

## Laboratory Report

<b>Laboratory #:</b>	162472	<b>Call Name:</b>	Molly
<b>Order #:</b>	72079	<b>Registered Name:</b>	MW Rising Star
<b>Ordered By:</b>	Darah Gestes	<b>Breed:</b>	Bernedoodle
<b>Ordered:</b>	Jan. 15, 2020	<b>Sex:</b>	Female
<b>Received:</b>	Feb. 6, 2020	<b>DOB:</b>	Jan. 2019
<b>Reported:</b>	Feb. 19, 2020	<b>Registration #:</b>	-
		<b>Microchip #:</b>	932002000584926

### Results:

Disease	Gene	Genotype	Interpretation
Degenerative Myelopathy	<i>SOD1</i>	WT/WT	Normal (clear)
Degenerative Myelopathy (Bernese Mountain Dog Type)	<i>SOD1</i>	WT/M	Carrier
GM2 Gangliosidosis (Poodle Type)	<i>HEXB</i>	WT/WT	Normal (clear)
Neonatal Encephalopathy with Seizures	<i>ATF2</i>	WT/WT	Normal (clear)
Osteochondrodysplasia	<i>SLC13A1</i>	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration	<i>PRCD</i>	WT/WT	Normal (clear)
Von Willebrand Disease I	<i>VWF</i>	WT/WT	Normal (clear)

WT, wild type (normal); M, mutant; Y, Y chromosome (male)

### Interpretation:

Molecular genetic analysis was performed for seven specific mutations reported to be associated with disease in dogs. We identified two normal copies of the DNA sequences in six of the mutations tested. Thus, this dog is not at an increased risk for the diseases associated with these six mutations. However, we identified one normal copy and one mutant copy of the DNA sequences for *SOD1*. Thus, this dog is a carrier of Degenerative Myelopathy (Bernese Mountain Dog Type).

### Recommendations:

Degenerative Myelopathy (Bernese Mountain Dog Type) is inherited in an autosomal recessive fashion. Based on this, and the fact that this dog showed a mutation in one copy of the *SOD1* gene, this dog is a carrier of this disease. Although dogs that carry only one copy of this mutation will not be clinically affected, if bred with another carrier, the pairing could produce affected offspring. To avoid producing affected offspring, this dog should be bred with dogs that are normal (WT/WT) for this gene. Dogs related to this dog have an increased risk to be affected by or carry the mutated gene. Additional testing for this mutation is indicated for related dogs.

Paw Print Genetics® has genetic counseling available to you at no additional charge to answer any questions about these test results, their implications and potential outcomes in breeding this dog.



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**Helen F Smith, PhD**  
Assistant Laboratory Director



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**Casey R Carl, DVM**  
Associate Medical Director

Normal results do not exclude inherited mutations not tested in these or other genes that may cause medical problems or may be passed on to offspring. These tests were developed and their performance determined by Paw Print Genetics<sup>®</sup>. This laboratory has established and verified the tests' accuracy and precision. Because all tests performed are DNA-based, rare genomic variations may interfere with the performance of some tests producing false results. If you think these results are in error, please contact the laboratory immediately for further evaluation. In the event of a valid dispute of results claim, Paw Print Genetics will do its best to resolve such a claim to the customer's satisfaction. If no resolution is possible after investigation by Paw Print Genetics with the cooperation of the customer, the extent of the customer's sole remedy is a refund of the fee paid. In no event shall Paw Print Genetics be liable for indirect, consequential or incidental damages of any kind. Any claim must be asserted within 60 days of the report of the test results.

# Orthopedic Foundation for Animals Preliminary (Consultation) Report



A Not-For-Profit  
Organization

MIDWEST'S RISING STAR  
registered name

HYBRID  
breed

932002000584926  
tattoo/microchip/DNA profile

2123095  
application number

film/case no(s)

NOREG2123095  
registration number

F  
sex

1/1/2019  
date of birth

13  
age at evaluation in months

2/10/2020  
date of report

Owner  
DARAH GESTES  
470 BARKWOOD LN  
GLEN CARBON, IL 62034

Veterinarian  
VETERINARY GROUP OF CHESTERFIELD  
17709 EDISON AVE STE A  
CHESTERFIELD, MO 63005

## RADIOGRAPHIC EVALUATION OF PELVIC PHENOTYPE WITH RESPECT TO HIP DYSPLASIA

\* The study must be repeated when the animal is 24 months of age or older to qualify for an OFA number.

\_\_\_\_\_ **EXCELLENT HIP JOINT CONFORMATION\***  
superior hip joint conformation as compared with other  
individuals of the same breed and age

✓ \_\_\_\_\_ **GOOD HIP JOINT CONFORMATION\***  
well formed hip joint conformation as compared with other  
individuals of the same breed and age

\_\_\_\_\_ **FAIR HIP JOINT CONFORMATION\***  
minor irregularities of the hip joint conformation as compared  
with other individuals of the same breed and age

\_\_\_\_\_ **BORDERLINE HIP JOINT CONFORMATION**  
marginal hip joint conformation of indeterminate status with  
respect to hip dysplasia at this time – **Repeat study in six  
months**

\_\_\_\_\_ **MILD HIP DYSPLASIA**  
radiographic evidence of minor dysplastic changes of the hip  
joints

\_\_\_\_\_ **MODERATE HIP DYSPLASIA**  
well defined radiographic evidence of dysplastic changes of  
the hip joints

\_\_\_\_\_ **SEVERE HIP DYSPLASIA**  
radiographic evidence of marked dysplastic changes of the  
hip joints

### HIP JOINTS - STANDARD VD VIEW RADIOGRAPHIC FINDINGS

- \_\_\_\_\_ subluxation
- \_\_\_\_\_ remodeling of femoral head/neck
- \_\_\_\_\_ osteoarthritis/degenerative joint disease
- \_\_\_\_\_ shallow acetabula
- \_\_\_\_\_ acetabular rim/edge change
- \_\_\_\_\_ unilateral pathology \_\_\_\_\_ left \_\_\_\_\_ right
- \_\_\_\_\_ transitional vertebra
- \_\_\_\_\_ spondylosis
- \_\_\_\_\_ panosteitis
- \_\_\_\_\_ other

### ELBOW JOINTS – FLEXED LATERAL VIEW

✓ \_\_\_\_\_ negative for elbow dysplasia    ✓ \_\_\_\_\_ L    ✓ \_\_\_\_\_ R

### ELBOW DYSPLASIA

Grade I    L \_\_\_\_\_ R \_\_\_\_\_  
Grade II    L \_\_\_\_\_ R \_\_\_\_\_  
Grade III    L \_\_\_\_\_ R \_\_\_\_\_

### RADIOGRAPHIC FINDINGS

degenerative joint disease (DJD)    L \_\_\_\_\_ R \_\_\_\_\_  
ununited anconeal process (UAP)    L \_\_\_\_\_ R \_\_\_\_\_  
fragmented coronoid process (FCP)    L \_\_\_\_\_ R \_\_\_\_\_  
osteochondrosis    L \_\_\_\_\_ R \_\_\_\_\_

Consultation by: Greg Keller DVM  
G.G. KELLER/DVM, MS, DACVR  
CHIEF OF VETERINARY SERVICES

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## Coat Color and Trait Certificate

<b>Call Name:</b>	Molly	<b>Laboratory #:</b>	162472
<b>Registered Name:</b>	MW Rising Star	<b>Registration #:</b>	-
<b>Breed:</b>	Bernedoodle	<b>Microchip #:</b>	932002000584926
<b>Sex:</b>	Female	<b>Certificate Date:</b>	Feb. 19, 2020
<b>DOB:</b>	Jan. 2019		

**This canine's DNA showed the following genotype(s):**

Coat Color/Trait Test	Gene	Genotype	Interpretation
Cu Locus (Curly Hair)	<i>KRT71</i>	Cu/Cu <sup>C</sup>	Wavy/Curly coat (carrier)

**Interpretation:**

This dog carries one copy of **Cu<sup>C</sup>** and one copy of **Cu** which results in a wavy or curly coat. However, the overall coat type of this dog is dependent on the combination of this dog's genotypes at the L, Cu, and IC loci. This dog will pass **Cu<sup>C</sup>** on to 50% of its offspring and **Cu** to 50% of its offspring.

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