

## EQUINE DISEASE PANEL TEST REPORT

<b>Provided Information:</b> Name: <b>NONSTOP MILLIONAIRE</b> Registration: <b>5901222</b>	<b>Case: NQ107753</b> Date Received: 01-Apr-2024 Report Issue Date: 10-Apr-2024 Report ID: 3759-9953-0406-6039 <p style="text-align: right; font-size: small;">Verify report at <a href="http://vgl.ucdavis.edu/verify">vgl.ucdavis.edu/verify</a></p>
DOB: <b>03/28/2018</b> Sex: <b>Mare</b> Breed: <b>Quarter Horse</b>	
Sire: <b>TEJONS WHITE GOLD</b> Dam: <b>NONSTOP TO PARIS</b> Reg: <b>5617565</b> Reg: <b>4247320</b> Microchip: Microchip:	

### RESULT

### INTERPRETATION

Condition	Result	Interpretation
<b>Glycogen Branching Enzyme Deficiency (GBED)</b>	<b>N/N</b>	Normal. No copies of the GBED allele detected.
<b>Hereditary Equine Regional Dermal Asthenia (HERDA)</b>	<b>N/N</b>	Normal. No copies of the HERDA allele detected.
<b>Hyperkalemic Periodic Paralysis (HYPP)</b>	<b>N/N</b>	Normal. No copies of the HYPP allele detected.
<b>Myosin-Heavy Chain Myopathy (MYHM)</b>	<b>N/N</b>	Normal. No copies of the MYHM allele detected. Horse does not have increased susceptibility for immune mediated myositis or nonexertional rhabdomyolysis caused by the MYHM allele.
<b>Malignant Hyperthermia (MH)</b>	<b>N/N</b>	Normal. No copies of the MH allele detected.
<b>Polysaccharide Storage Myopathy Type 1 (PSSM1)</b>	<b>N/N</b>	Normal. No copies of the PSSM1 allele detected.