# UNIVERSITY OF CALIFORNIA, DAVIS

BERKELEY • DAVIS • IRVINE • LOS ANGELES • MERCED • RIVERSIDE • SAN DIEGO • SAN FRANCISCO • SANTA BARBARA •

VETERINARY GENETICS LABORATORY
SCHOOL OF VETERINARY MEDICINE
ONE SHIELDS AVENUE
DAVIS, CALIFORNIA 95616-8744

TELEPHONE: (530) 752-2211
FAX: (530) 752-3556

## AQHA GENETIC DISEASE PANEL TEST RESULTS

AMERICAN QUARTER HORSE ASSOCIATION
P.O. BOX 200
AMARILLO, TX 79168-0001

*Case:* QHA3424
*Date Received:* 20-Feb-201
*Print Date:* 27-Feb-2018
*Report ID:* 4896-8411-4129-
Verify report at www.vgl.ucdavis.edu/myvgl/v

*Horse:* ***MAGICAL VOODOO*** *Reg:* ***5702218***

*YOB:* ***2014*** *Sex:* ***Stallion*** *Breed:* ***Quarter Horse*** *Alt. ID:* ***6658218***

*Sire:* ***SHINERS VOODOO DR*** *Reg:* 5101811

*Dam:* ***VERY SMART SWEETHART*** *Reg:* 5165985

| | | |
|---|---|---|
| **GBED** | **N/N** | N/N - Normal - Does not possess the disease-causing GBED gene |
| **HERDA** | **N/HRD** | N/HRD - Carrier - horse carries one copy of the HERDA gene |
| **HYPP** | **N/N** | N/N - Normal - Does not possess the disease-causing HYPP gene |
| **MH** | **N/N** | N/N - Normal - horse does not have the MH gene |
| **PSSM1** | **N/N** | N/N - Normal - horse does not have the PSSM1 gene |

GBED - Glycogen Branching Enzyme Deficiency. Fatal disease of newborn foals caused by defect in glycogen storage. Affects heart and skeletal muscles and brain. Inh
disease.

HERDA - Hereditary Equine Regional Dermal Asthenia. Skin disease characterized by hyperextensible skin, scarring, and severe lesions along the back of affected hors
around 2 years of age. Inherited as a recessive disease.

HYPP - Hyperkalemic Periodic Paralysis. Muscle disease caused by defect in sodium channel gene that causes involuntary muscle contraction and increased level of po
Inherited as dominant disease. Two copies of defective gene produce more severe signs than one copy.

MH - Malignant Hyperthermia. Rare but life-threatening skeletal muscle disease triggered by exposure to volatile anesthetics (halothane), depolarizing muscle relaxant
and stress. Presumed inheritance as dominant disease.

PSSM1 - Polysaccharide Storage Myopathy Type 1. Muscle disease characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include mus
twitching, sweating, weakness and reluctance to move. Inherited as a dominant disease.

GBED testing performed under a license agreement with the University of Minnesota.
HERDA testing performed under a license agreement with the University of California, Davis.
PSSM1 testing performed under a license agreement with the American Quarter Horse Association.

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*Dam:* ***VERY SMART SWEETHART*** *Reg:* 5165985

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| **HYPP** | **N/N** | N/N - Normal - Does not possess the disease-causing HYPP gene |
| **MH** | **N/N** | N/N - Normal - horse does not have the MH gene |
| **PSSM1** | **N/N** | N/N - Normal - horse does not have the PSSM1 gene |

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PSSM1 testing performed under a license agreement with the American Quarter Horse Association.

*Name:* **MAGICAL VOODOO**

*Registration:* **584383**

*Date Recei*
*Report Issu*
*Report ID:*

*DOB:* **01/01/2014** *Sex:* **Stallion** *Breed:* **Quarter Horse**

*Sire:* SHINERS VOODOO DR
*Reg:* 5101811
*Microchip:*

*Dam:* VERY SMART SV
*Reg:* 5165985
*Microchip:*

| RESULT | | INTERPRETATION | RESULT | |
|---|---|---|---|---|
| RED FACTOR | **e/e** | Only red factor detected. Basic color is red in the absence of modifying genes. | SPLASHED WHITE | |
| AGOUTI | **A/a** | 1 copy of agouti. If present, black pigment is restricted to the points. | TOBIANO | |
| CREAM | **N/Cr** | 1 copy of Cream dilution detected. | LEOPARD | |
| PEARL | **N/N** | No copies of Pearl dilution detected. | PATTERN-1 | |
| SILVER | **N/N** | No copies of Silver dilution detected. | BRINDLE 1 | |
| DUN | **nd2/nd2** | Horse is not Dun dilute. Primitive markings are absent. | TIGER EYE | |
| CHAMPAGNE | **N/N** | No copies of Champagne dilution detected. | MUSHROOM (SHETLAND PONY) | |
| LETHAL WHITE OVERO | | Not requested. | GRAY | **Absent** |
| SABINO 1 | | Not requested. | ROAN | |
| DOMINANT WHITE (W5, W10, W20, W22) | | Not requested. | | |

UCDAVIS
VETERINARY MEDICINE
Veterinary Genetics Laboratory

# MYOSIN-HEAVY CHAIN MYOPATHY (MYHM) TEST REPORT

***Provided Information:***

*Name:* **MAGICAL VOODOO**

*Registration:* **584383**

***Case:*** **NQ86118**
***Date Received:*** 29-Aug-2022
***Report Issue Date:*** 18-Apr-2023
***Report ID:*** 2541-5288-2437-7162

Verify report at www.vgl.ucdavis.edu/verify

*DOB:* **01/01/2014** *Sex:* **Stallion** *Breed:* **Quarter Horse**

*Sire:* SHINERS VOODOO DR
*Reg:* 5101811
*Microchip:*

*Dam:* VERY SMART SWEETHART
*Reg:* 5165985
*Microchip:*

## RESULT

| | |
|---|---|
| **Myosin-Heavy Chain Myopathy (MYHM)** | **N/N** |

## INTERPRETATION

Normal. No copies of the MYHM allele detected. Horse does not have increased susceptibility for immune m
myositis or nonexertional rhabdomyolysis caused by the MYHM allele.