Name: phillip hutchison Sex: M (DOB: 2020-09-19) Address: Breed: Golden Retriever Microchip: KC registration: KC name: AGLABS GOLDEN GOOSE Results date: Jul 12, 2022 Date Received: Jun 27, 2022 The DNA for this testing was collected by the owner/breeder. Mutation Disorder COL7A1 (c.5716G>A) Dystrophic epidermolysis bullosa PKP1 (c.202+1G>C) Ectodermal dysplasia SLC4A3 (c.2601\_2602insC) Golden Retriever PRA1 PNPLA1 (c.1445\_1447delACGInsTACTACTA) Ichthyosis (Golden Retriever) DMD () Muscular Dystrophy CLN5 (c.934\_935delAG) Neuronal Ceroid Lipofuscinosis 5

Guide to results

Sensory Ataxic Neuropathy

Osteogenesis Imperfecta

Progressive Retinal Atrophy (GR-PRA2)

Progressive Retinal Atrophy (PRA-PRCD)

Clear means we have tested for the mutation known to cause the disorder and have not found it. goose will not develop CLEAR disorder and does not carry it.

> Carrier means we have tested for the mutation known to cause the disorder and have found one copy of it. goose will develop the disorder but can pass the mutation onto their offspring. If you choose to breed from this animal, care mus taken to ensure that any potential mate has tested CLEAR for the mutation goose carries, otherwise approximately 25° puppies will be affected with the disorder.

If your animal has any "at risk" mutations it means that our geneticists have determined that goose is likely to develop disorder. You should read the information about the disorder carefully, along with the notes left by our geneticists. advisable that you show this report to your vet during your next routine visit, unless otherwise indicated by the geneticist

COL1A1 (c.1276G>C)

tRNA-Tyr (m.5305delA)

TTC8 (c.669delA)

PRCD (c.5G>A)

Animal Genetics

Generated on: 26/01/23 Genetic Testing Report AGLABS GOLDEN GOOSE OF FALLOWFEN

Result

CLEAR

CLEAR

CLEAR

CLEAR

CLEAR

CLEAR

CLEAR

Owned By

Phillip Hutchison

AT RISK

Name: AGLABS GOLDEN GOOSE OF FALLOWFEN Lab Reference #: 650535

Breed: Golden Retriever

Microchip:

phenotype: Sex: Male Birth: Sep 19, 2020

Disorder Results (2 of 2)

n/n Ich (GR) n/n Ich-2

Clear: Dog is negative for the mutation associated with Ichthyosis. Dog is clear of the mutation associated with Ichthyosis 2.

British Veterinary Association/Kennel Club/International Sheep Dog Society (BVA/KC/ISDS)  CANINE HEALTH SCHEMES EYE EXAMINATION CENTURE ASSOCIATION OF THE PROPERTY OF THE PRO
CANINE HEALTH SCHEMES EYE EXAMINATION CERTIFICATE  KC no.
GOOSE KC no.  AGUARS GOLDEN GOOSE  Microchip no  Date of previous examination  Sex M F Date of birth 19-9-30
. 1 20
Owner's email address
out address O16.56 812.133 Vet's email address
with the used where gonioscopy is required.  In special to facilitate a complete examination of the eye and that a local execution of the eye and
be the dod submitted and persons and may be published. Any appeal against the results seclided below must be made to the BVA (for details see EPWP1).  Is necessary to facilitate a complete examination of the eye and that a local anaesthetic search and information provided in this form will be used to administer the eye examination service and will be retained for 7 years for accounting purposes on an electronic search and the provided in the search and
Date 47 - Z.7
Comea Drainage Angle Iris Lens Vitreous Fundus
Indirect Biomicroscopy Gogloscopy Tonometry Other  Comea Drainage Angle Iris Lens Vitreous Fundus  LEFT Comments NO BREED RELATED ADNEXAL OR OCULAR CONDITIONS
Z196 PLA BILATERALLY
L. IRISTIRIS
BILLAT - ALSTICHIASIS
FUNDUS
DNA sample taken on this date: Yes No Little confirm that the scanned microchip number
matches the number on the certificate Information for owners/Appeals leaflet (EPWP1) issued
INHERITED EYE DISEASE STATUS
The section applies to the known inherited ocular conditions specified in the Procedure Notes. These results will be sent to the KC and/or ISDS as appropriate.  CLINICALLY CLINICALLY CLINICALLY NON-CONGENITAL UNAFFECTED AFFECTED AFFECTED.
CONGENITALINEURALIA (HC) Hereditary cataract
Citie eye shormaly  Ordinary lens luxation  (PAG) Primary lens luxation  (PAG) Primary open angle glaucoma  (IOP) Intracoular pressure R mmHg L mmHg
Meliodi retinal dysplasia  (PRA) Progressive retinal atrophy (RPED) Retinal pigment epithelial dystrophy
TICL) Total refinal dysplantal predictory cataract DCI congestal hereditary cataract CCI congestal hereditar
- IM 2 3 Result
0 = normal, 1 = mility alreaded, 2
Consider sector with coular conditions not currently specified in the Procedure Notes.  Posterior Cortical Cataract Posterior Cortical Cataract Posterior Polar Subcapsular Cataract
Ocular Melanosis Pectinate ligament abnormality PhPV  Other conditions (specify) C PhPV
Lens luxation Anterior Capsular Cataract Anterior Cordical Cataract
Mithodia defects  Committed deposition  Perinuclear Cataract  Nuclear Cataract  Nuclear Cataract  Nuclear Cataract
2 Course Schemewith the results as shown 1, 1–7-2
(a) (A A)
Name Panellist Scale of Panellist Scale of Panellist Scale of PLA Testing, which is valid for 3 years  This certificate is valid for 12 months from date of signature with the exception of PLA Testing, which is valid for 3 years  BVA telephone 020 7908 6380 SVA 01/20
District White Access College Day Dink - retained by panellist
British Veterinary Association/Kennel Club/International Sheep Dog Society (BVA/KC/ISDS)  CANINE HEALTH SCHEMES EYE EXAMINATION CERTIFICATE
CANINE HEALTH SCHLINES ET E
Pet name GCOSE KC no.  KC registered name AGCASS GOLDEN GLOSE OF PANOWED Date of previous examination 19-9-20
Pet name GCOSE KC no.  KC registered name AGLAGS GOLDEN CLOSE OF PANOWERS Date of previous examination 19-9.20  Breed Retroace (GOLDEN) Colour GOLD Sex M F Date of birth 19-9.20  Description pages and address MR * MRS HUTCHINSON,
Pet name GCOSE KC no.  KC registered name AGCASS GOLDEN GLOSE OF PANOWED Date of previous examination 19-9-20

I understand and agree that the	will be used where gonioscopy is required.  ormation provided in this form will be used to admir	hister the eye examination service and will elevant information relating to CHS service	to estained for 7 years for accounting purposes on an electronic
system. My personal information	may be used from time at the things of the t	m301	Date 21.7.23.
Signature of Owner/Agent			
/	EXAMINATION	ON OF THE EYE AND ADNEXA	
Aydriatic Ophthalmoscopy I	Direct Indirect Biomicroscopy	Gonioscopy Tonometry O	ner
Parts Examined: Adnexa	Cornea Drainage Angle Iris	Lens La Vitreous La Fundos La	mments NO BREED RELATED ADNEXAL
RIGHT		LEFT	OR OCULAR CONDITIONS
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	O ) FONDUS	0	DNA sample taken on this date: Yes \( \subseteq \text{No } \subseteq \)
			I confirm that the scanned microchip number matches the number on the certificate
			Information for owners/Appeals leaflet (EPWP1) issued
-	INHERD	TED EYE DISEASE STATUS	
	nherited ocular conditions specified in the Proceed		he KC and/or ISDS as appropriate.
	CLINICALLY CLINICALLY	NON-COM	IGENITAL CLINICALLY CLINICALLY UNAFFECTED AFFECTE
CONGENITAL/NEONATA	UNAFFECTED AFFECTED	(HC) Hereditary cataract	
Collie eye anomaly     Choroidal hypoplasia	4	(PLL) Primary lens luxation	Н
- Coloborna		(POAG) Primary open angle gi (IOP) Intraocular pressure	R mmHg L mmHg
Multifocal retinal dysplasia     Total retinal dysplasia		(PRA) Progressive retinal at	ophy
C) Congenital hereditary catar		(RPED) Retinal pigment epithe	
PV) Persistent hyperplastic prin ) Pectinate ligament abnorma		'Clinically affected' signifies t the inherited disease(s) specif	ed, whereas 'Clinically
de 0 1 2 3 Result		unaffected' signifies that then	is no such evidence.
	nioscopy Grading Result: normal, 1 = mildly affected, 2 = moderately affe	cted: 3 = severely affected.	
0=1	normal, 1 = milaly anected, 2 - moderatory and		
ally affected with ocular conditi	ions not currently specified in the Procedure No	ø€s.	
chiasis	Persistent pupillary membrane	Posterior Cortical Cataract	GPRA-like appearance
pic cilia	Ocular Melanosis	Posterior Polar Subcapsular Cat Posterior Capsular Cataract	
asis	Pectinate ligament abnormality Lens luxation	Posterior Capsular Cataract PHPV	Other conditions (specify)
pion	Anterior Capsular Cataract	Optic nerve hypoplasia	<u> </u>
ined entropion/ectropion	Anterior Cortical Cataract	Posterior segment coloborna	+
ocular defects	Perinuclear Cataract Nuclear Cataract	Choroidal hypoplasia MRD-like appearance	<b>T</b>
UE-1d deposition	7	0 0	
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