

Jodie & Warren Woroniecki 7075 28th St. Hebron, ND 58638 701-878-4088 Check us out online at---www.WoronieckiRanchQuarterHorses.com Or email, call or stop by the ranch. woronieckiranch@westriv.com

5 Panel Information as it Pertains to Woroniecki Ranch Quarter Horses

At Woroniecki Ranch Quarter Horses we order a genetic kit through AQHA and the results are sent to VGL laboratory of the School of Veterinary Medicine at the University of California, Davis. 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) doesn't allow a foal to store enough sugar in its cells for energy, function of the brain, heart and skeletal muscles. 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 deciding 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) causes the skin on a horse's back to literally peel away. The skin will slough becoming loose and tented to never return to its original position. HERDA is a recessive trait and only horses that inherit both recessive genes from each parent (HDR/HDR) will be afflicted. Carries (N/HDR) and non-carries (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/HDR) it is highly advised to not breed to another carrier to avoid producing afflicted offspring

Hyperkalemic Periodic Paralysis (HYPP) is a muscle condition that leads to weak muscles or severe twitching of the muscles. In most cases symptoms include tremors, weakness, cramping, sweating and inability to relax. In severe cases horse can collapse from a heart attack or respiratory failure and die. 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.

Malignant Hyperthermia (MH) is a rare but deadly disorder triggered by the use of anesthesia, muscle relaxant succinylcholine and stress. The horse will often experience high heart rate along with rapid breathing and extreme fever. This can also lead to death in some cases. Some horses are also a carrier of PSSM along with MH. MH is a dominant trait and carriers will be afflicted if undergoing surgery or extreme stress. It is highly recommended NOT to breed a carrier.

Polysaccharide Storage Myopathy (PSSM1) is when the muscles store too much glycogen causing muscle stiffness and muscle tying up. Most horses experience pain with strenuous exercise. 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.

Whiskey Up Sedna JW 2024 Bay Filly		(AQHA)	
GBED Status HERDA Status HYPP Status MH Status PSSM1 Status	N/N N/N N/N N/N N/N	Carries one copy of the GBED gene. If breeding mare, breed to N/N stallions	

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SANTA BARBARA . SANTA CRUZ

VETERINARY GENETICS LABORATORY SCHOOL OF VETERINARY MEDICINE ONE SHIELDS AVENUE DAVIS, CALIFORNIA 95616-8744

AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001 Case:QHA192968Date Received:11-May-2015Print Date:15-May-2015Report ID:0187-9866-2321-9024Verify report at www.vgl.ucdavis.edu/myvgl/verify.html

Horse: WHOOP UP WHISKEY

Reg: 5359075

TELEPHONE: (530) 752-2211

FAX: (530) 752-3556

YOB: 2010 Sex: Stallion Breed: Quarter	er Horse Alt. ID: 6281311	
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Sire: PADDYS IRISH WHISKEY	Reg: 2983308	
Dam: MY LITTLE SUGAR BABE	Reg: 3179872	

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



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Polysaccharide Storage Myopathy (PSSM1) is when the muscles store too much glycogen causing muscle stiffness and muscle tying up. Most horses experience pain with strenuous exercise. **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.

HQ Jacked Up Lights 2018 Sorrel Mare	(AQHA)	All NN by parentage. Parents' tests included.
GBED Status N/N HERDA Status N/N HYPP Status N/N MH Status N/N PSSM1 Status N/N		

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SANTA BARBARA + SANTA CRUZ

AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001			Case: Date	QHA168728 04-Dec-2014
			Print Date: Report ID: Verify report at	29-Apr-2019 0756-2277-8786-5100 I www.vgl.ucdavis.edu/myvgl/verify.htm
Horse: CD DYN YOB: 2012 Sex		uarter Horse Alt. ID: 6504701	Reg: 5522360	
Sire: CD LIGI Dam: ARC DY		0	3767592 4330746	
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		
НҮРР	N/N	N/N - Normal - Does not possess the disease-causing HYPP gene		
МН	N/N	N/N - Normal - horse does not have the MH gene		
PSSMI	N/N	N/N - Normal - horse does not have the PSSMI 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

HERDA - Hereditary Equine Regional Dennal 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 (succinyleholine), 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.

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AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICA P.O. BOX		HORSE ASSOCIATION	Case:	QHA245239
AMARILLO, TX 79168-0001			Date	28-Apr-2016
			Print Date:	29-Apr-2019
			Report ID: Verify report a	3967-6017-9256-7057 www.vgl.ucdavis.edu/myvgl/verify.htm
	WOOGI BOOGA Mare Breed: Qua	LOO rter Horse Ali. ID: 6415971	Reg: 5455160	
Sire: JACK TO	THE HILT	Reg	4647367	
Dam: DUTCHE	ESS JO JACK	Reg	3575533	
GBED	N/N	N/N - Normal - Docs not possess the disease-causing GBED gene		
HERDA	N/N	N/N - Normal - horse does not have the HERDA gene		
НҮРР	N/N	N/N - Normal - Does not possess the disease-causing HYPP gene		
МН	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		

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GLYCOGEN BRANCHING ENZYME DEFICIENCY

(GBED)

TEST REPORT

Provided Information:

Name: WHISKEY UP SEDNA JW

Registration: AQHA Pending

Case: Date Received: Report Issue Date: Report ID:

HQ JACKED UP LIGHTS

NQ114588

09-Sep-2024 18-Sep-2024 1652-6576-1151-8135

Registration. AGRA Fending

Verify report at vgl.ucdavis.edu/verify

DOB: 05/09/2024 Sex: Mare Breed: Quarter Horse

Sire: WHOOP UP WHISKEY

Reg: 5359075

Glycogen Branching

Enzyme Deficiency (GBED)

Microchip:

RESULT

N/G

INTERPRETATION

5900145

Carrier. One copy of the GBED allele detected.

Dam:

Reg: 590 Microchip: