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TEST	PREZZO (compreso di DEPOSITO e IVA)
Hyperuricosuria (SLC)	60,0
Malignant hyperthermia (MH)	60,0
Degenerative myelopathy exon 2 (DM exon2	60,0
Acatalasemia	60,0
Achromatopsia (day blindness) (ACHM-1)	60,0
Adult onset neuropathy (AON)	80,0
Akutes Lungenversagen (ARDS)	60,0
Alaskan Husky enzephalopathy (AHE)	60,0
Alaskan Malamute polyneuropathy (AMPN)	60,0
Amelogenesis imperfecta (AI)	60,0
Alexander disease	60,0
Bardet-Biedl syndrome (BBS)	60,0
Brachyuria (stumpy tail)	60,0
C-Lokus (albino)	60,0
C3 deficiency (C3)	60,0
Canine leucocyte adhesion deficiency (CLAD)	60,0
Canine multi-focal retinopathy (CMR1/2/3)	80,0
Canine Multiple System Degeneration (CMSD)	60,0
Cerebral dysfunction (CDFS)	60,0
Chondrodysplasia (dwarfism)	60,0
Cleft lip/palate and syndactyly (CLPS)	60,0
Collie eye anomaly (CEA) Optigen	160,0
Collie eye anomaly (CEA) partnerlaboratory	60,0
Combi: Ataxia (LOA + SCA)	90,0
Combi: Ataxia + PLL (LOA + SCA + PLL)	110,0
Combi: DM exon 2 + MDR1	100,0
Combi: Dry Eye Curly Coat + Episodic Falling	65,0
Combi: JE + LSD + Furnishing	110,0
Cone degeneration (CD) Optigen	170,0
Congenital hypothyroidism with goiter (CHG)	60,0
Congenital ichtyosis	60,0
Congenital myasthenic syndrome (CMS)	60,0
Congenital stationary night blindness (CSNB)	60,0
Copper toxicosis (CT)	60,0
Craniomandibular osteopathy (CMO)	60,0
Cystinuria	60,0

DIREZIONE E SEDE CENTRALE
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TEST	PREZZO (compreso di DEPOSITO e IVA)
Dwarfism (pituitary form)	60,0
Dwarfism (Skeletal dysplasia 2)(SD2)	60,0
Dystrophic Epidermolysis bullosa (DEB)	60,0
Ectodermal dysplasia/Skin fragility syndrome	60,0
Epidermolytic hyperkeratosis (EHK)	60,0
Episodic falling (EF)	60,0
Exercise induced collapse (EIC)	80,0
Factor VII deficiency	60,0
Familial nephropathy (FN)	60,0
Familial Nephropathy (FN) partnerlaboratory	80,0
Fanconi syndrome	60,0
Finnish Hound ataxia (FHA)	60,0
Fucosidosis	60,0
Gallbladder mucoceles	60,0
Glanzmann thrombasthenia (GT)	60,0
Globoid cell leucodystrophy (Krabbe-disease)	60,0
Glycogen sorage disease GSD Ia	60,0
Glycogen storage disease GSD II (Pompe)	60,0
Glycogen storage disease type IIIa (GSD3)a	60,0
GM1-Gangliosidosis (GM1)	60,0
GM2-gangliosidosis (Sandhoff disease) (GM2)	60,0
Grey Collie syndrome (GCS)	60,0
Hemophilia A (factor VIII deficiency)	80,0
Hemophilia B (factor IX deficiency)	80,0
Hemorragic diathesis (Scott syndrom)	60,0
Hereditary ataxia (HA)	60,0
Hereditary Cataract (HSF4)	60,0
Hereditary myopathy (CNM)	60,0
Hereditary nasal parakeratosis (HNPK)	60,0
Hereditary neuropathy (GHN)	60,0
Hereditäre nasale Parakeratose (HNPK)	110,0
Hypomyelination/Shaking puppy syndrome (SPS)	60,0
Ichthyosis	60,0
Imerslund-Gräsbeck syndrome (IGS)	60,0

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Junctional epidermolysis bullosa (JEB)	60,0
Juvenile epilepsy (JE)	60,0
Juvenile laryngeal paralysis & polyneuropathy (JLPP)	60,0
Juvenile myoclonic epilepsy (JME)	60,0
L-2-hydroxyglutaric aciduria (L2HGA)	60,0
Lagotto storage disease (LSD)	60,0
Late onset ataxia (LOA)	60,0
Leonberger polyneuropathy (LPN1)	60,0
Leonberger Polyneuropathy 2 (LPN2)	60,0
Leukocyte adhesion deficiency type III (LAD3)	60,0
Lundehund syndrome	60,0
Macrothrombocytopenia (MTC)	60,0
May-Hegglin anomaly (MHA)	60,0
MDR1-gene defect(Ivermectin hypersensibility)	60,0
Mucopolysaccharidosis type IIIa (MPS3a)	60,0
Mucopolysaccharidosis type VII (MPS 7)	60,0
Muscular dystrophy (MD)	60,0
Musladin-Lueke syndrome (MLS)	60,0
Myostation mutation ("bully"-gene)	60,0
Myotonia congenita	60,0
Narcolepsy	60,0
Necrotizing meningoencephalitis (NME/PDE)	60,0
Nemalin myopathy	60,0
Neonatal cortical cerebellar abiotrophy (NCCD	60,0
Neonatal encephalopathy with seizures (NEWS)	60,0
Neuroaxonal dystrophy (NAD)	60,0
Neuronal ceroid lipifuscinosis (NCL)	60,0
Neuronal ceroid lipifuscinosis (NCL) partnerlab	60,0
Obesity	60,0
Osteogenesis imperfecta (brittle bone dis.)	60,0

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TEST	PREZZO (compreso di DEPOSITO e IVA)
Paket Dobermann	130,0
Paket Landseer	130,0
Paket Rhodesian Ridgeback	130,0
Paket Rottweiler	130,0
Paroxysomal dyskinesia (PxD)	60,0
Persistent Müllerian duct snydrome (PMDS)	60,0
Phosphofructokinase deficiency (PFKD)	60,0
Polycystic kidney disease (PKD)	60,0
Postoperative hemorrhage (P2Y12)	60,0
Prekallikrein deficiency (KLK)	60,0
Primary ciliary dyskinesia (PCD)	60,0
Primary hyperoxaluria (PH)	60,0
Primary lens luxation (PLL)	60,0
Primary open angle glaucoma (POAG)	60,0
Primäres Weitwinkelglaukom u. Linsenluxation	60,0
Progressive retinal atrophia (crd2-PRA)	60,0
Progressive retinal atrophia (rcd2-PRA)	60,0
Progressive Retinal Atrophy (Bas_PRA)	60,0
Progressive retinal atrophy (CNGA1-PRA)	60,0
Progressive retinal atrophy (crd-PRA)	60,0
Progressive retinal atrophy (g-PRA)	60,0
Progressive retinal atrophy (GR_PRA1)	60,0
Progressive retinal atrophy (GR_PRA2)	60,0
Progressive Retinal Atrophy (pap_PRA1)	60,0
Progressive retinal atrophy (PRA)	60,0
Progressive retinal atrophy (rcd4-PRA)	60,0
Progressive retinal atrophy (XL-PRA)	60,0
Protein losing nephropathy (PLN)	60,0
Pyruvat dehydrogenase phosphatase 1deficiency	60,0
Pyruvate kinase deficiency (PK)	60,0
Raine syndrome	60,0
Renal Cystadenocarcinoma and nodular fibrosis (RCND)	60,0
Retinal dysplasia (OSD) Optigen	170,0
Retinal dysplasia (OSD) Partnerlaboratory	60,0
Sensorische Neuropathie (SN)	60,0
Severe combined immunodeficiency (SCID)	60,0
Shar Pei Autoinflammatory Disease (SPAID)	60,0

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TEST	PREZZO (compreso di DEPOSITO e IVA)
Renal Cystadenocarcinoma and nodular fibrosis (RCND)	60,0
Retinal dysplasia (OSD) Optigen	170,0
Retinal dysplasia (OSD) Partnerlaboratory	60,0
Sensorische Neuropathie (SN)	60,0
Severe combined immunodeficiency (SCID)	60,0
Shar Pei Autoinflammatory Disease (SPAID)	60,0
Spinal dysraphism	60,0
Spinocerebellar ataxia (SCA)	60,0
Spondylocostal dysostosis (Comma defect)	60,0
Spongy degeneration with cerebellar ataxia	60,0
Spongy degeneration with cerebellar ataxia 1	60,0
Startle disease	80,0
Subakute nekrotisierende Enzephalopathie(SNE)	60,0
Thrombopathia	60,0
Trapped neutrophil syndrome (TNS)	60,0
van den Ende-Gupta syndrome (VDEGS)	60,0
Vitamin D dependent rickets (VDR)	60,0
von-Willebrand disease type I (vWD1)	60,0
von-Willebrand disease type II (vWD 2)	60,0
von-Willebrand disease type III (vWD3)	60,0
Warburg micro syndrome 1 (WARBM1)	60,0
X-chromosomal severe immuno defiziency	60,0
X-linked myopathy (XL-MTM)	60,0
Test per pelo	60,0
A-locus (alleles: Ay, Aw, at, a)	60,0
B-locus (brown, chocolate, liver(nose))	60,0
Coat length I (long or short hair)	60,0
Curly (curled hair)	60,0
D-locus D1 (dilution)	60,0
E-locus (yellow, lemon, red, cream, apricot)	60,0
EM-locus (melanistic mask)	60,0
Furnishing (wire hair)	60,0
K-locus (only the allele KB)	60,0
M-locus (atypicl merle, merle, cryptic merle)**	60,0
S-locus (piebald, white spotting)	60,0

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