

Wednesday, February 14, 2018

8:22 AM UNIVERSITY OF CALIFORNIA, DAVIS

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VETERINARY GENETICS LABORATORY  
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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		<b>Date Received:</b> 12-May-2017 <b>Print Date:</b> 16-May-2017 <b>Report ID:</b> 6359-9692-9056-5084 <small>Verify report at <a href="http://www.vgl.ucdavis.edu/myvgl/verify.html">www.vgl.ucdavis.edu/myvgl/verify.html</a></small>
<b>Horse:</b> 00012233422		<b>Reg:</b>
<b>DOB:</b> 2017 <b>Sex:</b> Stallion <b>Breed:</b> Quarter Horse <b>An ID:</b> 6842661		
<b>Sire:</b> ATTENCION		<b>Reg:</b> 5176974
<b>Dam:</b> MISS COOL PERFECTION		<b>Reg:</b> 3911300
<b>GBED</b>	<b>N/N</b>	N/N - Normal - Does not possess the disease-causing GBED gene
<b>HERDA</b>	<b>N/N</b>	N/N - Normal - horse does not have the HERDA gene
<b>HYPP</b>	<b>N/N</b>	N/N - Normal - Does not possess the disease-causing HYPP gene
<b>MH</b>	<b>N/N</b>	N/N - Normal - horse does not have the MH gene
<b>PSSM1</b>	<b>N/N</b>	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 a recessive disease.

**HERDA** - Hereditary Lupine Regional Dermatitis. Skin disease characterized by depigmented skin, scuffing, and severe lesions along the back of affected horses. Typical on colts around 2 years of age. Inherited as a recessive disease.

**HYPP** - Type 1 Aleutian Periodic Paralysis. Muscle disease caused by defect in sodium channel gene that causes intermittent muscle weakness 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 that induce depolarizing muscle relaxants (succinylcholine) and stress. Presumed inheritance is dominant disease.

**PSSM1** - Polyunsaturated Storage Myopathy Type 1. Muscle disease characterized by accumulation of abnormal complex lipids 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.