

## EQUINE DISEASE PANEL TEST REPORT

<b>Provided Information:</b>		<b>Case:</b> <b>NQ74973</b>
<b>Name:</b> METALATOR		<b>Date Received:</b> 15-Oct-2021
<b>Registration:</b> 6026626		<b>Report Issue Date:</b> 08-Nov-2021
		<b>Report ID:</b> 7370-6738-5346-9013
		Verify report at <a href="http://www.vgl.ucdavis.edu/verify">www.vgl.ucdavis.edu/verify</a>
<b>DOB:</b> 02/07/2020 <b>Sex:</b> Stallion <b>Breed:</b> Quarter Horse		

### RESULT

<b>Glycogen Branching Enzyme Deficiency (GBED)</b>	N/N
<b>Hereditary Equine Regional Dermal Asthenia (HERDA)</b>	N/HRD
<b>Hyperkalemic Periodic Paralysis (HYPP)</b>	N/N
<b>Myosin-Heavy Chain Myopathy (MYHM)</b>	N/N
<b>Malignant Hyperthermia (MH)</b>	N/N
<b>Polysaccharide Storage Myopathy Type 1 (PSSM1)</b>	N/N

### INTERPRETATION

Normal - Does not possess the disease-causing GBED gene

Carrier - horse carries one copy of the HERDA gene

Normal - Does not possess the disease-causing HYPP gene

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

Normal - horse does not have the MH gene

Normal - horse does not have the PSSM1 gene