

NAZWA CHOROBY	PUDEL	GOLDEN RETRIEVER	SETER IRLANDZKI	COCKER SPANIEL	MALTAŃCZYK	CAVALIER KING CHARLES SPANIEL	MALTIPOO	CAVAPOO	GOLDENDOODLE	IRISHDOODLE	COCKAPOO
Progressive retinal atrophy (rcd4-PRA)	v		v				v	v	v	v	v
Progressive retinal atrophy (prcd-PRA)	v			v	v		v	v	v	v	v
Squamous cell carcinoma of the toe - risk estimation	v						v	v	v	v	v
Degenerative myelopathy (DM) exon 1											
Degenerative myelopathy (DM exon 2)			v	v							
Degenerative myelopathy (DM exon 2)	v	v			v	v	v	v	v	v	v
GM2 Gangliosidosis	v						v	v	v	v	v
Intervertebral Disc Disease (Type I) CDDY	v				v	v	v	v	v	v	v
CDPA	v				v	v	v	v	v	v	v
Neonatal Encephalopathy with Seizures, NEWS	v						v	v	v	v	v
Osteochondrodysplasia (OCD)	v						v	v	v	v	v
Von Willebrand Disease Type I, Type I vWD	v						v	v	v	v	v
Von Willebrand Disease Type I, Type I vWD			v								
BADANIE ORTOPEDYCZNE	v	v	v	v	v	v	v	v	v	v	v
USG SERCA	v	v	v	v	v	v	v	v	v	v	v
Episodic Falling Syndrome						v		v			
Dry eye curly coat syndrome (CCS)						v		v			
Muscular Dystrophy MD		v				v		v			
Macrothrombocytopenia (MTC)	v			v	v	v	v	v	v	v	v
MCAD deficiency						v		v			
Myxomatous mitral valve disease (MMVD)	v					v	v	v	v	v	v
Xanthinurie Type II				v		v		v			v
Congenital myasthenic syndrome (CMS)		v							v		
Dystrophic Epidermolysis Bullosa		v									
Sensory Ataxic Neuropathy (SAN)		v									
Congenital eye malformations (CEM)		v									
Ichthyosis*		v							v		
Ichthyosis type 2		v							v		
Neuronal ceroid lipofuscinosis (NCL)		v							v		
Osteogenesis imperfecta (Brittle bone disease)		v							v		
Progressive retinal atrophy (GR-PRA1)		v							v		
Progressive retinal atrophy (GR-PRA2)		v							v		
Progressive retinal atrophy (rcd1-PRA)			v							v	
Canine leukocyte adhesion deficiency (CLAD)			v							v	
Globoid cell leukodystrophy (Krabbe disease)			v							v	
Acral mutilation syndrome (AMS)				v							v
Familial nephropathy (FN)*				v							v
Gallbladder mucoceles				v							v
Paradoxical pseudomyotonia (PP)				v							v
X-chromosomal retinal dysplasia				v							v
Achromatopsia (day blindness) (ACHM)											
Alexander disease (AxD)											
Centronuclear myopathy (CNM)											
Congenital myasthenic syndrome (CMS)											
Copper storage disease - Copper toxicosis (CT)*											
Cystinuria											
Dyserythropoietic anemia and myopathy (DAMS)											
Exercise induced collapse (EIC)											
Haemophilia A (Factor VIII deficiency)											
Hereditary nasal parakeratosis (HNPK)											
Laryngeal paralysis with polyneuropathy type 3 (LPPN3)											
Macular corneal dystrophy (MCD)											
Myotonia congenita											
Narcolepsy											
Obesity (ADI)											
Pyruvate kinase deficiency (PK)											
Retinal dysplasia (OSD)*											
Skeletal dysplasia 2 (Dwarfism) (SD2)											
Stargardt disease (STGD, retinal degeneration)											
X-linked myopathy (XL-MTM)											
hyperurikosuria (HUU/SLC)					v		v				
Malignant Hyperthermia (MH)					v		v				
Glycogen storage disease (GSD Ia)					v		v				

niebieski kolor czcionki - podstawowe
czerwony kolor czcionki - pakiety dodatkwoe
Czarny kolor czcionk pozostałe - extra
prime + badania cech zewnetrznych