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HYPP REPORT

FAX: (530) 752-3556

MARY ANN ROHRBAUGH 2020 TALL FIR DR DOVER. PA 17315		Case: Date Received:	NQ44219 26-Jul-2018
		Print Date: Report ID:	30-Jul-2018 1617-0926-5859-1193
		Verify report at www.vgl.ucdavis.edu/myvgl/verify.htm	
Horse: VISTAR	<i>Reg:</i> 50507		
DOB: 05/09/2009 Sex: Mare Breed: Pony of the Americas			
Sire: BORN A STAR	<i>Reg:</i> 49421		
Dam: IMPRESSA RUE	<i>Reg:</i> B45536		

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).