

AQHA

AMERICAN QUARTER HORSE ASSOCIATION

TO SPEED PROCESSING,
PLEASE RETURN THIS FORM
WITH ADDITIONAL INFOR-
MATION AT YOUR EARLIEST
CONVENIENCE.

February 16, 2015

Reg Key: 4445872

BRIAN & LISA FULTON
29393 272ND AVE
VALENTINE, NE 69201-2172

Cust ID: 2080256

RE: A STREAK OF FLING
3806003

Dear MEMBER:

The results for the disease panel test that you ordered for A STREAK OF FLING are enclosed.

The DNA profile for this panel was compared to the original DNA. For more information regarding these diseases, please refer to AQHA rules REG109.3-REG109.7.

If you have any questions, please contact us at (806) 376-4811. Our office hours are Monday through Friday, 8 a.m. to 5 p.m. Central time.

Sincerely,

Registration Department

Enclosure

HYP302

P.O. Box 200 ■ Amarillo, Texas ■ 79168 ■ 1600 Quarter Horse Drive ■ Amarillo, Texas ■ 79104
(806) 376-4811



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: QHA176446 Date Received: 26-Jan-2015 Print Date: 29-Jan-2015 Report ID: 9044-0502-1017-3073 Verify report at www.vgl.ucdavis.edu/myvgl/verify.html
Horse: A STREAK OF FLING YOB: 1999 Breed: QH Sex: S Alt. ID: 4445872 Reg: 3806003	
Sire: STREAKIN SIX Dam: MOON FLING	Reg: 1301847 Reg: 1678394

GBED	N/N
HERDA	N/N
HYPP	N/N
MH	N/N
PSSMI	N/N

N/N - Normal - Does not possess the disease-causing GBED gene
 N/N - Normal - horse does not have the HERDA gene
 N/N - Normal - Does not possess the disease-causing HYPP gene
 N/N - Normal - horse does not have the MH gene
 N/N - Normal - horse does not have the PSSMI gene

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

HERDA - Hereditary Equine Regional Dermal Asthenia. Skin disease characterized by hyperextensible skin, scarring, and severe lesions along the back of affected horses. Typical onset is 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 potassium in blood. 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 relaxants (succinylcholine), and stress. Presumed inheritance as dominant disease.

PSSMI - Polysaccharide Storage Myopathy Type 1. Muscle disease characterized by accumulation of abnormal complex sugars in skeletal muscles. Signs include muscle pain, stiffness, skin 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.
 PSSMI testing performed under a license agreement with the American Quarter Horse Association.