

### REGISTERED NAME FROZEN KINDA

REGISTRATION NUMBER 5164742



DATE ISSUED NOVEMBER 07, 2016 OWNER NAME BIZET MELANIE

MELANIE BIZET DE CHEMIN MOULIN BUCKET 1 1495 SART-DAMES AVELINES BELGIUM





MARKINGS LEFT HIND PASTERN WHITE. DARK SPOTS ON LEFT HIND CORONET. NO OTHER MARKINGS.

The name on the front of this certificate listed as CURRENT OWNER is the present owner of this horse as shown on the records of American Quarter Horse Association. If ownership changes have occurred, up to three previous owners are listed below. All other ownership records are on file in the AQHA office.

2/14/13MAIDA SILVIA7/15/10DORAZIO DANIELA2/28/09ANGELUCCI Q H

00123 ROMA, ITALY 00133 ROME, ITALY 00052 CERVETERI, ITALY

(Physical Address) 1600 Quarter Horse Drive Amarillo, TX 79104 Telephone: (806)376-4811

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(Mailing Address) P.O.Box 200 Amarillo, Texas 79168

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Reg: 5164742



SANTA BARBARA 🔹 SANTA CRUZ

# AQHA GENETIC DISEASE PANEL TEST RESULTS

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

Case: Date Received: Print Date: Report 1D:

## QHA271536 31-Oct-2016

 Date:
 04-Nov-2016

 ort ID:
 4237-6145-3064-1020

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

Horse: FROZEN KINDA

YOB: 2007 Sex: Mare Breed: Quarter Horse Alt. 1D: 6073687

<i>Sire:</i> FROZEN SAILOR <i>Dam:</i> SHES MY KINDA CHIC		Reg: 3406970 Reg: 3574385	
GBED	N/N	N/N - Normal - Does not possess the disease-causing GBED gene	
HERDA	N/N	N/N - Normal - horse does not have the HERDA gene	
НҮРР	N/N	N/N - Normal - Does not possess the disease-causing HYPP gene	
МН	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. 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.

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