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AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001		Case: QHA144171 Date Received: 16-May-2014 Print Date: 19-May-2014 Report ID: 6859-5225-7354-1160 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
Horse: SMART SAILING VIXEN YOB: Breed: QH Sex: M Alt. ID: 6545308		Reg: 5548461
Sire: TAMU NU BAR CODY Dam: FOXIE WILLA MAY		Reg: 4205232 Reg: 3211105

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 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.

Provided Information: Name: SMART SAILING VIXEN Registration: 5548461	Case: NQ64187 Date Received: 08-Jan-2021 Report Issue Date: 14-Jan-2021 Report ID: 3137-5204-7457-2075 Verify report at www.vgl.ucdavis.edu/verify
DOB: 04/21/2013 Sex: Mare Breed: Quarter Horse	
Sire: TAMU NU BAR CODY Reg: 4205232 Microchip:	Dam: FOXIE WILLA MAY Reg: 3211105 Microchip:

RESULT

INTERPRETATION

**Myosin-Heavy Chain
Myopathy (MYHM)**

N/My

1 copy of the MYHM mutation is present. Horse may develop IMM following infection or vaccination, or nonexertional rhabdomyolysis. Horse can pass on the mutation to 50% of offspring.