



Jodie & Warren Woronecki
7075 28th St.
Hebron, ND 58638
701-878-4088

Check us out online at----
www.WoroneckiRanchQuarterHorses.com
Or email, call or stop by the ranch.
woroneckiranch@westriv.com

5 Panel Information as it Pertains to Woronecki Ranch Quarter Horses

At Woronecki 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. **Carriers (N/HDR) 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/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.**

San Ella Drift JW (APHA 1,048,375)

2014 Buckskin Mare

GBED Status N/N

HERDA Status N/N

HYPP Status N/N

MH Status N/N

PSSM1 Status N/N

Equine Genetic Testing Report

Submitted By
Jodie & Warren Woroniecki
Woroniecki Ranch Quarter Horses
7075 28th St
Hebron, ND 58638



Subject Horse

Date Received: 10/28/2019

Horse Name: **San Ella Drift JW** Lab Reference #: **00130709**
 Breed: **Paint Horse** Registration: **1,048,375**
 Phenotype: **Buckskin** Birth: **2014**
 Sex: **Mare**

Sire
Sire Name: **Walter O Rielly**
Breed: **Paint Horse**
Registration: **613,648**
Phenotype: **Perlino**

Dam
Dam Name: **Leos San Ella JB**
Breed: **Paint Horse**
Registration: **747,042**
Phenotype: **Bay**

Coat Color and Pattern Testing

| | | | |
|---|------------------|---------|---|
| X | Tobiano | nn | Negative for Tobiano. |
| X | Frame Overo | nn | Negative for Frame Overo (LWO). |
| X | Sabino 1 | nn | Negative for the Sabino 1 gene. |
| X | Splashed White 1 | nn | Negative for the Splashed White SW1 mutation. |
| X | Splashed White 2 | nn | Negative for the Splashed White SW2 mutation. |
| X | Splashed White 3 | nn | Negative for the Splashed White SW3 mutation. |
| X | Appaloosa (LP) | lp/lp | Tested negative for the main Appaloosa LP gene and is NOT affected by CSNB. |
| X | PATN1 | n/n | Negative: Horse does not carry the PATN-1 gene mutation. |
| X | Red/Black Factor | Ee | Heterozygous. Horse is Black based but carries a recessive copy of the Red gene. |
| X | Agouti | AA | Homozygous for Agouti. Horse carries two copies of the Agouti gene. |
| X | Cream Dilution | nCr | Heterozygous. Single dilute. Horse carries one copy of the Cream Dilution gene. |
| X | Dun Dilution | nd1/nd2 | 1 copy of nd1 and 1 copy of nd2. Horse is not Dun diluted. Varying levels of primitive markings possible. |
| X | Silver Dilution | nn | Negative for Silver Dilution. |
| X | Champagne | nn | Negative for Champagne Dilution. |
| X | Pearl Dilution | nn | Negative for Pearl Dilution. |
| | Gray | | Not Tested |

Genetic Disorders

| | | | |
|---|--------|-------|--|
| X | HYPP | n/n | Clear: Negative for the HYPP gene mutation. |
| X | HERDA | N/N | Clear: Negative for the HERDA gene mutation. |
| X | GBED | N/N | Clear: Negative for the GBED gene mutation. |
| X | MH | n/n | Clear: Negative for the MH gene mutation found in Quarter horses and related breeds. |
| X | IMM | N/IMM | Both the normal and mutant alleles MYH1 gene were detected. Horse has a susceptibility to developing |
| X | PSSM 1 | n/n | Clear: Negative for the PSSM Type 1 gene mutation. |
| | FIS | | Not Tested |
| | JEB1 | | Not Tested |
| | JEB2 | | Not Tested |
| | CA | | Not Tested |
| | LFS | | Not Tested |
| | SCID | | Not Tested |
| | OAAM1 | | Not Tested |
| | WFFS1 | | Not Tested |

Additional Comments

None

Genetic Marker Results

Run Date: *Not Tested*

| | | | | | | |
|-------|-------|-------|-------|-------|------|---------|
| - | - | - | - | - | - | - |
| AHT4 | AHT5 | ASB17 | ASB2 | ASB23 | AME | CA425UK |
| - | - | - | - | - | - | - |
| HMS3 | HMS6 | HMS7 | HTG10 | HTG4 | LEX3 | LEX33 |
| - | - | - | - | - | - | - |
| VHL20 | UMD11 | HMS1 | HMS2 | HTG6 | HTG7 | |