## UNIVERSITY OF CALIFORNIA. DAVIS

BENKELEY + DAVIS + BRYINE + LOS ANGELES + MERCED + RIVERSIDE + SAN DIEGO + SAN FRANCISCO

SANTA BARBARA . SANTA CRUZ

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

TELEPHONE. (\$30) 752-2211 PAX: (530) 752-3556

## AOHA GENETIC DISEASE PANEL TEST RESULTS

**AMERICAN QUARTER HORSE ASSOCIATION** P.O. BOX 200

AMARILLO, TX 79168-0001

QHA47405 Case:

28-Feb-2012 Date Received:

Report Date: Report ID:

06-Mar-2012 6634-4360-8022-9055 Verify report at https://www.vgl.ucdarla.eduknyvgl/verify.html

Horse: FOUR SIX HANCOCK

Reg: 3812306

YOB: 97

Breed: QH

Sex: S

Alt. ID: 4149490

Sire: TANQUERY GIN Dam: HANCOCK CINCO Reg: 1090327 Reg: 0867986

GBED	NN	N/N - No
HERDA	N/N	N/N - No
НҮРР	N/N	N/N - No
МН	N/N	N/N - No
PSSM1	N/N	N/N - Na

ormal - Does not possess the disease-causing GBED gene

orumi - harve does not have the HERDA gene

ormal - Does not possess the disease-causing HYPP gene

ormal - horse does not have the MH gene

arnal - horse does not have the PSSMI gene

GBED - Olycogen Branching Enzyme Deficiency. Fatal disease of newborn fouls caused by defect in glycogen storage. Affects heart and skeletal muscles and brain. Inherited as recessive disease

HERDA - Hereditary Equiae Regional Dermal Authenia. Skin disease characterized by hyperentensible skin, scarring, and severe lesions along the back of affected horses. Typical equet is around 2 years of age. Inherited as a recessive disease.

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

MH - Malignant Hyperthermia. Rase but life-threatening skeletal muscle disease triggered by exponent to volatile anesthetics (haloftame), depolarizing muscle relaxants (succinylcholine), and stress. Presumed inheritance as dominant disease.

PSBM1 - Polysaocharide Storage Myopathy Type 1. Muscle direase characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include councie pain, stiffners, akin twitching, aweating, weakness and reluctance to move. Inherited as a dominant disease.

GBED testing performed under a license agreement with the University of Minascota. 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.