### ORTHOPEDIC FOUNDATION FOR ANIMALS, INC.

BLACKFORK'S CHASING AFTER YOU registered name

LABRADOR RETRIEVER

breed

film/test/lab #

900113001672055 tattoo/microchip/DNA profile

2261877 application number

06/30/2021 date of report

RESULTS:

Based upon the radiograph submitted, the consensus was that no evidence of elbow dysplasia was recognized.

**NORMAL** 

SS12185303

registration no.

05/13/2019

O.F.A. NUMBER

age at evaluation in months

LR-EL104231F25-VPI

This number issued with the right to correct or revoke by the Orthopedic Foundation for Animals.

F

sex

wnel



G.G.KELLER. D.V.M., M.S., DACVR CHIEF OF VETERINARY SERVICES

A Not-For-Profit Organization

www.ofa.org

This electronic OFA certificate was generated on: 06/30/2021

This certification can be verified on the OFA website by entering the dog's registration number into the orange search box located at the top of the page or by scanning the QR code above.

If there are any errors on this certificate, please email CORRECTIONS@OFFA.ORG to request a correction.

Orthopedic Foundation for Animals, Inc. 2300 E. Nifong Blvd. Columbia, MO 65201-3806

OFA website: www.ofa.org E-mail address: ofa@offa.org Phone number: 573-442-0418 Fax number: 573-875-5073

### ORTHOPEDIC FOUNDATION FOR ANIMALS, INC.

BLACKFORK'S CHASING AFTER YOU registered name

LABRADOR RETRIEVER

film/test/lab #

900113001672055 tattoo/microchip/DNA profile

2261877 application number

06/30/2021 date of report

RESULTS:

Based upon the radiograph submitted, the consensus was that no evidence of hip dysplasia was recognized. The hip joint conformation was evaluated as:

wner



F sex

05/13/2019

25

age at evaluation in months



LR-255948E25F-VPI

O.F.A. NUMBER

This number issued with the right to correct or revoke by the Orthopedic Foundation for Animals.

**EXCELLENT** 

OFA eCert



Verify certificate with QR scan

www.ofa.org

G.G.KELLER. D.V.M., M.S., DACVR CHIEF OF VETERINARY SERVICES

This electronic OFA certificate was generated on: 06/30/2021

大大大大大大大大大大大大大大大大大大大大大大大大大大大大大大大大**大** 

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#### ORTHOPEDIC FOUNDATION FOR ANIMALS, INC.

BLACKFORK'S CHASING AFTER YOU

registered name

LABRADOR RETRIEVER

sex/breed

700151 film/test/lab #

900113001672055 tattoo/microchip/DNA profile

2261877 application number

03/24/2022 date of report

RESULTS:

Based upon the exam dated 06/15/2021, this dog has been found to be free of observable inherited eye disease and has been issued an Eye Certification Registry Number which is valid for one year from the time of the exam.

**NORMAL** 

OFA eCert



www.ofa.org

SS12185303 registration no.

05/13/2019 date of birth

age at evaluation in months



A Not-For-Profit Organization

LR-EYE24815/25F-VPI

O.F.A. NUMBER

This number issued with the right to correct or revoke by the Orthopedic Foundation for Animals.

> G.G.KELLER. D.V.M., M.S., DACVR CHIEF OF VETERINARY SERVICES

This electronic OFA certificate was generated on: 03/24/2022

This certification can be verified on the OFA website by entering the dog's registration number into the orange search box located at the top of the page or by scanning the QR code above.

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OFA website: www.ofa.org E-mail address: ofa@offa.org Phone number: 573-442-0418 Fax number: 573-875-5073



# **DNA Test Report**

Dog Information

Blackfork's Chasing After You (Chloe)

NAM

Labrador Retriever

GENETIC BREED

AKC: SS12185303 REGISTRATION

Female SEX

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DATE OF BIRTH

n/a MICROCHIP Audry Steelman
OWNER NAME

Canine Genetic Health Screen

TEST

July 26th, 2021 TEST DATE

## **BREED HEALTH TESTS**

Centronuclear Myopathy       PTPLA       NN       Clear       ②         Degenerative Myelopathy, DM       SOD1(A)       GG       Clear       ②         Exercise-Induced Collapse       DNM1       GG       Clear       ②         Hereditary Nasal Parakeratosis, HNPK       SUV39H2       GG       Clear       ②         Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU       SLC2A9 (Exon 5)       GG       Clear       ②         Macular Corneal Dystrophy, MCD       CHST6       CC       Clear       ②         Narcolepsy       HCRTR2       AA       Clear       ②         Progressive Retinal Atrophy, prcd       PRCD Exon 1       CC       Clear       ②         Pyruvate Kinase Deficiency       PKLR Exon 7 SNP Variant 1       CC       Clear       ②         Skeletal Dysplasia 2, SD2       COL11A2       GG       Clear       ②         Achromatopsia       CNGA3 (Exon 7 Deletion)       NN       Clear       ☆         Alexander Disease       GFAP (Exon 4)       GG       Clear       ☆         Congenital Myasthenic Syndrome, CMS       COLQ (Exon 14)       TT       Clear       ☆         Colden Retriever Progressive Retinal Atrophy 2, GR-PRA2       TTC8 Exon 8       NN       NN	DISEASE	GENE	GENOTYPE	RESULT	TESTING RECOMMENDED BY
Exercise-Induced Collapse  Hereditary Nasal Parakeratosis, HNPK  SUV39H2  GG  Clear  Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU  SLC2A9 (Exon 5)  GG  Clear  Macular Corneal Dystrophy, MCD  CHSTG  CC  Clear  Narcolepsy  HCRTR2  AA  Clear  Progressive Retinal Atrophy, prcd  PRCD Exon 1  GG  Clear  Skeletal Dysplasia 2, SD2  COL11A2  GG  Clear  COL11A2  GG  Clear  Alexander Disease  GFAP (Exon 4)  GG  Clear  COL1ear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear  Colar  Colar	Centronuclear Myopathy	PTPLA	NN	Clear	<b>(</b>
Hereditary Nasal Parakeratosis, HNPK  SUV39H2  GG  Clear  Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU  SLC2A9 (Exon 5)  GG  Clear  Macular Corneal Dystrophy, MCD  CHST6  CC  Clear  Progressive Retinal Atrophy, prcd  PRCD Exon 1  GG  Clear  Pryuvate Kinase Deficiency  PKLR Exon 7 SNP Variant 1  CC  Clear  COL11A2  GG  Clear  Achromatopsia  CNGA3 (Exon 7 Deletion)  NN  Clear  Alexander Disease  GFAP (Exon 4)  GG  Clear  CC  Clear  COL11A2  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear  TT  Clear	Degenerative Myelopathy, DM	SOD1(A)	GG	Clear	•
Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU  SLC2A9 (Exon 5)  GG  Clear  Macular Corneal Dystrophy, MCD  CHST6  CC  Clear  Marcolepsy  HCRTR2  AA  Clear  Progressive Retinal Atrophy, prcd  PRCD Exon 1  GG  Clear  Pyruvate Kinase Deficiency  PKLR Exon 7 SNP Variant 1  CC  Clear  Skeletal Dysplasia 2, SD2  COL11A2  GG  Clear  CNGA3 (Exon 7 Deletion)  NN  Clear  Adexander Disease  GFAP (Exon 4)  GG  Clear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear	Exercise-Induced Collapse	DNM1	GG	Clear	•
Macular Corneal Dystrophy, MCD  CHST6  CC  Clear  Narcolepsy  HCRTR2  AA  Clear  Progressive Retinal Atrophy, prcd  PRCD Exon 1  CC  Clear  Skeletal Dysplasia 2, SD2  Achromatopsia  CNGA3 (Exon 7 Deletion)  NN  Clear  Alexander Disease  GFAP (Exon 4)  GG  Clear  COLiar  Congenital Myasthenic Syndrome, CMS  CHST6  CHST7  CHST7  CHEAT  CC  Clear  CC  CC  CLear  CC  CLear  CC  CLear  CC  CC  CC  CLear  CC  CC  CC  CC  CC  CC  CC  CC  CC	Hereditary Nasal Parakeratosis, HNPK	SUV39H2	GG	Clear	•
Narcolepsy  HCRTR2  AA  Clear  Progressive Retinal Atrophy, prcd  PRCD Exon 1  GG  Clear  Cle	Hyperuricosuria and Hyperuricemia or Urolithiasis, HUU	SLC2A9 (Exon 5)	GG	Clear	•
Progressive Retinal Atrophy, prcd  PRCD Exon 1  GG  Clear  Pyruvate Kinase Deficiency  PKLR Exon 7 SNP Variant 1  CC  Clear  COL11A2  GG  Clear  COL21A2  Achromatopsia  CNGA3 (Exon 7 Deletion)  NN  Clear  Alexander Disease  GFAP (Exon 4)  GG  Clear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear	Macular Corneal Dystrophy, MCD	CHST6	CC	Clear	•
Pyruvate Kinase Deficiency  PKLR Exon 7 SNP Variant 1  CC  Clear  Skeletal Dysplasia 2, SD2  COL11A2  GG  Clear  Achromatopsia  CNGA3 (Exon 7 Deletion)  NN  Clear  Alexander Disease  GFAP (Exon 4)  GG  Clear  Conine Elliptocytosis  SPTB Exon 30  CC  Clear  Clear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear	Narcolepsy	HCRTR2	AA	Clear	•
Skeletal Dysplasia 2, SD2  COL11A2  GG Clear  CNGA3 (Exon 7 Deletion)  NN Clear  Alexander Disease  GFAP (Exon 4)  GG Clear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT Clear	Progressive Retinal Atrophy, prcd	PRCD Exon 1	GG	Clear	•
Achromatopsia CNGA3 (Exon 7 Deletion) NN Clear  Alexander Disease GFAP (Exon 4) GG Clear  Canine Elliptocytosis SPTB Exon 30 CC Clear  Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Pyruvate Kinase Deficiency	PKLR Exon 7 SNP Variant 1	CC	Clear	•
Alexander Disease GFAP (Exon 4) GG Clear Canine Elliptocytosis SPTB Exon 30 CC Clear Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Skeletal Dysplasia 2, SD2	COL11A2	GG	Clear	•
Canine Elliptocytosis  SPTB Exon 30  CC  Clear  Congenital Myasthenic Syndrome, CMS  COLQ (Exon 14)  TT  Clear	Achromatopsia	CNGA3 (Exon 7 Deletion)	NN	Clear	×
Congenital Myasthenic Syndrome, CMS COLQ (Exon 14) TT Clear	Alexander Disease	GFAP (Exon 4)	GG	Clear	×
	Canine Elliptocytosis	SPTB Exon 30	СС	Clear	x
Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2 TTC8 Exon 8 NN Clear	Congenital Myasthenic Syndrome, CMS	COLQ (Exon 14)	TT	Clear	x
	Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2	TTC8 Exon 8	NN	Clear	x



# **DNA Test Report**

Dog Information

Blackfork's Chasing After You (Chloe)

Female

**Labrador Retriever** 

**GENETIC BREED** 

AKC: SS12185303

n/a

REGISTRATION

MICROCHIP

DATE OF BIRTH

**Audry Steelman** OWNER NAME

**Canine Genetic Health Screen** 

TEST

July 26th, 2021

TEST DATE

## **BREED HEALTH TESTS**

DISEASE	GENE	GENOTYPE	RESULT	TESTING RECOMMENDED BY
Myotubular Myopathy 1, X-linked Myotubular Myopathy, XL-MTM	MTM1 (Exon 7)	cc	Clear	ਮ
Progressive Retinal Atrophy, crd4/cord1	RPGRIP1 (Exon 2)	NN	Clear	x
Ulrich-like Congenital Muscular Dystrophy	COL6A3	GG	Clear	낡