

AMERICAN KENNEL CLUB • FOUNDED 1884

# Certified Pedigree

**Sire** GCHS CH PICCADILLY'S A SHADE DARKER  
SR95048501 (03-18) OFA24G OFEL24 BLK  
AKC DNA #V841100

**CH THE SUN ALWAYS SHINES**  
SS11545503 OFA27G OFEL24  
LABRADOR RETRIEVER MALE YLW  
Date Whelped: 03/12/2019  
Breeder: SHEILA NORGREN

**Dam** CH EAST HILL POPPY  
SR92183406 (07-19) OFA24G OFEL24 BLK



AMERICAN  
KENNEL CLUB®

*Genia Di Nardo*  
Executive Secretary

The Seal of The American Kennel Club affixed hereto certifies that this pedigree was compiled from official Stud Book records on November 16, 2021.

GCHB CH PICCADILLY'S SEEK SHADE DS  
SR70870303 (02-14) OFA24G OFEL24 BLK AKC  
DNA #V706017

ERINHILL'S SECRET RENDEZVOUS AT  
BAYSKOR  
SR76126903 (04-16) OFA24G OFEL24 BLK

NORTHWOODS HEARTBREAKER OF BANNER  
SR65387602 (06-12) OFA25G OFEL25 BLK AKC  
DNA #V674194

MARSH RIDGE MAD DASH TO EASTHILL  
SR79892804 (06-16) OFA24G OFEL24 BLK

CH ENDEAVOR PICCADILLY GOTTA WEAR  
SHADES JH NA NAJ  
SR40600101 (06-09) OFA24G OFEL24 BLK  
AKC DNA #V601280

PRETON ANITA AT WINDFALL  
SR58676801 (04-12) OFA55G OFEL55 BLK  
(BRA) AKC DNA #V649739

CH FORTUNE'S CURIOUS GEORGE  
SR29330802 (03-08) OFA24E OFEL24 BLK  
AKC DNA #V501469

CH VICTORIA'S SECRET AT ERINHILL  
SR54803401 (04-13) OFA27G OFEL27 BLK

CH LOBUFF BOBWHITE AT  
CHUCKLEBROOK  
SN91419803 (07-03) OFA24G OFEL24 YLW  
AKC DNA #V266976

BANNERS SHEZAN ANGEL NORTHWOOD  
SR41372102 (03-11) BLK

GCHS CH SUNNYDAZE RUNNING ACROSS  
THE MILES JH  
SR60224403 (03-12) OFA25G OFEL25 YLW  
AKC DNA #V643060

EASTHILLS SPLASH-N-DASH  
SR59489202 (09-12) OFA26G OFEL26 BLK

# THE AMERICAN KENNEL CLUB

## CHAMPIONSHIP CERTIFICATE

*This certifies that*

RETRIEVER (LABRADOR)  
THE SUN ALWAYS SHINES ~ SS11545503

*bred by*

SHEILA NORGREN

*owned by*

SHEILA NORGREN

*having completed the requirements on*

AUGUST 1, 2021

*has been officially recorded a*

**CHAMPION**

*by The American Kennel Club*



AMERICAN  
KENNEL CLUB®

*Gina DiNardo*  
Executive Secretary

**THE SUN ALWAYS SHINES**

SS11545503

M YELLOW LABRADOR RETRIEVER

Born Mar 12 2019

Sire: PICCADILLY'S A SHADE DARKER SR95048501

Dam: EAST HILL POPPY SR92183406

Registry	Test Date	Report Date	Age (m)	Conclusion	OFA Number
ADVANCED CARDIAC	Feb 20 2021	Mar 18 2021	23	NORMAL AUSC+ECHO	LR-ACA2648/23M-VPI
ELBOW	Mar 17 2021	Mar 29 2021	24	NORMAL	LR-EL 102053M24-VPI
HIPS	Jul 7 2021	Jul 19 2021	27	GOOD	LR-256352G27M-VPI

2/23/22, 11:44 AM

Animal Details

**THE SUN ALWAYS SHINES**

Registration:	SS11545503 (AKC)	Sire:	SR95048501	 <p><a href="#">Add a photo to your dog's record.</a></p> <p><a href="#">Click here to learn more.</a></p>
Breed:	LABRADOR RETRIEVER	Dam:	SR92183406	
Sex:	M	Titles:		
Color:	YELLOW	CHIC #:	N/A	
Birthdate:	Mar 12 2019	Addtl. Reg. #		
DNA Profile:				

TEST RESULTS					
OFA Number	Registry	Test Date	Report Date	Age (mos)	Final Conclusion
LR-ACA2648/23M-VPI	ADVANCED CARDIAC	Feb 20 2021	Mar 18 2021	23	NORMAL AO/CONG, AUSC/ECHO
LR-EL 102053M24-VPI	ELBOW	Mar 17 2021	Mar 29 2021	24	NORMAL
LR-256352G27M-VPI	HIPS	Jul 7 2021	Jul 19 2021	27	GOOD

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## Canine Genetic Health Certificate™

**Call Name:** RA  
**Registered Name:** The Sun Always Shines  
**Breed:** Labrador Retriever  
**Sex:** Male  
**DOB:** March 2019

**Laboratory #:** 153300  
**Registration #:** -  
**Certificate Date:** March 11, 2021

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

Disease	Gene	Genotype	Interpretation
Centronuclear Myopathy	PTPLA	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome (Labrador Retriever Type)	COLQ	WT/WT	Normal (clear)
Exercise-Induced Collapse	DNM1	WT/M	Carrier
Hereditary Nasal Parakeratosis	SUV39H2	WT/WT	Normal (clear)
Macular Corneal Dystrophy (Labrador Retriever Type)	CHST6	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Golden Retriever 2	TTC8	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration	PRCD	WT/WT	Normal (clear)
Retinal Dysplasia/Oculoskeletal Dysplasia 1	COL9A3	WT/WT	Normal (clear)
Skeletal Dysplasia 2	COL11A2	WT/WT	Normal (clear)
Stargardt Disease	ABCA4	WT/M	Carrier

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

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

**Christina J Ramirez, PhD, DVM, DACVP**  
Medical Director

Paw Print Genetics® 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.



## Canine Genetic Health Certificate™

**Call Name:** RA  
**Registered Name:** CH The Sun Always Shines  
**Breed:** Labrador Retriever  
**Sex:** Male  
**DOB:** March 2019

**Laboratory #:** 376901  
**Registration #:** SS11545503  
**Certificate Date:** Jan. 20, 2023

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

Disease	Gene	Genotype	Interpretation
Copper Toxicosis (Labrador Retriever Type) ATP7A	ATP7A	M/Y	Carrier Male
Copper Toxicosis (Labrador Retriever Type) ATP7B	ATP7B	WT/WT	Normal (clear)

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

**Blake C Ballif, PhD**  
Laboratory & Scientific Director

**Christina J Ramirez, PhD, DVM, DACVP**  
Medical Director

Paw Print Genetics® performed the testing on the dog listed on this certificate. 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. The results included in this report relate only to the items tested using the sample provided. These tests were developed and their performance determined by Paw Print Genetics. This laboratory has established and verified the test(s) accuracy and precision with >99.9% sensitivity and specificity. The presence of mosaicism may not be detected by this test. Non-paternity may lead to unexpected results. This is not a breed identification test. 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, 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 any consequential or punitive damages.



## Coat Color and Trait Certificate

**Call Name:** RA  
**Registered Name:** The Sun Always Shines  
**Breed:** Labrador Retriever  
**Sex:** Male  
**DOB:** March 2019

**Laboratory #:** 153300  
**Registration #:** -  
**Certificate Date:** Feb. 17, 2020

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

Coat Color/Trait Test	Gene	Genotype	Interpretation
L Locus (Long Hair/Fluffy)	FGF5	Sh/Lh	Shorthaired (carries longhaired)

### Interpretation:

This dog carries one copy of **Sh** and one copy of **Lh** which results in short hair. 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 **Sh** on to 50% of its offspring and **Lh** to 50% of its offspring.

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.

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

**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®. 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.





## Canine Genetic Health Certificate™

**Call Name:** RA  
**Registered Name:** CH The Sun Always Shines  
**Breed:** Labrador Retriever  
**Sex:** Male  
**DOB:** March 2019

**Laboratory #:** 376901  
**Registration #:** SS11545503  
**Certificate Date:** Jan. 20, 2023

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

Disease	Gene	Genotype	Interpretation
Copper Toxicosis (Labrador Retriever Type) ATP7A	ATP7A	M/Y	Carrier Male
Copper Toxicosis (Labrador Retriever Type) ATP7B	ATP7B	WT/WT	Normal (clear)

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

**Blake C Ballif, PhD**  
Laboratory & Scientific Director

**Christina J Ramirez, PhD, DVM, DACVP**  
Medical Director

Paw Print Genetics® performed the testing on the dog listed on this certificate. 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. The results included in this report relate only to the items tested using the sample provided. These tests were developed and their performance determined by Paw Print Genetics. This laboratory has established and verified the test(s) accuracy and precision with >99.9% sensitivity and specificity. The presence of mosaicism may not be detected by this test. Non-paternity may lead to unexpected results. This is not a breed identification test. 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 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 any consequential or punitive damages.



# Orthopedic Foundation for Animals

2300 E Nifong Blvd, Columbia, MO 65201-3806

Phone: (573) 442-0418; Fax: (573) 875-5073

www.ofa.org. A not-for-profit organization

Registered name:  
SUN Always shines  
Breed: Labrador Sex: M

ID Number (if any): ☐ Tattoo ☐ Microchip

Registration Number: ☐ ARC ☐ Other

Date of Birth (mm/dd/yy): Date of Exam (mm/dd/yy):

Owner Name: Sheila Norgren

Co-Owner Name: Phone:

Owner Address: 900 Gully Rd

City: Howell State: MI Zip/postal code: 48873

E-Mail (use both lines if needed):

I hereby certify that the animal examined is the animal described on this application, and understand that the results of this exam will be submitted by the examining ophthalmologist to the database for statistical gathering purposes. I understand that only passing results will be released to the public unless the initials of a registered owner or authorized agent appear in the authorization box below which permits the OFA to release non-passing results to the public.

Sheila Norgren  
Signature of owner or authorized agent/representative

I hereby authorize the OFA to release the results of the evaluation of the animal described on this application to the public if the results are non-passing (initials)

- ☐ I DID verify microchip/tattoo on this dog  
☒ I DID NOT verify microchip/tattoo on this dog  
☐ NO MICROCHIP / TATTOO PRESENT

I certify that I have performed this ophthalmic examination using pharmacological mydriasis, ophthalmoscopy, and biomicroscopy.

Signature: [Signature] ACVO # EC596 Date 11/6/19

Diplomate, American College of Veterinary Ophthalmologists

FEES AND CREDIT CARD INFORMATION ON THE BACK OF THE WHITE (OWNER) COPY



506712

## Companion Animal Eye Registry (CAER)

RIGHT EYE	GLOBE	LEFT EYE
<input type="checkbox"/>	microphthalmos	<input type="checkbox"/>
<input type="checkbox"/>	keratoconjunctivitis sicca	<input type="checkbox"/>
<input type="checkbox"/>	glaucoma	<input type="checkbox"/>
<b>EYELIDS</b>		
<input type="checkbox"/>	entropion	<input type="checkbox"/>
<input type="checkbox"/>	ectropion	<input type="checkbox"/>
<input type="checkbox"/>	distichiasis	<input type="checkbox"/>
<input type="checkbox"/>	ectopic cilia	<input type="checkbox"/>
<input type="checkbox"/>	imperforate lacrimal punctum	<input type="checkbox"/>
<b>NICTITANS</b>		
<input type="checkbox"/>	cartilage anomaly/eversion	<input type="checkbox"/>
<input type="checkbox"/>	gland prolapse	<input type="checkbox"/>
<input type="checkbox"/>	plasmoma/atypical pannus	<input type="checkbox"/>
<b>CORNEA</b>		
<input type="checkbox"/>	dystrophy — epithelial/stromal	<input type="checkbox"/>
<input type="checkbox"/>	dystrophy — endothelial	<input type="checkbox"/>
<input type="checkbox"/>	pannus	<input type="checkbox"/>
<input type="checkbox"/>	pigmentary keratitis/keratopathy	<input type="checkbox"/>
<b>UVEA</b>		
<input type="checkbox"/>	uveal cyst	<input type="checkbox"/>
<input type="checkbox"/>	iris coloboma	<input type="checkbox"/>
<input type="checkbox"/>	iris hypoplasia	<input type="checkbox"/>
<input type="checkbox"/>	iris sphincter dysplasia	<input type="checkbox"/>
<input type="checkbox"/>	pigmentary uveitis	<input type="checkbox"/>
<input type="checkbox"/>	uveal melanoma	<input type="checkbox"/>
<b>peristent pupillary membranes</b>		
<b>LENS</b>		
<input type="checkbox"/>	anterior cortex	<input type="checkbox"/>
<input type="checkbox"/>	posterior cortex	<input type="checkbox"/>
<input type="checkbox"/>	equatorial cortex	<input type="checkbox"/>
<input type="checkbox"/>	anterior sutures	<input type="checkbox"/>
<input type="checkbox"/>	posterior sutures	<input type="checkbox"/>
<input type="checkbox"/>	nucleus	<input type="checkbox"/>
<input type="checkbox"/>	capsular	<input type="checkbox"/>
<input type="checkbox"/>	generalized/complete	<input type="checkbox"/>
<input type="checkbox"/>	resorbing/hypermature	<input type="checkbox"/>
<b>suspect not inherited</b>		
<input type="checkbox"/>	subluxation/luxation	<input type="checkbox"/>
<b>VITREOUS</b>		
<input type="checkbox"/>	PHPV/PHTVL	<input type="checkbox"/>
<input type="checkbox"/>	persistent hyaloid artery	<input type="checkbox"/>
<input type="checkbox"/>	degeneration	<input type="checkbox"/>

Ophthalmologist Name: **Dr. Michael West EC596**  
Ophthalmologist Address: **BluePearl Veterinary Partners**  
**Southfield, MI**  
City: State: Zip/postal code:  
Phone: **248-354-6660**  
Email:

RIGHT EYE	FUNDUS	LEFT EYE
<input type="checkbox"/>	retinal detachment	<input type="checkbox"/>
<input type="checkbox"/>	retinal atrophy—generalized	<input type="checkbox"/>
<input type="checkbox"/>	retinopathy	<input type="checkbox"/>
<input type="checkbox"/>	retinal dysplasia	<input type="checkbox"/>
<input type="checkbox"/>	choroidal hypoplasia	<input type="checkbox"/>
<input type="checkbox"/>	coloboma	<input type="checkbox"/>
<input type="checkbox"/>	optic nerve coloboma	<input type="checkbox"/>
<input type="checkbox"/>	optic nerve hypoplasia	<input type="checkbox"/>
<input type="checkbox"/>	micropapilla	<input type="checkbox"/>

**OTHER CONDITIONS**  
☐ Unlisted conditions suspected as **inherited**. Describe in comments  
☐ Unlisted conditions suspected as **not inherited**

**NORMAL**

Comments


WHITE = Owner/OFA Registration copy; PINK = ACVO Diplomate copy; YELLOW = ACVO Research copy

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## Laboratory Report

**Laboratory #:** 153300  
**Order #:** 101846  
**Ordered By:** Sheila Norgren  
**Ordered:** March 1, 2021  
**Received:** March 2, 2021  
**Reported:** March 11, 2021

**Call Name:** RA  
**Registered Name:** The Sun Always Shines  
**Breed:** Labrador Retriever  
**Sex:** Male  
**DOB:** March 2019  
**Registration #:** -

### Results:

Disease	Gene	Genotype	Interpretation
Centronuclear Myopathy	PTPLA	WT/WT	Normal (clear)
Congenital Myasthenic Syndrome (Labrador Retriever Type)	COLQ	WT/WT	Normal (clear)
Macular Corneal Dystrophy (Labrador Retriever Type)	CHST6	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Golden Retriever 2	TTC8	WT/WT	Normal (clear)
Progressive Retinal Atrophy, Progressive Rod-Cone Degeneration	PRCD	WT/WT	Normal (clear)
Retinal Dysplasia/Oculoskeletal Dysplasia 1	COL9A3	WT/WT	Normal (clear)
Skeletal Dysplasia 2	COL11A2	WT/WT	Normal (clear)
Stargardt Disease	ABCA4	WT/M	Carrier

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

### Interpretation:

Molecular genetic analysis was performed for eight specific mutations reported to be associated with disease in dogs. We identified two normal copies of the DNA sequences in seven of the mutations tested. Thus, this dog is not at an increased risk for the diseases associated with these seven mutations. However, we identified one normal copy and one mutant copy of the DNA sequences for *ABCA4*. Thus, this dog is a carrier of Stargardt Disease.

### Recommendations:

Stargardt Disease is inherited in an autosomal recessive fashion. Based on this, and the fact that this dog showed a mutation in one copy of the *ABCA4* 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.