

## DECISION TREE FOR DIAGNOSTIC TESTING FOR EXERTIONAL RHABDOMYOLYSIS

\*Disclosure of financial interest: Drs. McCue, Mickelson and Valberg are the patent owners for the genetic testing for *GYS1*. A portion of the proceeds from this test will go towards their continued research as well as patent royalties.

Based on our current understanding of PSSM and known genetic mutations we are currently recommending the following approach in the diagnostic process for PSSM:

**Signs of exertional rhabdomyolysis:** Muscle stiffness, cramping, pain, muscle fasciculation exercise intolerance, firm muscles, lameness, discolored brown urine (myoglobinuria), or progressive weakness, **and suspect PSSM**

