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HYPP REPORT

<p>KELLI BUTLER 9300 N LEWIS AVE SPERRY, OK 74073</p>	<p>Case: NQ40057 Date Received: 04-Jan-2018 Print Date: 05-Jan-2018 Report ID: 1494-1920-6333-1057 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html</p>
<p><i>Horse:</i> 17F PF JUST GRAND <i>Reg:</i> <i>DOB:</i> 02/01/2017 <i>Sex:</i> Stallion <i>Breed:</i> Quarter Horse</p>	
<p><i>Sire:</i> IEM THE ONE <i>Reg:</i> <i>Dam:</i> PF JUST GRAND <i>Reg:</i></p>	

HYPP Test Result

N/N

Result Codes:

- H/H Hyperkalemic - Homozygous for HYPP (two copies of the HYPP gene).
N/H Hyperkalemic - Heterozygous (one normal and one HYPP gene).
N/N Normal - Does not possess the disease-causing HYPP gene.

The disease is inherited as an autosomal dominant trait, which means that a heterozygote (N/H) bred to a normal (N/N) will result in approximately half of the offspring being affected and half being normal. The homozygote (H/H) is usually severely affected with the disease.

The test indicates the presence or absence of a base pair substitution in the skeletal muscle sodium channel gene. The abnormal gene codes for a defective sodium channel protein that causes the disease Hyperkalemic Periodic Paralysis (HYPP).