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Deraldine's Tea

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Breed Analysis

This section provides a summary of your pet's results for 200+ Breeds we test for. Need a refresher on Breed Genetics? Click the “Learn More” button to review our Onboarding.

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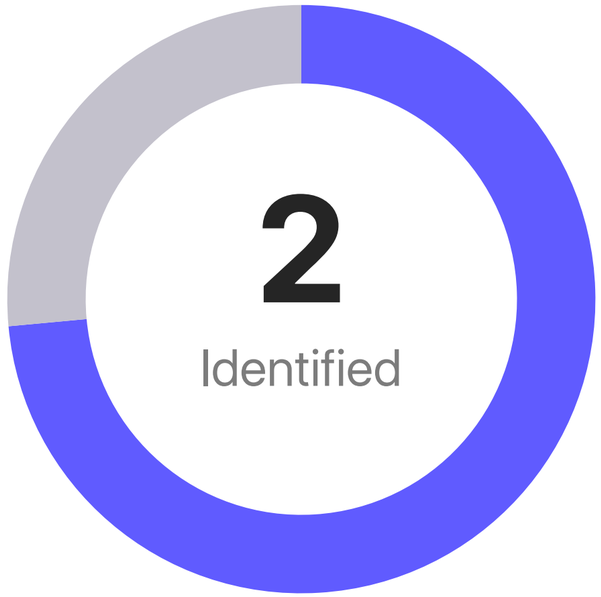
Deraldine's Tea



German Hunting Terrier

10

3.6

Similarity Score

German Hunting Terrier

Read Breed Information

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Genetic Health Predisposition

This section provides a brief summary of your pet's results for 280+ health conditions. Need a refresher on Genetic Health? Click the "Learn More" button to review our Onboarding.

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Predisposition Health Markers

At Risk 1 | Carrier 0 | Clear 213

All

NAME
Brachycephaly

Showing 1 items out of 214 results found

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Genetic Health Predisposition

This section provides a brief summary of your pet's results for 280+ health conditions. Need a refresher on Genetic Health? Click the "Learn More" button to review our Onboarding.

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Predisposition Health Markers

At Risk 1 | Carrier 0 | Clear 213

All

NAME
2,8-DHA urolithiasis
Achromatopsia, CNGA3-related
Achromatopsia, CNGB3-related
Acral mutilation syndrome
Alexander disease
Amelogenesis imperfecta, ENAM-...
Australian Labradoodle dystrop...
Bandera's neonatal ataxia (BNA...
Benign familial juvenile epile...
Beta-mannosidosis
Bleeding disorder, P2RY12-rela...
C3 deficiency
Canine Scott syndrome (CSS)
Cardiomyopathy, YARS2-related
Centronuclear myopathy
Cerebellar ataxia, KCNIP4-rela...
Cerebellar ataxia, RAB24-relat...
Cerebellar ataxia, SEL1L-relat...
Chondrodysplasia, FGF4-related
Chondrodysplasia, ITGA10-relat...
Ciliary dyskinesia, CCDC39-rel...
Ciliary dyskinesia, NME5-relat...
Cone-rod dystrophy 1 (crd1)
Cone-rod dystrophy 2 (crd2)
Cone-rod dystrophy 4 (crd4)
Congenital eye malformation
Congenital Hypothyroidism, SLC...
Congenital hypothyroidism, TPO...
Congenital keratoconjunctiviti...
Congenital myasthenic syndrome...
Congenital myasthenic syndrome...
Congenital myasthenic syndrome...
Congenital stationary night bl...
Craniomandibular osteopathy
Cystinuria, type IA
Cystinuria, type IIA
Cystinuria, type IIB
Dandy-Walker-like malformation...
Deafness, bilateral, and vesti...
Deafness, unilateral and vesti...
Decreased litter size
Degenerative myelopathy
Dental hypomineralization
Diffuse cystic renal dysplasia...
Dilated cardiomyopathy, RBM20-...
Dilated cardiomyopathy, TTN-re...
Disorder of sexual development...
Disproportionate dwarfism
Duchenne muscular dystrophy
Dwarfism, GH1-related
Dwarfism, LHX3-related
Dystrophic epidermolysis bullo...
Early retinal degeneration (er...
Ehlers-Danlos syndrome, classi...
Ehlers-Danlos syndrome, classi...
Ehlers-Danlos syndrome, Dermat...
Elliptocytosis
Epidermolysis bullosa simplex ...
Erythrocytic pyruvate kinase (...
Exercise induced metabolic myo...
Exercise-induced collapse synd...
Factor VII deficiency
Factor XI deficiency
Familial Adenomatous Polyposis
Gangliosidosis, GM1
Gangliosidosis, GM2, type I
Gangliosidosis, GM2, type II
Gastrointestinal stromal tumor...
Generalized myoclonic epilepsy...
Glycogen storage disease, type...
Glycogen storage disease, type...
Glycogen storage disease, type...
Glycogen storage disease, type...
Goniodysgenesis and early-onse...
Haemophilia A
Haemophilia B
Hyperkeratosis, DSG1-related
Hyperkeratosis, FAM83G-related
Hyperkeratosis, KRT10-related
Hypocatalasia (Acatalasemia)
Hypomyelination
Hypophosphatasia
Hypotrichosis
Ichthyosis, PNPLA1-related
Ichthyosis, SLC27A4-related
Imerslund-Gräsbeck syndrome, C...
Inflammatory linear verrucous ...
Inflammatory myopathy
Inherited Myopathy of Great Da...
Junctional epidermolysis bullo...
Junctional epidermolysis bullo...
Juvenile-onset neuroaxonal dys...
Krabbe disease
L-2-hydroxyglutaricacidemia
Laryngeal paralysis and polyne...
Leber congenital amaurosis
Leigh-like subacute necrotizin...
Lens luxation
Lethal acrodermatitis
Leukocyte adhesion deficiency,...
Leukocyte adhesion deficiency,...
Leukodystrophy, CYTB-related
Leukodystrophy, TSEN54-related
Leukoencephalomyelopathy
Ligneous membranitis
Limb-girdle muscular dystrophy...
Limb-girdle muscular dystrophy...
Lundehund syndrome
Malignant hyperthermia
May-Hegglin anomaly
Menkes disease
Methemoglobinaemia
Mitochondrial neurodegenerativ...
Mucopolysaccharidosis IIIA
Mucopolysaccharidosis VI
Mucopolysaccharidosis VII
Multifocal retinopathy 1
Multifocal retinopathy 2
Multifocal retinopathy 3
Multiple drug sensitivity (MDR...
Multiple system degeneration
Muscular dystrophy-dystroglyca...
Muscular dystrophy, COL6A3-rel...
Muscular hypertrophy (double m...
Musladin-Lueke syndrome
Myotonia
Myotubular myopathy
Nasal parakeratosis
Nephritis, X-linked
Nephropathy
Neuroaxonal dystrophy, PLA2G6-...
Neuroaxonal dystrophy, VPS11-r...
Neurodegenerative vacuolar sto...
Neurological defects, MYO5A-re...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuronal ceroid lipofuscinosis...
Neuropathy, sensory ataxic, MT...
neuropathy, sensory, RETREG1(F...
Non-syndromic hearing loss
Oculoskeletal dysplasia, COL9A...
Osteogenesis imperfecta, COL1A...
Osteogenesis imperfecta, COL1A...
Osteogenesis imperfecta, SERPI...
Pelizaeus-Merzbacher disease (...
Periodic Fever Syndrome
Persistent Müllerian duct synd...
Polycystic kidney disease
Polycythemia
Polyneuropathy, ARHGEF10-relat...
Polyneuropathy, GJA9-related
Polyneuropathy, NDRG1-related
Polyneuropathy, RAB3GAP1-relat...
Polyneuropathy, SBF2-related
Prekallikrein deficiency
Primary hyperoxaluria, type I ...
Primary open-angle glaucoma (P...
Primary open-angle glaucoma (P...
Progressive retinal atrophy, B...
Progressive retinal atrophy, C...
Progressive retinal atrophy, C...
Progressive Retinal Atrophy, C...
Progressive Retinal Atrophy, H...
Progressive Retinal Atrophy, I...
Progressive Retinal Atrophy, N...
Progressive retinal atrophy, R...
Progressive Retinal Atrophy, S...
Progressive retinal atrophy, S...
Progressive retinal atrophy, T...
Progressive retinal atrophy, X...
Progressive rod-cone degenerat...
Protein-losing nephropathy
Pulmonary surfactant metabolis...
Pyruvate dehydrogenase deficie...
Recurrent inflammatory pulmona...
Renal cystadenocarcinoma and n...
Respiratory distress syndrome
Rod-cone dysplasia 1 (rcd1), P...
Rod-cone dysplasia 3 (rcd3)
Severe combined immunodeficien...
Severe combined immunodeficien...
Severe combined immunodeficien...
Skeletal dysplasia 2 (SD2)
Spinocerebellar ataxia, CAPN1-...
Spinocerebellar ataxia, KCNJ10...
Spinocerebellar ataxia, SCN8A-...
Spinocerebellar ataxia, SPTBN2...
Spondylocostal dysostosis
Spongy degeneration with cereb...
Spongy degeneration with cereb...
Stargardt disease 1
Succinic semialdehyde dehydrog...
Thrombasthenia
Thrombocytopaenia
Thrombopathia
Trapped Neutrophil Syndrome
Ullrich congenital muscular dy...
Urolithiasis, SLC2A9-related
Ventricular arrhythmias and su...
Von Willebrand disease I
Von Willebrand disease II
Von Willebrand disease III
Wilson's disease, ATP7B-relate...
X-linked hypohidrotic ectoderm...
X-linked retinal dysplasia
Xanthinuria, type I
Xanthinuria, type II

Showing 213 items out of 214 results found