

## **Canine Genetic Health Certificate™**

Call Name: Wyatt

**Registered Name:** I'll Be Your Huckleberry of Out West

LS-N

**Breed:** Australian Shepherd

Sex: Male DOB: Dec. 2020

**Laboratory #:** 261854

**Registration #:** DN64671909 / E220997

Certificate Date: Nov. 17, 2021

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

Disease	Gene	Genotype	Interpretation
Coagulation Factor VII Deficiency	F7	WT/WT	Normal (clear)
Collie Eye Anomaly	NHEJ1	WT/WT	Normal (clear)
Cone Degeneration	CNGB3	WT/WT	Normal (clear)
Craniomandibular Osteopathy	SLC37A2	WT/WT	Normal (clear)
Degenerative Myelopathy	SOD1	WT/M	Carrier
Exercise-Induced Collapse	DNM1	WT/WT	Normal (clear)
Hereditary Cataracts (Australian Shepherd Type)	HSF4	WT/WT	Normal (clear)
Hyperuricosuria	SLC2A9	WT/WT	Normal (clear)
Intestinal Cobalamin Malabsorption (Australian Shepherd Type)	AMN	WT/WT	Normal (clear)
Intestinal Cobalamin Malabsorption (Border Collie Type)	CUBN	WT/WT	Normal (clear)
Multidrug Resistance 1	ABCB1	WT/M	Carrier (At-Risk)
Multifocal Retinopathy 1	BEST1	WT/WT	Normal (clear)

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

Blake C Ballif, PhD

Laboratory & Scientific Director

I han Cally

Casey R Carl, DVM

Associate Medical Director

Paw Print Genetics<sup>®</sup> performed the tests listed on this dog. See the Laboratory Report for interpretation and recommendations based on these findings. The genes/diseases reported here were selected by the client. 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. 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. Genetic counseling is available at Paw Print Genetics.



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Disease	Gene	Genotype	Interpretation
Neuronal Ceroid Lipofuscinosis 6	CLN6	WT/WT	Normal (clear)
Neuronal Ceroid Lipofuscinosis 8 (Australian Shepherd Type)	CLN8	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration	PRCD	WT/WT	Normal (clear)
Von Willebrand Disease I	VWF	WT/WT	Normal (clear)

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

NOTE: The following fields were adjusted at the client's request on Nov 17, 2021: Registration ID, Registered Name

Sho (Sally)

Blake C Ballif, PhD

Laboratory & Scientific Director

Ex RCI

Casey R Carl, DVM Associate Medical Director

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