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SANTA BARBARA • SANTA CRUZ

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

TELEPHONE: (530) 752-2211
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AQHA GENETIC DISEASE PANEL TEST RESULTS

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| AMERICAN QUARTER HORSE ASSOCIATION P.O. BOX 200 AMARILLO, TX 79168-0001 | | Case: QHA152777 Date Received: 04-Aug-2014 Print Date: 06-Aug-2014 Report ID: 5434-7261-8025-5071 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html |
| Horse: JESSES TOPAZ DOB: 01/01/2011 Breed: QH Sex: S Alt. ID: 6317408 | | Reg: 5374475 |
| Sire: MR JESS PERRY Dam: PADDYS TOPAZ | | Reg: 3145646 Reg: 4894615 |

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|-------|-----|---|
| 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.



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IMM AND MYH1 MYOPATHY (MYHM) GENETIC TEST REPORT

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| GLENN BLODGETT P.O. BOX 130 GUTHRIE, TX 79236 | Case: NQ51866 Date Received: 25-Jul-2019 Print Date: 29-Jul-2019 Report ID: 6900-0102-4312-6088 Verify report at www.vgl.ucdavis.edu/myvgl/verify.htm |
| Name: JESSES TOPAZ DOB: 01/01/2011 Sex: Stallion Breed: Quarter Horse | Reg: 5374475 |

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| IMM and MYH1 Myopathy | N/N |
|-----------------------|-----|

No copies of the MYHM mutation. Horse does not have increased susceptibility for IMM or nonexertional rhabdomyolysis.

For more detailed information on MYHM test results, please go to:
www.vgl.ucdavis.edu/services/horse/IMM.php