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Test Report

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**Analysis of the predisposition to dwarfism
in Miniature Horses (ACAN D1, D2, D3 improved, D4)**

Name: Grace N'Roses des Monts d'Auvergne
Lab-No.: 201806727 Identification: A225940
Gender: female Date of birth: 2016-03-12

Test	Inheritance	Result
ACAN	recessive	Normal (N/N)

Remark: The analysis was performed by a partner laboratory.
In literature 4 mutations in the ACAN gene were described which are responsible for dwarfism in Miniature Horses. The genotypes D1/D1, D1/D2, D1/D3 improved and D1/D4 are lethal. Therefore, breeding with D1 carriers should be avoided.

Legend: **Normal:** The animal does not carry the mutation. When used in breeding this animal will not spread the predisposition into the population.

Carrier: The animal is heterozygous for the predisposition. When used in breeding on average 50% of the offspring will inherit the mutated allele. Carriers (D1, D2, D3 improved, D4) of the predisposition will not show dwarfism-like symptoms.

Affected: The animal has two mutated alleles and will be affected by disproportionate dwarfism. When used in breeding all offspring will inherit a mutated allele.

Lethal: Affected animals with a D1 genotype (D1/D1, D1/D2, D1/D3 improved or D1/D4) are not viable.

Unknown: The genotypes D3 improved /D3 improved and D4/D4 have not been detected in any animal yet. Therefore, the effect of these mutations remains unclear.

Comment: -

Rheinbach, 2018-07-30

Please note:

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Provided Information:

Name: **GRACE N ROSES DES MONTS D'AUVERGNE**
 Registration: **A225940**

Case: **NQ58945**
 Date Received: 30-Jun-2020
 Report Issue Date: 10-Jul-2020
 Report ID: 9088-8354-6302-2095

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

YOB: Sex: **Mare** Breed: **Miniature Horse**

Skeletal Atavism Result

N/N



Interpretation

N/N	Normal. No copies of the skeletal atavism mutations detected.
N/Del1	Carrier. 1 copy of a skeletal atavism mutation detected.
N/Del2	Carrier. 1 copy of a skeletal atavism mutation detected.
Del1/Del1	Affected. 2 copies of mutation detected.
Del2/Del2	Affected. 2 copies of mutation detected.
Del1/Del2	Affected. 2 copies of mutation detected.