</b>
<b>Sire: PADDYS IRISH WHISKEY</b> <b>Dam: MY LITTLE SUGAR BABE</b>		<b>Reg: 2983308</b> <b>Reg: 3179872</b>

GBED	N/G	N/G - Carrier - Heterozygous (one normal and one GBED gene)
HERDA	N/N	N/N - Normal - horse does not have the HERDA gene
HYPP	N/N	N/N - Normal - Does not possess the disease-causing HYPP gene
MH	N/N	N/N - Normal - horse does not have the MH gene
PSSM1	N/N	N/N - Normal - horse does not have the PSSM1 gene

GBED - Glycogen Branching Enzyme Deficiency. Fatal disease of newborn foals caused by defect in glycogen storage. Affects heart and skeletal muscles and brain. Inherited as recessive disease.

HERDA - Hereditary Equine Regional Dermal Asthenia. Skin disease characterized by hyperextensible skin, scarring, and severe lesions along the back of affected horses. Typical onset is around 2 years of age. Inherited as a recessive disease.

HYPP - Hyperkalemic Periodic Paralysis. Muscle disease caused by defect in sodium channel gene that causes involuntary muscle contraction and increased level of potassium in blood. Inherited as dominant disease. Two copies of defective gene produce more severe signs than one copy.

MH - Malignant Hyperthermia. Rare but life-threatening skeletal muscle disease triggered by exposure to volatile anesthetics (halothane), depolarizing muscle relaxants (succinylcholine), and stress. Presumed inheritance as dominant disease.

PSSM1 - Polysaccharide Storage Myopathy Type 1. Muscle disease characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include muscle pain, stiffness, skin twitching, sweating, weakness and reluctance to move. Inherited as a dominant disease.

GBED testing performed under a license agreement with the University of Minnesota.

HERDA testing performed under a license agreement with the University of California, Davis.

PSSM1 testing performed under a license agreement with the American Quarter Horse Association.

## EQUINE DISEASE TEST REPORT

<b>Provided Information:</b>		<b>Case:</b>	<b>HRD6503</b>
<b>Name:</b>	<b>WHOOOP UP WHISKEY</b>	<b>Date Received:</b>	10-Apr-2012
<b>Registration:</b>	<b>5359075</b>	<b>Report Issue Date:</b>	18-Mar-2025
		<b>Report ID:</b>	5718-7192-8743-4015
		<b>Reissue of:</b>	0294-0065-4700-9066
		Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a>	
<b>DOB: 06/03/2010   Sex: Stallion   Breed: Quarter Horse</b>			
<b>Sire:</b>	PADDYS IRISH WHISKEY	<b>Dam:</b>	MY LITTLE SUGAR BABE
<b>Reg:</b>	2983308	<b>Reg:</b>	3179872
<b>Microchip:</b>		<b>Microchip:</b>	

### RESULT

### INTERPRETATION

<b>Hereditary Equine Regional Dermal Asthenia (HERDA)</b>	<b>N/N</b>	Normal. No copies of the HERDA allele detected.
<b>Myosin-Heavy Chain Myopathy (MYHM)</b>	<b>N/N</b>	Normal. No copies of the MYHM allele detected. Horse does not have increased susceptibility for immune mediated myositis or nonexertional rhabdomyolysis caused by the MYHM allele.

## EQUINE JUVENILE SPINOCEREBELLAR ATAXIA TEST REPORT

<b>Provided Information:</b>  Name: <b>WHOOOP UP WHISKEY</b> Registration: <b>5359075</b>	<b>Case: HRD6503</b> Date Received: 10-Apr-2012 Report Issue Date: 09-Jan-2025 Report ID: 2894-5255-0443-5144  <div style="text-align: right; font-size: small;">Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a></div>						
DOB: <b>06/03/2010</b> Sex: <b>Stallion</b> Breed: <b>Quarter Horse</b>							
<table style="width: 100%;"> <tr> <td style="width: 50%;">Sire: PADDYS IRISH WHISKEY</td> <td style="width: 50%;">Dam: MY LITTLE SUGAR BABE</td> </tr> <tr> <td>Reg: 2983308</td> <td>Reg: 3179872</td> </tr> <tr> <td>Microchip:</td> <td>Microchip:</td> </tr> </table>		Sire: PADDYS IRISH WHISKEY	Dam: MY LITTLE SUGAR BABE	Reg: 2983308	Reg: 3179872	Microchip:	Microchip:
Sire: PADDYS IRISH WHISKEY	Dam: MY LITTLE SUGAR BABE						
Reg: 2983308	Reg: 3179872						
Microchip:	Microchip:						

### RESULT

### INTERPRETATION

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

## MYOSIN-HEAVY CHAIN MYOPATHY (MYHM) TEST REPORT

<b>Provided Information:</b> Name: <b>CABALLERO DRIFT JW</b> Registration: <b>5741394</b>		<b>Case: NQ125772</b> Date Received: 18-Jun-2025 Report Issue Date: 30-Jun-2025 Report ID: 1925-5273-7137-4151 Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a>
DOB: 05/01/2016 Sex: Mare Breed: Quarter Horse		
Sire: WALTER O RIELLY Reg: 4343282 Microchip:		Dam: MUJER DEWEY DIAL JW Reg: 4975025 Microchip:

### RESULT

### INTERPRETATION

Myosin-Heavy Chain Myopathy (MYHM)	N/N
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Normal. No copies of the MYHM allele detected. Horse does not have increased susceptibility for immune mediated myositis or nonexertional rhabdomyolysis caused by the MYHM allele.

## EQUINE JUVENILE SPINOCEREBELLAR ATAXIA TEST REPORT

<b>Provided Information:</b>  Name: <b>CABALLERO DRIFT JW</b>  Registration: <b>5741394</b>	<b>Case: NQ125772</b>  Date Received: 18-Jun-2025 Report Issue Date: 25-Jun-2025 Report ID: 1100-1320-6369-2139  <div style="text-align: right; font-size: small;">Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a></div>
DOB: <b>05/01/2016</b> Sex: <b>Mare</b> Breed: <b>Quarter Horse</b>	
Sire: WALTER O RIELLY Reg: 4343282 Microchip:	Dam: MUJER DEWEY DIAL JW Reg: 4975025 Microchip:

### RESULT

### INTERPRETATION

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



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## AQHA GENETIC DISEASE PANEL TEST RESULTS

<b>AMERICAN QUARTER HORSE ASSOCIATION</b> P.O. BOX 200 AMARILLO, TX 79168-0001		<b>Case: QHA192970</b> <b>Date Received: 11-May-2015</b> <b>Print Date: 15-May-2015</b> <b>Report ID: 9633-4604-1417-2011</b> Verify report at <a href="http://www.vgl.ucdavis.edu/myvgi/verify.html">www.vgl.ucdavis.edu/myvgi/verify.html</a>
<b>Horse: WALTER O RIELLY</b> YOB: 2000 Sex: Stallion Breed: Quarter Horse Alt. ID: 5067985		<b>Reg: 4343282</b>
<b>Sire: BRADYWOOD</b> <b>Dam: LACYS KEEPSAKE</b>		<b>Reg: 3190953</b> <b>Reg: 3169660</b>

GBED	N/N	N/N - Normal - Does not possess the disease-causing GBED gene
HERDA	N/N	N/N - Normal - horse does not have the HERDA gene
HYPP	N/N	N/N - Normal - Does not possess the disease-causing HYPP gene
MH	N/N	N/N - Normal - horse does not have the MH gene
PSSM1	N/N	N/N - Normal - horse does not have the PSSM1 gene

GBED - Glycogen Branching Enzyme Deficiency. Fatal disease of newborn foals caused by defect in glycogen storage. Affects heart and skeletal muscles and brain. Inherited as recessive disease.

HERDA - Hereditary Equine Regional Dermal Asthenia. Skin disease characterized by hyperextensible skin, scarring, and severe lesions along the back of affected horses. Typical onset is around 2 years of age. Inherited as a recessive disease.

HYPP - Hyperkalemic Periodic Paralysis. Muscle disease caused by defect in sodium channel gene that causes involuntary muscle contraction and increased level of potassium in blood. Inherited as dominant disease. Two copies of defective gene produce more severe signs than one copy.

MH - Malignant Hyperthermia. Rare but life-threatening skeletal muscle disease triggered by exposure to volatile anesthetics (halothane), depolarizing muscle relaxants (succinylcholine), and stress. Presumed inheritance as dominant disease.

PSSM1 - Polysaccharide Storage Myopathy Type I. Muscle disease characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include muscle pain, stiffness, skin twitching, sweating, weakness and reluctance to move. Inherited as a dominant disease.

GBED testing performed under a license agreement with the University of Minnesota.

HERDA testing performed under a license agreement with the University of California, Davis.

PSSM1 testing performed under a license agreement with the American Quarter Horse Association.



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## AQHA GENETIC DISEASE PANEL TEST RESULTS

<b>AMERICAN QUARTER HORSE ASSOCIATION</b> P.O. BOX 200 AMARILLO, TX 79168-0001	<b>Case: QHA206630</b> <b>Date Received: 31-Aug-2015</b> <b>Print Date: 07-Sep-2015</b> <b>Report ID: 2507-1961-5218-7124</b> Verify report at <a href="http://www.vgl.ucdavis.edu/myvgl/verify.html">www.vgl.ucdavis.edu/myvgl/verify.html</a>
<b>Horse: MUJER DEWEY DIAL JW</b> <b>YOB: 2007 Sex: Mare Breed: Quarter Horse Alt. ID: 5848427</b>	<b>Reg: 4975025</b>
<b>Sire: MUJER TACKY JAY</b> <b>Dam: MISS LORI JAY</b>	<b>Reg: 2580521</b> <b>Reg: 3020482</b>

<b>GBED</b>	<b>N/N</b>
<b>HERDA</b>	<b>N/N</b>
<b>HYPP</b>	<b>N/N</b>
<b>MH</b>	<b>N/N</b>
<b>PSSM1</b>	<b>N/N</b>

N/N - Normal - Does not possess the disease-causing GBED gene

N/N - Normal - horse does not have the HERDA gene

N/N - Normal - Does not possess the disease-causing HYPP gene

N/N - Normal - horse does not have the MH gene

N/N - Normal - horse does not have the PSSM1 gene

**GBED** - Glycogen Branching Enzyme Deficiency. Fatal disease of newborn foals caused by defect in glycogen storage. Affects heart and skeletal muscles and brain. Inherited as recessive disease.

**HERDA** - Hereditary Equine Regional Dermal Asthenia. Skin disease characterized by hyperextensible skin, scarring, and severe lesions along the back of affected horses. Typical onset is around 2 years of age. Inherited as a recessive disease.

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**PSSM1** - Polysaccharide Storage Myopathy Type 1. Muscle disease characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include muscle pain, stiffness, skin twitching, sweating, weakness and reluctance to move. Inherited as a dominant disease.

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