



VETERINARY GENETICS LABORATORY
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CEREBELLAR ABIOTROPHY TEST REPORT

SASCHA QUADERER BANGARTEN 23 9490 VADUZ SWITZERLAND	<i>Case:</i> CBA312 <i>Date Received:</i> 16-Mar-2009 <i>Report Date:</i> 19-Mar-2009 <i>Report ID:</i> 5698-3353-9138-5037
<i>Name:</i> MARJAN ALBIDAYER <i>YOB:</i> 08 <i>Breed:</i> AR <i>Sex:</i> S	<i>Reg:</i> DE408082007408008 <i>Alt. ID:</i>
<i>Sire:</i> MARAJJ <i>Dam:</i> PUSTYNNA DROGA	<i>Reg:</i> 812158604 <i>Reg:</i>

CEREBELLAR ABIOTROPHY RESULT

N/N

Result Codes:

N/N	Normal: horse does not possess markers associated with CA
N/CA	Carrier (1 copy of the CA gene): horse has markers associated with CA and is considered to be phenotypically normal and a carrier of the disease gene
CA/CA	Affected (2 copies of the CA gene): horse has markers associated with CA and is considered to have the disease
N/U	CA status cannot be determined. Atypical combination of DNA markers. Possibility of CA-carrier status cannot be excluded with confidence.

Cerebellar Abiotrophy (CA) is a neurological genetic disease found in Arabian horses that is characterized by head tremor and lack of balance equilibrium. CA is inherited as a recessive trait and thus breedings between two carriers (N/CA) have a 25% chance of producing an affected foal. Breedings between carrier and normal (N/N) horses produce only normal foals, but 50% of these are expected to be carriers. While CA is not fatal, affected animals can be a danger to themselves and their handlers because of their symptoms.

This is not a direct genetic test for CA. The analysis is based on markers associated with this disease.