

# AMERICAN KENNEL CLUB

NAME

HAPPY TAILS WESLEY

NUMBER

PR27325002

BREED

POODLE

SEX

MALE

COLOR

RED & WHITE

DATE OF BIRTH

JULY 14, 2023

SIRE

QC GEYSER'S GIBSON

PR26137005 11-23 (AKC DNA #V10056687)

DAM

QC CREAM CAROLINE

PR25599909 11-23

BREEDER

IVAN D WEAVER

OWNER

MATTHEW YODER

4460 TOWNSHIP ROAD 617

MILLERSBURG OH 44654-9188



AMERICAN  
KENNEL CLUB®

CERTIFICATE ISSUED  
MARCH 15, 2024

*This certificate invalidates all previous certificates issued.*

If a date appears after the name and number of the sire and dam, it indicates the issue of the Stud Book Register in which the sire or dam is published.

**For Transfer Instructions, see back of Certificate.**

*This Certificate issued with the right to correct or revoke by the American Kennel Club.*

## REGISTRATION CERTIFICATE

990000009934753 - Wesley



## Demographic Information

AT/AT e/e B/B K4/K4 SP/SP CLEAR

Call Name	Comet	DOB	July 14, 2023
Registered Name	-	Registration #	-
Breed	Poodle	Tattoo	-
Sex	Male	Microchip	4753
Owner	Ivan Weaver	Laboratory #	417244
		Report Date	Sept. 15, 2023

These tests were developed and performed by Paw Print Genetics®, Spokane WA.

## Explanation of Results

- Normal** A 'Normal' result means that your dog does not have the mutation that causes the associated genetic disease.
- Carrier** A 'Carrier' result indicates that your dog has inherited one copy of the mutation that has been reported to cause this genetic disease. Your dog may not be clinically affected by this mutation because two copies of the mutation are usually required to cause disease.
- Carrier / At-Risk** A 'Carrier / At-Risk' result indicates that your dog inherited one copy of the mutation that has been reported to cause this genetic disease. Based on the mode of genetic inheritance for this particular disease, inheriting one mutant copy of the gene may result in the disease. Dogs with one copy of the mutation may have a milder phenotype as compared to cats with two copies of this mutation.
- At-Risk / Affected** An 'At-Risk / Affected' result indicates that your dog inherited one or two copies of the mutation that has been reported to cause this genetic disease. Based on the mode of genetic inheritance for this particular disease, inheriting one or two mutant copies of the gene may result in the disease.

## No Result

'No Result' indicates that we were unable to obtain a genotype for your dog for this specific disease or trait and does not mean that your dog is a carrier or at-risk for this disease. There are a variety of reasons why a specific test may not provide a reportable result. Unique variations in the genetic code of some individuals may exist and cause certain regions of the genome to not perform properly with a specific test. In addition, suboptimal sampling of the dog's cheek cells could also result in poor sample performance due to inadequate cell counts, bacterial and fungal growth, or the presence of other test inhibitors. An acceptable level of tests with no results has been determined by Paw Print Genetics. Dogs with at least 90% of the test results are determined to be acceptable and reportable. If your dog has an unacceptable level of tests with no results, you will be contacted for a new sample to repeat the testing.

Please review our testing terms and disclaimers regarding your results.

WT: **wild type (normal)** M: **mutant** Y: **Y chromosome (male)**

## Breed Profile

Disease Name	Genotype	Interpretation
Congenital Methemoglobinemia	WT/WT	Normal (Clear)
Degenerative Myelopathy	WT/WT	Normal (Clear)
Degenerative Myelopathy (Bernese Mountain Dog Variant)	0	
Degenerative Myelopathy (Common Variant)	0	
Ehlers-Danlos Syndrome	WT/WT	Normal (Clear)
Ehlers-Danlos Syndrome (Variant 1)	0	
Ehlers-Danlos Syndrome (Variant 2)	0	
GM2 Gangliosidosis (Poodle Type)	WT/WT	Normal (Clear)
Hereditary Cataracts	WT/WT	Normal (Clear)
Intervertebral Disc Disease Risk Factor and Chondrodystrophy (CDDY with IVDD)	WT/WT	Normal (Clear)
Multidrug Resistance 1	WT/WT	Normal (Clear)
Neonatal Encephalopathy with Seizures	WT/WT	Normal (Clear)
Osteochondrodysplasia	WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration (prcd)	WT/WT	Normal (Clear)
Progressive Retinal Atrophy, Rod-Cone Dysplasia 4	WT/WT	Normal (Clear)
Von Willebrand Disease I	WT/WT	Normal (Clear)

WT: **wild type (normal)** M: **mutant** Y: **Y chromosome (male)**

**Submitted By**Happy Tail Pets, LLC  
Happy Tail Pets LLC**Subject Dog**Dog Name: **4753\_Wesley**  
Breed: **Poodle**  
Phenotype: **Red & White**  
Sex: **Male**  
Birth: **Jul 14, 2023**Lab Reference #: **840009**  
Microchip: **990000009934753****Disorder Results (6 of 16)**

CDPA	<b>N/N</b>	Clear: Dog is negative for the CDPA mutation.
CDDY	<b>N/N</b>	Clear: Dog is negative for the mutation associated with CDDY.
DM	<b>n/n</b>	Clear: Dog is negative for mutation associated with Degenerative Myelopathy.
NEwS	<b>n/n</b>	Clear: Dog is negative for mutation associated with NEwS.
PRA-prcd	<b>n/n</b>	Negative: Dog is negative for the mutation associated with prcd-PRA.
vWD1	<b>n/n</b>	Clear: Dog is negative for the mutation associated with von Willebrand's Disease Type I.

**Color Results (5 of 16)**

A-Locus	<b>at/at</b>	Dog has two copies of the gene causing tan points.
B-Locus	<b>B/B</b>	Dog does not carry the mutation for most forms of chocolate coloration.
D-Locus	<b>D/D</b>	Negative: Dog is negative for the mutation associated with a diluted coat color.
E-Locus	<b>e/e</b>	Dog has two copies of cream/yellow.
K-Locus	<b>n/n</b>	Dog is negative for the KB allele, and the coat coloration will be based on the agouti genotype.

**Pattern Results (1 of 16)**

S-Locus	<b>S/S</b>	Homozygous: Dog has two copies of S-Locus resulting in a nearly solid white, parti, or piebald coat color.
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**Trait Results (4 of 16)**

Curl 1&2	<b>C<sup>1</sup>/C<sup>1</sup></b>	The dog has two copies of the hair curl allele. The dog will have curly hair, and will always pass on a copy of the hair curl allele to any offspring. All offspring of this dog will have curly hair.
Furnishings	<b>F/F</b>	Furnished: Dog has two copies of the furnishings mutation and will always produce offspring with a furnished coat.
Hair Length (1-5)	<b>l<sup>1</sup>/l<sup>1</sup></b>	Two copies of the long-hair allele, dog will have longer than average hair per the breed standard.
Shedding	<b>n/n</b>	Dog has no copies of the shedding allele. The dog will have a low propensity towards shedding.



# "WESLEY"

## HAPPY TAIL'S WESLEY



DNA Test Report

Test Date: July 22nd, 2024

[embk.me/happytailswesley](http://embk.me/happytailswesley)

### BREED ANCESTRY



### GENETIC STATS

Predicted adult weight: **22 lbs**

Life stage: **Young adult**

Based on your dog's date of birth provided.

### TEST DETAILS

Kit number: EM-55950226

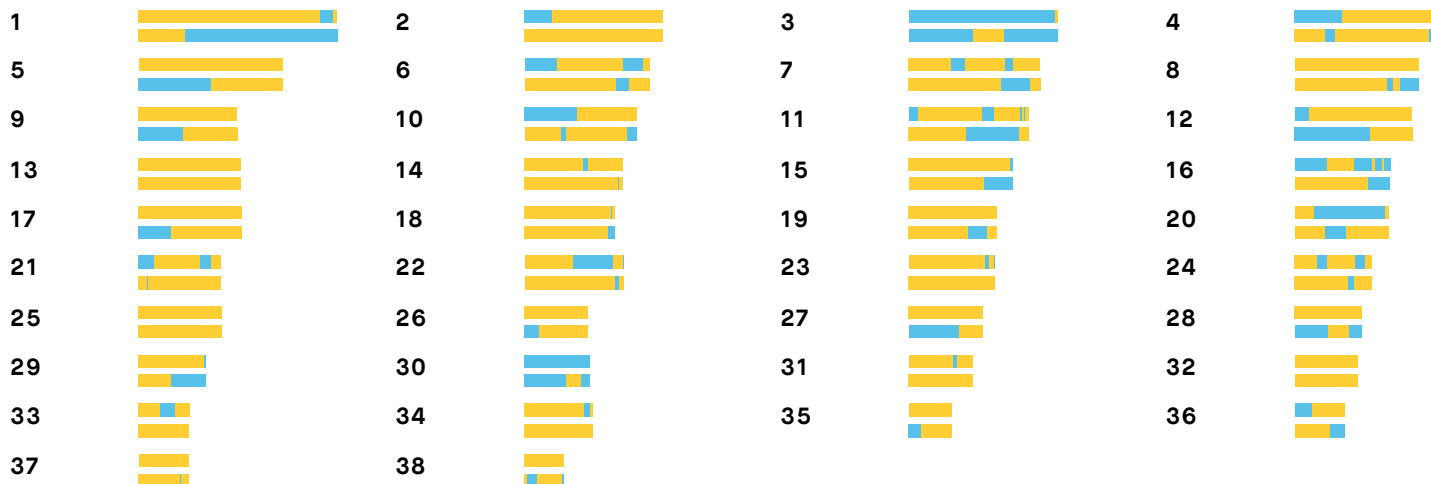
Swab number: 31220612405627

### BREED ANCESTRY BY CHROMOSOME

Our advanced test identifies from where Wesley inherited every part of the chromosome pairs in his genome.

Breed colors:

Poodle (Small) Poodle (Standard)





# "WESLEY"

## HAPPY TAIL'S WESLEY



DNA Test Report

Test Date: July 22nd, 2024

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### BREED-RELEVANT RESULTS

Research studies indicate that these results are more relevant to dogs like Wesley, and may influence his chances of developing certain health conditions.

<input checked="" type="checkbox"/> Degenerative Myelopathy, DM (SOD1A)	Clear
<input checked="" type="checkbox"/> GM2 Gangliosidosis (HEXB, Poodle Variant)	Clear
<input checked="" type="checkbox"/> Intervertebral Disc Disease (Type I) (FGF4 retrogene - CFA12)	Clear
<input checked="" type="checkbox"/> Neonatal Encephalopathy with Seizures, NEWS (ATF2)	Clear
<input checked="" type="checkbox"/> Osteochondrodysplasia (SLC13A1, Poodle Variant)	Clear
<input checked="" type="checkbox"/> Progressive Retinal Atrophy, prcd (PRCD Exon 1)	Clear
<input checked="" type="checkbox"/> Von Willebrand Disease Type I, Type I vWD (VWF)	Clear

Registration: American Kennel Club (AKC)

PR27325002

