

Code	Test Name
H627	Aberrant Autophagy (LSD)
H758	Abortion (embryonic lethality), BTBD17-related
H386	Achromatopsia 2 (Day Blindness) - German Shepherd
H387	Achromatopsia 2 (Day Blindness) - Labrador Retriever
H904	Achromatopsia 3 (Day Blindness)
H918	Achromatopsia 3 (Day Blindness) - Pointer type
H336	Acral Mutilation Syndrome (AMS)
H338	Acute Respiratory Distress Syndrome (ARDS)
H893	Adult Onset Deafness 1
H822	Adult Onset Deafness EAOD (4 associated markers)
H389	Alexander Disease
H351	Amelogenesis Imperfecta
H679	Amelogenesis Imperfecta 2
H680	Amelogenesis Imperfecta 3
H866	Bardet-Biedl syndrome 2
H788	Beta Mannosidosis
H450	Bleeding disorder due to P2RY12 defect
H487	Brachyury (Bobtail)
H412	C3 Deficiency
H807	Cardiomyopathy and juvenile mortality (CJM) - Belgian Shepherd
H663	Centronuclear Myopathy (CNM) - Border Collie
H749	Centronuclear Myopathy (CNM) - Labrador Retriever
H413	Cerebellar Abiotrophy
H410	Cerebellar Ataxia
H331	Cerebellar Ataxia (SDCA1)
H341	Cerebellar Ataxia (SDCA2)
H804	Cerebellar Ataxia / NCL-A
H653	Cerebellar Ataxia 2
H905	Cerebellar Ataxia, KCNIP4
H411	Cerebellar Ataxia, progressive early-onset
H318	Cerebellar Cortical Degeneration
H355	Cerebellar Hypoplasia Resembling
H356	Chondrodysplasia, disproportionate short-limbed
H346	Chondrodystrophy and Intervertebral Disc Disease
H709	CLAD, Type I
H484	CLAD, Type III
H344	Cleft Lip / Palate and Syndactyly (CLPS)

Code	Test Name
H871	CMR1 (Canine Multifocal Retinopathy)
H730	CMR2 (Canine Multifocal Retinopathy)
H705	Collie Eye Anomaly CEA, CH
H625	Congenital Cornification Disorder
H416	Congenital Hypothyroidism 1 - Dog
H488	Congenital Hypothyroidism 3 - Dog
H694	Congenital Methemoglobinemia - All breeds
H626	Congenital Myasthenic Syndrome (CMS) - Jack Russell Terrier
H339	Congenital Myasthenic Syndrome (CMS) - Labrador Retriever
H425	Congenital Myasthenic Syndrome (CMS) - Old Danish Pointer
H728	Congenital Stationary Night Blindness (CSNB) - Briard
H814	Congenital Stationary Night Blindness (CSNB) - Beagle
H701	Coppertoxicosis
H312	Cranio-mandibular Osteopathy (CMO)
H901	Cranio-mandibular Osteopathy (CMO) - Basset Hound
H357	crd (Cone rod dystrophy) 1
H358	crd (Cone rod dystrophy) 2
H766	crd4-PRA (previously cord1-PRA)
H929	Cystinuria - English and French Bulldog
H703	Cystinuria I - A - 1
H644	Cystinuria, type II - A
H643	Cystinuria, type II - A - 1
H645	Cystinuria, type II - B
H889	Deafness with Vestibular Dysfunction - DVD
H308	Degenerative Myelopathy 2 (DM2)
H673	Degenerative Myelopathy, DM
H327	Dental Hypomineralization
H489	Dermatofibrosis
H865	Dilated Cardiomyopathy (PLN)
H434	Dilated Cardiomyopathy DCM1 (PDK4)
H459	Dilated Cardiomyopathy DCM3 (RBM20)
H876	Disproportionate Dwarfism
H375	Dog_Skin Fragility
H739	Dominant PRA
H913	Dry Eye Curly Coat Syndrome
H497	Dystrophic Epidermolysis Bullosa (RDEB)
H878	Early Onset PRA - Spanish Water Dog

Code	Test Name
H756	Early-onset adult deafness (Rhodesian Ridgeback)
H385	Ectodermal Dysplasia X-linked
H689	Ehlers-Danlos syndrome type 1 - All breeds
H686	Ehlers-Danlos syndrome type 1 - Labrador Retriever
H732	Ehlers-Danlos Syndrome, type 7 - Doberman
H687	Elliptocytosis
H363	Epidermolytic Hyperkeratosis
H486	Epilepsy, BFJ
H672	Exercise Induced Collapse, EIC
H467	Exercise Induced Metabolic Myopathy
H727	Eye malformation, congenital
H435	Factor VII deficiency
H713	Familial Thyroid Follicular Cell Carcinoma I & II
H863	Fanconi syndrome - FS
H869	Fecundity
H676	FN, Familial Nephropathy - (English) Cocker Spaniel
H633	FN, Familial Nephropathy - English Springer Spaniel
H736	Fucosidosis
H702	Gangliosidosis (GM1) - Shiba Inu
H361	Gangliosidosis (GM2 Type I) - Japanese Chin
H490	Gangliosidosis (GM2 Type II) - Poodle Type
H440	Glanzmanns Thrombasthenia (GT) 1 - Dog
H496	Glaucoma (POAG)
H737	Globoid Cell Leukodystrophy / Krabbes Disease
H415	Glycogen Storage Disease GSD I
H472	Goniodysgenesis and glaucoma
H915	gPRA
H868	GR_PRA1
H473	GR_PRA2
H752	Gray Collie Syndrome (Cyclic Neutropenia)
H347	GSDII (Pompe Disease)
H931	Haemophilia A (factor VIII deficiency)
H491	Haemophilia A (Factor VIII)
H607	Haemophilia B (factor IX deficiency)
H780	Haemophilia A - German Shepherd
H392	Hemorrhagic diathesis (Scott Syndrome)
H781	Hereditary Ataxia - Australian Shepherd

Code	Test Name
H809	Hereditary Cataract (HC) - HSF4
H699	Hereditary Cataract 2 (HSF4)
H777	Hereditary Footpad Hyperkeratosis, DSG1-related
H492	Hereditary Footpad Hyperkeratosis, FAM83G-related
H811	Hyperuricemia (HUU)
H364	Hypocatalasia
H365	Hypomyelination
H636	Hypophosphatasia
H873	Ichthyosis - Golden Retriever Type 1
H304	Ichthyosis 3
H378	Ichthyosis 4
H384	Ichthyosis 5
H858	Ichthyosis type 1 & 2 - Golden Retriever
H366	IGS (Selective Cobalamin Malabsorption) 1
H367	IGS (Selective Cobalamin Malabsorption) 2
H867	Inflammatory Myopathy (Myositis)
H726	Inflammatory Pulmonary Disease
H718	Junctionalis Epidermolysis Bullosa, LAMA3-related
H778	Junctionalis Epidermolysis Bullosa, LAMB3-related
H335	Juvenile Myoclonic Epilepsy
H329	Juvenile Laryngeal Paralysis Polyneuropathy (JLPP)
H724	L2-HGA
H641	Laryngeal paralysis (LP)
H693	Laryngeal paralysis and polyneuropathy, CNTNAP1-related
H463	Lethal Acrodermatitis (LAD)
H708	Leukodystrophy
H395	Limb girdle muscular dystrophy (LGMD) 1
H864	Lundehund syndrome - LS
H879	Macrothrombocytopenia (MTC)
H317	Macular Corneal Dystrophy
H746	Malignant Hyperthermia (MH) - All breeds
H882	May-Hegglin Anomaly (MHA) - Pug
H629	MDR1 Multi Drug Resistance
H754	MDR1 Multi Drug Resistance - Phenobarbital Resistance
H786	Metabolizer of a Cognitive Enhancer
H824	Modifier of copper toxicosis, ATP7A-related
H451	Mucopolysaccharidose Type VII - 2

Code	Test Name
H418	Mucopolysaccharidosis Type IIIa
H886	Mucopolysaccharidosis type VI (MPS)
H748	Mucopolysaccharidosis Type VII
H307	Multifocal Retinopathy 3 (cmr3) 2
H747	Muscular Dystrophy (GRMD)
H735	Muscular Dystrophy, COL6A3-related
H359	Muscular Dystrophy, Duchenne type (DMD)
H419	Muscular Dystrophy, Duchenne type (MDM)
H688	Muscular Dystrophy-dystroglycanopathy
H424	Musladin-Lueke Syndroom (MLS)
H391	Myasthenia gravis-like disease
H368	Myopathy
H690	Myotonia - Labrador Retriever
H738	Myotonia Congenita
H498	Myotonia Congenita 2
H776	Myotubular Myopathy
H707	Narcolepsy 1
H697	Narcolepsy 2
H698	Narcolepsy 3
H460	Nasal parakeratosis
H812	Neonatal Encephalopathy (NEWS)
H370	Nephritis
H890	Neuroaxonal Dystrophy (NAD) - Papillon
H888	Neuroaxonal Dystrophy (NAD) - Rottweiler
H462	Neuroaxonal Dystrophy (NAD) - Spanish Water Dog
H428	Neuroaxonal Dystrophy (NAD), MFN2-related
H716	Neurological defects with dilute coat colour
H494	Neuronal ceroid lipofuscinosis (NCL) 1
H429	Neuronal ceroid lipofuscinosis (NCL) 10
H499	Neuronal ceroid lipofuscinosis (NCL) 2
H380	Neuronal ceroid lipofuscinosis (NCL) 5 GR
H330	Neuronal ceroid lipofuscinosis (NCL) 6
H721	Neuronal ceroid lipofuscinosis (NCL), 5
H337	Neuronal ceroid lipofuscinosis NCL, 8-2
H652	Neuronal ceroid lipofuscinosis, 8-1
H895	Obesity
H898	Osteochondrodysplasia (OC)

Code	Test Name
H430	Osteogenesis imperfecta
H431	Osteogenesis imperfecta 2
H381	Osteogenesis Imperfecta 3
H634	Paroxysmal Dyskinesia
H792	Paroxysmal Dyskinesia (PD) - Markiesje
H717	PFK (Phosphofruktokinase Deficiency)
H872	Pituitary dwarfism
H457	POAG / PLL Primary Glaucoma and Lens Luxation
H509	Polycystic Kidney Disease (PKD) - Bull Terrier
H914	Polyneuropathy 1
H495	Polyneuropathy 2
H379	Polyneuropathy 3 (LPN1)
H342	Polyneuropathy 4 (LPN2)
H390	PRA 2
H682	PRA 5
H394	PRA BBS4
H372	PRA crdPRA
H382	PRA erd
H373	PRA type 3
H704	prcd PRA
H439	Prekallikrein deficiency
H414	Primary Ciliary Dyskinesia (PCD), CCDC39-related
H383	Primary Glaucoma
H374	Primary hyperoxaluria
H849	Primary Lens Luxation - PLL
H371	Progressive Retinal Atrophy (Bas-PRA) - Basenji
H808	Progressive Retinal Atrophy (PRA) HIVEP3
H445	Pyruvate Kinase Deficiency (PK) - Basenji
H455	Pyruvate Kinase Deficiency (PKDef) - Beagle
H741	Pyruvate Kinase Deficiency (PKDef) - Labrador Retriever
H454	Pyruvate Kinase Deficiency (PKDef) - Pug
H768	rcd1 PRA
H769	rcd1a PRA
H770	rcd3 PRA
H511	rcd4-PRA
H725	Retinal Dysplasia
H794	Retinal Dysplasia Retinal Folds RD OSD 1

Code	Test Name
H695	Sensory Neuropathy - All breeds
H423	Severe Combined Immunodeficiency - SCID
H456	Severe Combined Immunodeficiency - SCID 2
H891	Shar-Pei Fever (SPAID)
H510	Skeletal Dysplasia 2 (SD2)
H303	Spinocerebellar ataxia
H683	Spinocerebellar Ataxia 2
H328	Spinocerebellar ataxia, LOA (Late Onset Ataxia)
H803	Spondylocostal Dysostosis (Comma Defect)
H907	Stargardt disease 1
H441	Thrombocytopaenia
H442	Thrombopathia
H448	Thrombopathia 2
H449	Thrombopathia 3
H787	Trapped Neutrophil Syndrome (TNS)
H432	Tremor, X-linked
H932	Ventricular Arrhythmias and Sudden Death
H433	Vitamin D-deficiency rickets, type II
H642	Von Willebrand disease 3 - 2
H677	Von-Willebrands Disease Type 1
H345	Von-Willebrands Disease Type 2-2
H744	Von-Willebrands Disease Type 3
H696	Warburg Micro Syndrome 1 (WARBM1)
H825	Wilson Disease
H772	X Linked PRA1 (XL PRA1)
H427	X-linked Myotubular Myopathy (MTM)
H745	X-SCID
H671	Xanthinuria, type 1 - All breeds
H789	Xanthinuria, type 2 - Manchester Terrier