14/9/2015

8700 2322 0028 434

Nuphar's Deep In A Dream, Barbet

Registered name: Nuphar's Deep In A Dream
Nickname: Lotus
Registration ID: NHSB 2884576
Microchip: 528140000482532
Breed: Barbet
Gender: Female

Owner: Anne Plomp
Country: Netherlands
Testing date: 12/9/2015
No DNA identification profile.

Dog's identity verified from microchip or tattoo by veterinarian or other authorized person during sample taking: **No**

Test results - Known disorders in the breed

Disorder	Type	Mode of inheritance	Result
Malignant Hyperthermia (MH)	Pharmacogenetics	Autosomal Dominant	Clear

When obtaining a carrier or affected test result, we recommend that you contact your veterinarian for more detailed information on the condition and possible treatment.

On behalf of Genoscoper Laboratories,

Jonas Donner

SIGNATURE

Jonas Donner, PhD, Head of Research and Development
at Genoscoper Laboratories

14/9/2015

8700 2322 0028 434

Nuphar's Deep In A Dream, Barbet

Registered name: Nuphar's Deep In A Dream
Nickname: Lotus
Registration ID: NHSB 2884576
Microchip: 528140000482532
Breed: Barbet
Gender: Female

Owner: Anne Plomp
Country: Netherlands
Testing date: 12/9/2015
No DNA identification profile.

Dog's identity verified from microchip or tattoo by veterinarian or other authorized person during sample taking: **No**

Test results - Traits - page 1/2

Trait	Genotype	Description
Colour Locus E (Extensions)	E/E	The dog is homozygous for E allele.
Colour Locus B (Brown)	bc/bc	The dog is homozygous for bc allele.
Colour Locus K (Dominant Black)	KB/KB || KB/kbr || kbr/kbr	The dog is homozygous for three nucleotide deletion.
Colour Locus A (Agouti)	ay/ay	The dog is homozygous for ay-allele.
Colour Locus S (Piebald or extreme white spotting)	sp/sp	The dog is likely to be extreme white or to have piebald spotting.
Colour Locus H (Harlequin)	h/h	The dog is likely to be non-harlequin.

On behalf of Genoscoper Laboratories,

Jonas Donner

SIGNATURE

Jonas Donner, PhD, Head of Research and Development at Genoscoper Laboratories

8700 2322 0028 434

Nuphar's Deep In A Dream, Barbet

Registered name: Nuphar's Deep In A Dream
Nickname: Lotus
Registration ID: NHSB 2884576
Microchip: 528140000482532
Breed: Barbet
Gender: Female

Owner: Anne Plomp
Country: Netherlands
Testing date: 12/9/2015
No DNA identification profile.

Dog's identity verified from microchip or tattoo by veterinarian or other authorized person during sample taking: **No**

Test results - Traits - page 2/2

Trait	Genotype	Description
Furnishings / Improper Coat in Portuguese Water Dogs (marker test)	AA/TT	The dog is genetically likely to express furnishings.
Body mass, insulin-like growth factor 1 (IGF1) gene variant	A/G	The dog is heterozygous for this variant. This means that it carries one copy of the genetic variant typically associated with small body mass and one copy typically associated with large body mass. This genotype is often observed e.g. in Shetland Sheepdog, Border Collie and Welsh Corgi.
Coat length / "Fluffy" in Welsh Corgi	T/T	The dog carries two copies of the genetic variant typically associated with a long-haired coat. Dogs with this genotype typically have long coat.
Curly coat	T/T	The dog has a curly appearance and it carries two copies of the genetic variant typically associated with a curly coat.
Ear erectness (pricked ears versus floppy ears), variant chr10:11072007	C/C	Your dog is homozygous for (carries two copies of) a genetic variant typically associated with floppy ears. This genotype is common in breeds like English Springer Spaniel, Leonberger, Saluki, and Dachshunds. Interestingly, the C-allele of this variant is the ancestral allele frequent in wolf.
Natural Bobtail (T-box mutation)	C/C	The dog does not carry any copy of the bobtail mutation. It therefore likely has a long-tailed phenotype.
Snout/skull length (shortened head versus elongated head), bone morphogenetic protein 3 (BMP3) gene variant	C/C	Your dog is homozygous for the genetic variant typically found in breeds with an elongated head (e.g. Saluki, Collie, Irish Wolfhound).
Tiny size, insulin-like growth factor 1 receptor (IGF1R) gene variant	G/G	Your dog is homozygous for a genetic variant typically found in larger-sized breeds (height at the withers > 25.4 cm (10 inches)).

On behalf of Genoscoper Laboratories,

Jonas Donner

SIGNATURE

Jonas Donner, PhD, Head of Research and Development at Genoscoper Laboratories

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 1/8

Blood disorders

Disorder	Mode of inheritance	Result
Bleeding disorder due to P2RY12 defect	Autosomal Recessive	Clear
Canine Cyclic Neutropenia (Gray Collie Syndrome)	Autosomal Recessive	Clear
Canine Leucocyte Adhesion Deficiency (CLAD), type III	Autosomal Recessive	Clear
Congenital Macrothrombocytopenia; disease-linked SNP originally found in Norfolk and Cairn Terrier	Autosomal Recessive	Clear
Elliptocytosis		Clear
Factor IX Deficiency or Haemophilia B, Gly379Glu mutation	X-linked Recessive	Clear
Factor IX Deficiency or Haemophilia B; mutation originally found in Airedale Terrier	X-linked Recessive	Clear
Factor IX Deficiency or Haemophilia B; mutation originally found in German Wirehaired Pointer	X-linked Recessive	Clear
Factor IX Deficiency or Haemophilia B; mutation originally found in Lhasa Apso	X-linked Recessive	Clear
Factor IX Deficiency or Haemophilia B; mutation originally found in Rhodesian Ridgeback	X-linked Recessive	Clear
Factor VII Deficiency	Autosomal Recessive	Clear
Factor VIII deficiency or Haemophilia A; mutation originally found in Boxer	X-linked Recessive	Clear
Factor VIII deficiency or Haemophilia A; mutation originally found in German Shepherd Dog	X-linked Recessive	Clear
Factor VIII deficiency or Haemophilia A; p.Cys548Tyr mutation originally found in German Shepherd	X-linked Recessive	Clear
Glanzmann Thrombasthenia (GT), Type I; mutation originally found in Pyrenean Mountain Dog	Autosomal Recessive	Clear
Glycogen Storage Disease VII or Hereditary Phosphofructokinase (PFK) Deficiency	Autosomal Recessive	Clear
May-Hegglin Anomaly (MHA)	Autosomal Dominant	Clear
Prekallikrein Deficiency	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in Basenji	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in Beagle	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in Labrador Retriever	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in Pug	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in West Highland White Terrier	Autosomal Recessive	Clear
Pyruvate Kinase Deficiency of Erythrocyte; mutation originally found in West Highland White Terrier	Autosomal Recessive	Clear
Thrombopathia; mutation originally found in Basset Hound	Autosomal Recessive	Clear
Thrombopathia; mutation originally found in Eskimo Spitz	Autosomal Recessive	Clear
Thrombopathia; mutation originally found in Landseer	Autosomal Recessive	Clear
Trapped Neutrophil Syndrome (TNS)	Autosomal Recessive	Clear
Von Willebrand's Disease (vWD) Type II	Autosomal Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 2/8

Eye disorders

Disorder	Mode of inheritance	Result
Achromatopsia or Cone Degeneration (CD); CNGB3 gene deletion	Autosomal Recessive	Clear
Achromatopsia or Cone Degeneration (CD); mutation originally found in German Shorthaired Pointer	Autosomal Recessive	Clear
Autosomal Dominant Progressive Retinal Atrophy (ADPRA)	Autosomal Dominant	Clear
Canine Multifocal Retinopathy 1 (CMR1), Mastiff-related breeds mutation	Autosomal Recessive	Clear
Canine Multifocal Retinopathy 2 (CMR2); mutation originally found in Coton de Tulear	Autosomal Recessive	Clear
Canine Multifocal Retinopathy 3 (CMR3); mutation originally found in Lapponian Herder	Autosomal Recessive	Clear
Cone-rod Dystrophy (cord1-PRA / crd4)	Autosomal Recessive (Incomplete Penetrance)	Clear
Cone-rod dystrophy (crd SWD); mutation originally found in Standard Wire-haired Dachshund	Autosomal Recessive	Clear
Cone-Rod Dystrophy 1 (crd1); mutation originally found in American Staffordshire Terrier	Autosomal Recessive	Clear
Cone-Rod Dystrophy 2 (crd2); mutation originally found in Pit Bull Terrier	Autosomal Recessive	Clear
Early Retinal Degeneration; mutation originally found in Norwegian Elkhound	Autosomal Recessive	Clear
Generalized Progressive Retinal Atrophy (gPRA)	Autosomal Recessive	Clear
Glaucoma; mutation originally found in Norwegian Elkhound	Autosomal Recessive	Clear
Golden Retriever Progressive Retinal Atrophy 1 (GR_PRA 1)	Autosomal Recessive	Clear
Primary Hereditary Cataract (PHC); mutation originally found in Australian Shepherd	Autosomal Dominant (Incomplete Penetrance)	Clear
Primary Lens Luxation (PLL)	Autosomal Recessive	Clear
Primary Open Angle Glaucoma; mutation originally found in Beagle	Autosomal Recessive	Clear
Progressive Retinal Atrophy (PAP1_PRA); mutation originally found in Papillon and Phalene	Autosomal Recessive	Clear
Progressive Retinal Atrophy (PRA), type III; mutation originally found in Tibetan Spaniel and Tibetan Terrier	Autosomal Recessive	Clear
Progressive Retinal Atrophy - adult onset; mutation originally found in Basenji	Autosomal Recessive	Clear
Rod-Cone Dysplasia 1 (rcd1); mutation originally found in Irish Setter	Autosomal Recessive	Clear
Rod-Cone Dysplasia 1a (rcd1a); mutation originally found in Sloughi	Autosomal Recessive	Clear
Rod-Cone Dysplasia 3 (rcd3)	Autosomal Recessive	Clear
X-Linked Progressive Retinal Atrophy 1 (XLPRA1)	X-linked Recessive	Clear
X-Linked Progressive Retinal Atrophy 2 (XLPRA2)	X-linked Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 3/8

Endocrine disorders

Disorder	Mode of inheritance	Result
Congenital hypothyroidism; mutation originally found in Toy Fox- and Rat Terrier	Autosomal Recessive	Clear

Immunological disorders

Disorder	Mode of inheritance	Result
Autosomal Recessive Severe Combined Immunodeficiency (ARSCID)	Autosomal Recessive	Clear
C3 deficiency	Autosomal Recessive	Clear
Severe Combined Immunodeficiency (SCID); mutation originally found in Frisian Water Dog	Autosomal Recessive	Clear
X-linked Severe Combined Immunodeficiency (XSCID); mutation originally found in Basset Hound	X-linked Recessive	Clear
X-linked Severe Combined Immunodeficiency (XSCID); mutation originally found in Cardigan Welsh Corgi	X-linked Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 4/8

Kidney disorders

Disorder	Mode of inheritance	Result
Cystinuria, Type II-A; mutation originally found in Australian Cattle Dog	Autosomal Dominant	Clear
Cystinuria, Type II-B; mutation originally found in Miniature Pinscher	Autosomal Dominant	Clear
Cystinuria; mutation originally found in Newfoundland Dog	Autosomal Recessive	Clear
Hyperuricosuria and Hyperuricemia (HUU) or Urolithiasis	Autosomal Recessive	Clear
Polycystic Kidney Disease (PKD)	Autosomal Dominant	Clear
Primary hyperoxaluria (PH); mutation originally found in Coton de Tulear	Autosomal Recessive	Clear
Renal Cystadenocarcinoma and Nodular Dermatofibrosis (RCND)	Autosomal Dominant	Clear
X-linked Hereditary Nephropathy (XLHN)	X-linked Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 5/8

Metabolic disorders

Disorder	Mode of inheritance	Result
Glycogen Storage Disease Type II (GSD II), or Pompe's disease	Autosomal Recessive	Clear
Glycogen Storage Disease, Type Ia (GSD Ia)	Autosomal Recessive	Clear
Glycogen Storage Disease, type IIIa (GSD IIIa)	Autosomal Recessive	Clear
Hypocatalasia or Acatalasemia	Autosomal Recessive	Clear
Imerslund-Gräsbeck Syndrome (IGS) or Intestinal cobalamin malabsorbtion; mutation originally found in Beagle	Autosomal Recessive	Clear
Imerslund-Gräsbeck Syndrome (IGS) or Intestinal cobalamin malabsorbtion; mutation originally found in Border Collie	Autosomal Recessive	Clear
Mucopolysaccharidosis Type IIIA (MPS IIIA); mutation originally found in Dachshund	Autosomal Recessive	Clear
Mucopolysaccharidosis Type IIIA (MPS IIIA); mutation originally found in New Zealand Huntaway	Autosomal Recessive	Clear
Mucopolysaccharidosis Type VII (MPS VII); mutation originally found in Brazilian Terrier	Autosomal Recessive	Clear
Mucopolysaccharidosis Type VII (MPS VII); mutation originally found in German Shepherd	Autosomal Recessive	Clear
Pyruvate Dehydrogenase Deficiency	Autosomal Recessive	Clear

Muscular disorders

Disorder	Mode of inheritance	Result
Cavalier King Charles Spaniel Muscular Dystrophy (CKCS-MD)	X-linked Recessive	Clear
Centronuclear Myopathy; mutation originally found in Great Dane	Autosomal Recessive	Clear
Centronuclear Myopathy; mutation originally found in Labrador Retriever	Autosomal Recessive	Clear
Duchenne-like Muscular Dystrophy, Pembroke Welsh Corgi-type	X-linked Recessive	Clear
Muscular Dystrophy, Duchenne type or Golden Retriever Muscular Dystrophy (GRMD)	X-linked Recessive	Clear
Muscular Hypertrophy (double musculing)	Autosomal Recessive	Clear
Myotonia; mutation originally found in Australian Cattle Dog	Autosomal Recessive	Clear
Myotonia; mutation originally found in Miniature Schnauzer	Autosomal Recessive	Clear
Myotubular Myopathy 1 or X-linked Myotubular Myopathy	X-linked Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 6/8

Neurological disorders

Disorder	Mode of inheritance	Result
Alaskan Husky Encephalopathy (AHE)	Autosomal Recessive	Clear
Bandera's Neonatal Ataxia (BNAt)	Autosomal Recessive	Clear
Benign Familial Juvenile Epilepsy or Remitting Focal Epilepsy	Autosomal Recessive	Clear
Cerebellar abiotrophy or neonatal cerebellar cortical degeneration (NCCD)	Autosomal Recessive	Clear
Cerebellar ataxia; mutation originally found in Old English Sheepdog and Gordon Setter	Autosomal Recessive	Clear
Fetal-onset Neuroaxonal Dystrophy (FNAD)	Autosomal Recessive	Clear
Hyperekplexia or Startle Disease	Autosomal Recessive	Clear
Hypomyelination and Tremor; mutation originally found in Weimaraner	Autosomal Recessive	Clear
L-2-Hydroxyglutaric aciduria (L2HGA); mutation 1 originally found in Staffordshire Bull Terrier	Autosomal Recessive	Clear
L-2-Hydroxyglutaric aciduria (L2HGA); mutation 2 originally found in Staffordshire Bull Terrier	Autosomal Recessive	Clear
Lagotto Storage Disease	Autosomal Recessive	Clear
Neonatal Encephalopathy with Seizures (NEWS)	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis 1 (NCL1)	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis 10 (NCL10)	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis 6 (NCL6)	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis 8 (NCL8)	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis 8 (NCL8), rare variant	Autosomal Recessive	Clear
Neuronal Ceroid Lipofuscinosis, type 12, mutation originally found in Tibetan terrier	Autosomal Recessive	Clear
Polyneuropathy; mutation originally found in Alaskan Malamute	Autosomal Recessive	Clear
Polyneuropathy; mutation originally found in Greyhound	Autosomal Recessive	Clear
Progressive early-onset cerebellar ataxia; mutation originally found in Finnish Hound	Autosomal Recessive	No call
Shaking Puppy (X-linked Generalized Tremor); mutation originally found in English Springer Spaniel	X-linked Recessive	Clear
Spinocerebellar ataksia (SCA); mutation originally found in Parson Russell Terrier	Autosomal Recessive	Clear
Spinocerebellar ataxia with myokymia and/or seizures	Autosomal Recessive	Clear

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 7/8

Neuromuscular disorders

Disorder	Mode of inheritance	Result
Congenital Myasthenic Syndrome (CMS)	Autosomal Recessive	Clear
Episodic falling (EF)	Autosomal Recessive	Clear
Globoid Cell Leukodystrophy (GLD) or Krabbe's disease, Terrier mutation	Autosomal Recessive	Clear
Globoid Cell Leukodystrophy (GLD) or Krabbe's disease; mutation originally found in Irish Setter	Autosomal Recessive	Clear
GM1 Gangliosidosis; mutation originally found in Alaskan Husky	Autosomal Recessive	Clear
GM1 Gangliosidosis; mutation originally found in Portuguese Water Dog	Autosomal Recessive	Clear
GM1 Gangliosidosis; mutation originally found in Shiba Dog	Autosomal Recessive	Clear
GM2 Gangliosidosis, mutation originally found in Japanese Chin	Autosomal Recessive	Clear
GM2 Gangliosidosis; mutation originally found in Toy Poodle	Autosomal Recessive	Clear

Skeletal disorders

Disorder	Mode of inheritance	Result
Chondrodysplasia (dwarfism); mutation originally found in Norwegian Elkhound and Karelian Bear Dog	Autosomal Recessive	Clear
Craniomandibular Osteopathy (CMO)	Autosomal Dominant (Incomplete Penetrance)	Clear
Hereditary Vitamin D-Resistant Rickets (HVDRR)	Autosomal Recessive	Clear
Oculoskeletal Dysplasia 2 or Dwarfism-Retinal Dysplasia 2	Autosomal Recessive	Clear
Osteochondrodysplasia; mutation originally found in Miniature Poodle	Autosomal Recessive	Clear
Osteogenesis imperfecta (OI) or Brittle Bone Disease; mutation originally found in Dachshund	Autosomal Recessive	Clear
Osteogenesis imperfecta; mutation originally found in Beagle		Clear
Osteogenesis imperfecta; mutation originally found in Golden Retriever		Clear
Skeletal Dysplasia 2 (SD2)	Autosomal Recessive	Clear

8700 2322 0028 434

Nuphar's Deep In A Dream, Barbet

Test results - Additional disorders found in other breeds - page 8/8

Skin disorders

Disorder	Mode of inheritance	Result
Anhidrotic Ectodermal Dysplasia or X-linked Ectodermal Dysplasia (XHED)	X-linked Recessive	Clear
Epidermolysis bullosa, dystrophic	Autosomal Recessive	Clear
Epidermolytic Hyperkeratosis or Ichthyosis in Norfolk Terrier	Autosomal Recessive	Clear
Hereditary Footpad Hyperkeratosis (HFH)	Autosomal Recessive	Clear
Lamellar Ichthyosis (LI)	Autosomal Recessive	Clear
Musladin-Lueke syndrome (MLS)	Autosomal Recessive	Clear

Other disorders

Disorder	Mode of inheritance	Result
Autosomal Recessive Amelogenesis Imperfecta (ARAI)	Autosomal Recessive	Clear
Cleft palate; mutation originally found in Nova Scotia Duck Tolling Retriever, reverse assay	Autosomal Recessive	Clear
Congenital Keratoconjuctivitis Sicca and Ichthyosiform Dermatosis (CKCSID) or Dry Eye Curly Coat Syndrome	Autosomal Recessive	Clear
Narcolepsy	Autosomal Recessive	Clear
Narcolepsy	Autosomal Recessive	Clear
Narcolepsy	Autosomal Recessive	Clear
Persistant Müllerian Duct Syndrome (PMDS), mutation originally found in Miniature Schnauzer	Autosomal Recessive	Clear
Primary Ciliary Dyskinesia (PCD)	Autosomal Recessive	Clear

On behalf of Genoscoper Laboratories,

Jonas Donner

SIGNATURE

Jonas Donner, PhD, Head of Research and Development at Genoscoper Laboratories

8700 2322 0028 434
Nuphar's Deep In A Dream, Barbet

APPENDIX
Explanation of the results of the tested disorders

Autosomal recessive inheritance (ARI)

Clear - A dog carries no copies of the tested mutation and has no or reduced likelihood of developing and passing on the disease/condition.
Carrier - A dog carries one copy of the tested mutation. Carriers typically have a normal, healthy appearance but pass on the mutation to approximately 50% of their offspring.
Affected - A dog carries two copies of the tested mutation and is at high or increased risk of developing the disease/condition.

Autosomal dominant inheritance (ADI)

Clear - A dog carries no copies of the tested mutation and has no or reduced likelihood of developing and passing on the disease/condition.
Affected - A dog carries one or two copies of the tested mutation and is at high or increased risk of developing the disease/condition.

X-linked recessive inheritance (X-linked)

Clear - A dog carries no copies of the tested mutation and has no or reduced likelihood of developing and passing on the disease/condition.
Carrier - Female carriers typically have a normal, healthy appearance but carry one copy of the tested mutation on one of their X chromosomes. As males only have one X chromosome, there are no male carriers.
Affected - Affected female dogs carry two mutated copies of the tested mutation. Affected males carry one copy of the tested mutation on their single X chromosome. Affected dogs are at high or increased risk of developing the disease/condition.

Please note that the descriptions above are generalized based on typically observed inheritance patterns. When obtaining a carrier or affected test result, always refer to the corresponding online test documentation for more detailed information on the condition and any exceptions.

Genoscoper Laboratories - Legal Notice