

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 Pis	tol JW	(AQHA)
2020 Bay Stallion		
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

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

Horse: WHOOP UP WHISKEY

TELEPHONE: (530) 752-2211 FAX: (530) 752-3556

AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001

Case:	QHA192968
Date Received:	11-May-2015
Print Date:	15-May-2015
Report ID:	0187-9866-2321-9024
Verify report at www	.vgl.ucdavis.edu/myvgl/verify.html

Reg: 5359075

YOB: 2010 Sex: Stallion Breed: Quarter Horse Alt. ID: 6281311

	Sire: PADDYS IRISH WHISKEY Dam: MY LITTLE SUGAR BABE		Reg: 2983308
			Reg: 3179872
	GBED	N/G	N/G - Carrier - Hetcrozygous (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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PSSM1

TELEPHONE: (530) 752-7211 FAX: (530) 752-3556 SANTA BARBARA + SANTA CRUZ

AQHA GENETIC DISEASE PANEL TEST RESULTS

P.O. BO	AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001			Case: Date Received: Report Date: Report ID: Verify report at https://ww	QHA53836 19-Apr-2012 25-Apr-2012 6975-5757-7332-0004 wwwgl.ucdavis.edu/myvgl/verify.html
Horse: ANYV YOB: 07	WARE Breed: QH	Sex: M	Reg Alt. ID: 57873	: 4921132 60	
Sire: PLAY Dam: READ	GUN YTOWARE		Reg: 317 Reg: 38		
GBED	N/N	N/N - Normal - Doe	s not possess the disease-	causing GBED gene	
HERDA	N/N	N/N - Normal - hors	se does not have the HERI	DA gene	
НҮРР	N/N	N/N - Normal - Doe	s not possess the disease-o	ausing HYPP gene	
МН	N/N	N/N - Normal - hore	e does not have the MH g	ene	

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.

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

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.

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N/N

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GBED REPORT

JODIE WORONIECKI 7075 28TH ST. HEBRON, ND 58638	Case: Date Received:	NQ58088 04-Jun-2020	
	<i>Print Date:</i> <i>Report ID:</i> Verify report at wy	10-Jun-2020 5044-9504-8488-1155 w.vgl.ucdavis.edu/myvgl/verify.htm	
Horse: WHISKEY UP PISTOL JW DOB: 05/10/2020 Sex: Stallion Breed: Quarter Horse	Reg: AQHA Pending		
Sire: WHOOP UP WHISKEY	<i>Reg:</i> 5359075		
Dam: ANYWARE	<i>Reg:</i> 4921132		

GBED Test Result

N/N

Result Codes:

- G/G Affected Homozygous for GBED (two copies of the GBED gene).
- N/G Carrier Heterozygous (one normal and one GBED gene).
- N/N Normal Does not possess the disease-causing GBED gene.

The condition is inherited as a recessive trait. This means that breedings between two carrier (N/G) horses have a 25% chance of producing an affected foal (G/G). Affected foals usually die at a young age or will need to be euthanized due to weakness. Breedings between carrier and normal (N/N) horses produce only normal foals but 50% of these are expected to be carriers.

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