



CATÁLOGO POR ESPECIES

SERVICIO DE GENÉTICA

exopolgenomics

A graphic element below the company name, consisting of a series of small, colorful dots (teal, red, grey) arranged in a slightly curved, horizontal line.

PRUEBAS GENÉTICAS



HUELLA GENÉTICA CANINA	Perfil de ADN (identidad, huella genetica) 19 Loci	HCADN
HUELLA GENÉTICA CANINA	Prueba de Paternidad - Progenitor	HCADN_P
HUELLA GENÉTICA CANINA	Prueba de paternidad - Cachorro	HCADN_C
HUELLA GENÉTICA FELINA	Perfil de ADN (identidad, huella genetica)	HCADN
HUELLA GENÉTICA FELINA	Prueba de Paternidad - Progenitor	HCADN_P
HUELLA GENÉTICA FELINA	Prueba de paternidad - Cachorro	HCADN_C
SEXADO DE AVES	Sexado de aves	SXPL
SEXADO DE AVES	Sexado de aves	SXCH

ENFERMEDADES GENÉTICAS CANINAS



CARDIOLÓGICAS	Arrhythmogenic Right Ventricular Cardiomyopathy	ARVC
CARDIOLÓGICAS	Dilated Cardiomyopathy	DCM
CARDIOLÓGICAS	Juvenile Dilated Cardiomyopathy	JDCM
CIRCULATORIAS	Thrombopathia	THR
CIRCULATORIAS	Thrombopathia due to P2Y12 defect	TP2Y12D
CIRCULATORIAS	Von Willebrands Disease Type I	VWF1
CIRCULATORIAS	Von Willebrands Disease Type II	VWF2
CIRCULATORIAS	Von Willebrands Disease Type III	VWF3

DERMATOLÓGICAS	Chronic Superficial Keratitis	CSK
DERMATOLÓGICAS	Coat Color Dilution Alopecia	BHFD
DERMATOLÓGICAS	Dermoid Sinus	DS
DERMATOLÓGICAS	Ectodermal Dysplasia (Hairless)	CED-HL
DERMATOLÓGICAS	Ectodermal Dysplasia (X-linked)	CED-XL
DERMATOLÓGICAS	Ectodermal Dysplasia-Skin Fragility Syndrome	CED-SFS
DERMATOLÓGICAS	Epidermolysis Bullosa, Dystrophic	RDEB
DERMATOLÓGICAS	Epidermolysis Bullosa, Junctional	JEB
DERMATOLÓGICAS	Harequin coat color	HCC
DERMATOLÓGICAS	Ichthiosis	ICT
DERMATOLÓGICAS	Ichthiosis	PNPLA1
DERMATOLÓGICAS	Ichthiosis - Jack Russel Terrier	ICT-JRT
DERMATOLÓGICAS	Ichthiosis -A (Golden Retriever)	ICT-A
INMUNOLÓGICAS	Complement C3 Deficiency	C3
INMUNOLÓGICAS	Cyclic Hematopoiesis	CN
INMUNOLÓGICAS	Leishmania Sensitivity	LESEN
INMUNOLÓGICAS	Severe Combined Immune-Deficiency X-linked	XSCID
INMUNOLÓGICAS	Severe Combined Immune-Deficiency-Autosoma	SCID
METABÓLICAS	MPS I	MPSI
METABÓLICAS	MPS IIIA	MPS IIIa
METABÓLICAS	MPS IIIB	MPS IIIb
METABÓLICAS	MPS VI	MPSVI
METABÓLICAS	MPS VII	MPSVII
METABÓLICAS	Pyruvate Dehydrogenase Phosphatase Deficiency	PDH, PDP-1
MUSCULOESQUELÉTICAS	Centronuclear Myopathy	CNM
MUSCULOESQUELÉTICAS	Chondrodysplasia	CHDY
MUSCULOESQUELÉTICAS	Chondrodysplasia ITGA	CITGA
MUSCULOESQUELÉTICAS	Cleft Palate	CP1
MUSCULOESQUELÉTICAS	Congenital Myasthenic syndrome	CMS
MUSCULOESQUELÉTICAS	Degenerative Myelopathy	DM

MUSCULOESQUELÉTICAS	Dystrophin Muscular Dystrophy	DMD
MUSCULOESQUELÉTICAS	Hereditary Footpad Hyperkeratosis	HFH
MUSCULOESQUELÉTICAS	Hereditary Nasal Parakeratosis	HNPK
MUSCULOESQUELÉTICAS	Malignant Hyperthermia	MH
MUSCULOESQUELÉTICAS	Musladin-Lueke Syndrome	MLS
MUSCULOESQUELÉTICAS	Myostatin Deficiency	MD
MUSCULOESQUELÉTICAS	Myotonia Congenita	MC
MUSCULOESQUELÉTICAS	Myotubular myopathy, x-linked	XLMTM
MUSCULOESQUELÉTICAS	Osteochondrodysplasia	OSTDR
MUSCULOESQUELÉTICAS	Osteogenesis Imperfecta COL1A	OI
MUSCULOESQUELÉTICAS	Pituitary Dwarfism	HD
MUSCULOESQUELÉTICAS	Short Tail (Bobtail)	SHRA-B
MUSCULOESQUELÉTICAS	Skeletal Dysplasia 2	SD2
MUSCULOESQUELÉTICAS	Spinal Dysraphism	NTD
NEUROLÓGICAS	Alaskan Husky Encephalopathy	AHE
NEUROLÓGICAS	Ataxia, Spinocerebellar	SCA
NEUROLÓGICAS	Canine multiple system degeneration	CMSD
NEUROLÓGICAS	Cerebelar ataxia- autophag	CAA
NEUROLÓGICAS	Cerebellar Ataxia	CA
NEUROLÓGICAS	Epilepsy (Lafora Type)	PME
NEUROLÓGICAS	Epilepsy, Phenobarbital resistant	EPR
NEUROLÓGICAS	Episodic falling	EFS
NEUROLÓGICAS	Exercise Induced Collapse	EIC
NEUROLÓGICAS	Familial Nephropathy	ARHN
NEUROLÓGICAS	Fucosidosis	FUDO
NEUROLÓGICAS	GM1 Gangliosidosis	GM1
NEUROLÓGICAS	GM2 Gangliosidosis	GM2
NEUROLÓGICAS	Juvenile Epilepsy	BFJE
NEUROLÓGICAS	L2-Hydroxyglutaric Aciduria	L2HGA
NEUROLÓGICAS	Narcolepsy	NARC

NEUROLÓGICAS	Necrotizing Meningoencephalitis	NME
NEUROLÓGICAS	Neonatal Ataxia	BNAt
NEUROLÓGICAS	Neonatal cerebellar cortical degeneration	NCCD
NEUROLÓGICAS	Neonatal Encephalopathy with Seizures	NEwS
NEUROLÓGICAS	Neuroaxonal dystrophy, fetal-onset	FNAD
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 1	NCL1
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 10	NCL10
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 2	NCL2
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 4A	NCL-4A
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 5	NCL5
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 6	NCL6
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis 8	NCL8
NEUROLÓGICAS	Neuronal Ceroid Lipofuscinosis A	NCL-A
NEUROLÓGICAS	Polyneuropathy	NDRG1
NEUROLÓGICAS	Polyneuropathy, Leonberger	LPN1
NEUROLÓGICAS	Pompe Disease	GSDII
NEUROLÓGICAS	Sensory ataxic neuropathy	SAN
NEUROLÓGICAS	Shaking Puppy (generalized tremor)	SP
NEUROLÓGICAS	Spongiform leukoencephalomyelopathy	SLEM
NEUROLÓGICAS	Startle Disease	SD
NEUROLÓGICAS	Warburg micro syndrome	WMS
OCULARES	Collie Eye Anomaly	CEA
OCULARES	Cone Degeneration	CD
OCULARES	Congenital Stationary Night Blindness	CSNB
OCULARES	Curly Coat Dry Eye	CKCSID
OCULARES	Early Hereditary Cataracts	EHC
OCULARES	Hereditary Cataracts (late-onset)	LHC/HC
OCULARES	Inherited myopathy of Great Danes	IMGD
OCULARES	Multifocal Retinopathy - Coton de Tulear	CMR2
OCULARES	Multifocal Retinopathy - Generica	CMR1

OCULARES	Oculo-Skeletal Dysplasia 1	OSD1
OCULARES	Oculo-Skeletal Dysplasia 2	OSD2
OCULARES	Oculocutaneous Albinism	OCA
OCULARES	Primary Lens Luxation	PLL
OCULARES	Primary Open Angle Glaucoma	POAG
OCULARES	Progressive Retinal Atrophy - adult onset	PARA-AO
OCULARES	Progressive Retinal Atrophy -GR_PRA1	GR-PRA1
OCULARES	Progressive Retinal Atrophy (Dominant)	ADPRA
OCULARES	Progressive retinal atrophy ERD	PRA -erd
OCULARES	Progressive retinal atrophy in Shetland sheepdog	CNGA1
OCULARES	Progressive Retinal Atrophy, crd SWD	CRD
OCULARES	Progressive Retinal Atrophy, crd-1	CRD1
OCULARES	Progressive Retinal Atrophy, crd-2	CRD2
OCULARES	Progressive Retinal Atrophy, crd-3	CRD3
OCULARES	Progressive Retinal Atrophy, Generalized	gPRA
OCULARES	Progressive Retinal Atrophy, GR_PRA2	GR PRA2
OCULARES	Progressive Retinal Atrophy, pcrd	pcrd
OCULARES	Progressive Retinal Atrophy, PRA1	PRA1
OCULARES	Progressive Retinal Atrophy, rcd-1	RCD1
OCULARES	Progressive Retinal Atrophy, rcd-1a	rcd1a
OCULARES	Progressive Retinal Atrophy, rcd-2	rcd2
OCULARES	Progressive Retinal Atrophy, rcd-3	rdc3
OCULARES	Progressive Retinal Atrophy, rcd-4	rcd4
OCULARES	Progressive Retinal Atrophy, Type A	pd
OCULARES	Progressive Retinal Atrophy, type III	P R A
OCULARES	Progressive Retinal Atrophy, X-linked	XLPR
SANGUÍNEAS	Congenital Macrothrombocytopenia	CGMTH
SANGUÍNEAS	Elliptocytosis	ELLCY
SANGUÍNEAS	Factor VII Deficiency	F7
SANGUÍNEAS	Factor XI Deficiency	F11

SANGUÍNEAS	Glanzmanns Thombasthenia Type 1	GTT1
SANGUÍNEAS	Globoid Cell Leukodystrophy	GLD
SANGUÍNEAS	Hemophilia A - Factor VIII Deficiency	FVIII
SANGUÍNEAS	Hemophilia B - Coagulation Factor IX Deficiency	HEMB
SANGUÍNEAS	Iron-Refractory Iron Deficiency Anemia	IRIDA
SANGUÍNEAS	Leukocyte Adhesion Deficiency	LAD
SANGUÍNEAS	May Hegglin Anomaly	MHA
SANGUÍNEAS	Prekallikrein Deficiency	PRDF
SANGUÍNEAS	Pyruvate Kinase Deficiency	PK
SANGUÍNEAS	Trapped Neutrophil Syndrome	TNS
SISTÉMICAS	Alport Syndrome	HN
SISTÉMICAS	Anal Furunculosis	AF
SISTÉMICAS	Autosomal Recessive Amelogenesis Imperfecta	ARAI
SISTÉMICAS	Catalase Deficiency	CAT
SISTÉMICAS	Cobalamin Malabsorption - Amnionless deficiency	IGS
SISTÉMICAS	Cobalamin Malabsorption - cubilin deficiency	IGS
SISTÉMICAS	Congenital Hypothyroidism with Goiter	CHG
SISTÉMICAS	Copper Toxicosis	CT
SISTÉMICAS	Cystinuria	CYS
SISTÉMICAS	Fanconi Syndrome	FS
SISTÉMICAS	Gallbladder mucocele formation	GMF
SISTÉMICAS	Glycogen storage disease IA	GSDI
SISTÉMICAS	Glycogen Storage Disease Type IIIa	GSDIIIa
SISTÉMICAS	Hyperoxaluria	PH1
SISTÉMICAS	Hyperuricosuria	HUU
SISTÉMICAS	Hyperuricosuria 2-8 hidoxyadenine	HU
SISTÉMICAS	Hypomyelination	HYMY
SISTÉMICAS	Multi drugs resistance	MDR1
SISTÉMICAS	Pancreatitis	PACR
SISTÉMICAS	Periodic Fever Syndrome	PFS

SISTÉMICAS	Persistent Mullerian Duct Syndrome	PMDS
SISTÉMICAS	Phosphofructokinase Deficiency	PFKD
SISTÉMICAS	Polycystic Kidney Disease	PKD
SISTÉMICAS	Primary ciliary dyskinesia	PCD
SISTÉMICAS	Protein Losing Nephropathy	PLN
SISTÉMICAS	Renal Cystadenocarcinoma and Nodular Dermatofibrosis	RCND

ENFERMEDADES GENÉTICAS FELINAS



CARDIOLÓGICAS	Hypertrophic Cardiomyopathy	HCM - RD
CARDIOLÓGICAS	Hypertrophic Cardiomyopathy	fHCM
METABÓLICAS	Mucopolidiosis II	MPSI
METABÓLICAS	MPS I	MPS IIIa
METABÓLICAS	MPS VI	MPS IIIb
METABÓLICAS	MPS VI	MPSVI
METABÓLICAS	MPS VII	MPSVII
MUSCULOESQUELÉTICAS	Burmese hypokalaemic periodic polymyopathy	BHP
NEUROLÓGICAS	Dihydropyrimidinase deficiency	DHP
NEUROLÓGICAS	GM1 Gangliosidosis	GM1
NEUROLÓGICAS	GM2 Gangliosidosis	GM2
NEUROLÓGICAS	Niemann - Pick C	NPC
NEUROLÓGICAS	Spinal Muscular Atrophy	SMA
OCULARES	Progressive Retinal Atrophy -rdAc	PRA-rdAc

OCULARES	Progressive Retinal Atrophy -Rdy	PRA-rdy
SANGUÍNEAS	Blood Type B	BloodB
SANGUÍNEAS	Factor XII Deficiency	F12
SANGUÍNEAS	Hemophilia B - Coagulation Factor IX Deficiency	F9
SANGUÍNEAS	Leukocyte Adhesion Deficiency	LAD
SANGUÍNEAS	Lipoprotein Lipase Deficiency	LPL
SANGUÍNEAS	Porphyria - AIP	PorAIP
SANGUÍNEAS	Porphyria - CEP	PorCEP
SANGUÍNEAS	Pyruvate Kinase Deficiency	PK
SANGUÍNEAS	Vitamin D Resistant Rickets	VDDR-1A
SISTÉMICAS	Congenital Hypothyroidism with Goiter	CHG
SISTÉMICAS	Glycogen Storage Disease IV	GSD IV
SISTÉMICAS	Hyperoxaluria	HyXal
SISTÉMICAS	Polycystic Kidney Disease	PKD



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