Slovgen s. r. o. , Diagnostické laboratórium, Ilkovičova 8, 841 04 Bratislava 4, tel : +421 **905 550 916**, IČO: 35 700 629, DIČ: 2020906151, č.ú. : IBAN: SK5511000000002626252786; SK7465000000000020239430
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Žiadanka o odberový materiál

ODBEROVÝ MATERIÁL JE ZDARMA

FORMULÁR VYPLNTE A POSITE NA EMAIL: bielikova@slovgen.sk

PART A: INFORMÁCIE O ZÁKAZNIKOVI

Zákazník:

Meno: ----------

Adresa (Ulica, PSČ, Mesto, Štát): ----------

Telefónne číslo: ----------

e-mail: ----------

PART B: INFROMÁCIE O ZVIERATI # povinné informácie

Druh#: ☐ pes ☐ mačka ☐ vták

Vzorka#: ☐ krv (EDTA) ☐ cytologická kefka ☐ krvná škvrna ☐ pierka

Počet odberových súprav :#
(Počet zvierat na analýzu:)

TAB 1: TYP ANALÝZY – vypĺňa objednávateľ analýzy **PODPIS:**

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ODDIEL C **Požadované vyšetrenie**: Vyberte si typ analýzy zo stĺpca 1 a vpíšte ho do tabuľky TAB 1

OCHORENIA PSOV (analýzy do 5 prac. dní, * analýza priamym sekvenovaním 10-15 dní) **NETLAČIŤ**

1) Analyses	2) Description
AUO-pack 1	for Australian Shepherd: CEA-SG#, DM-SG#, CMR1, HSF4, NBT, MDR1, NCL6, PRA-prcd
AUO-pack 2	for Australian Shepherd: CEA-SG#, DM-SG#, CMR1, HSF4, NBT, MDR1, PRA-prcd
AUO-pack 3	for Australian Shepherd: CEA-SG#, DM-SG#, CMR1, HSF4, MDR1, PRA-prcd
BC-pack 1	for Border Collie: CEA-SG#, GG, IGS, MDR1, NCL5, TNS, SN, Raine
BC-pack 2	for Border Collie: CEA-SG#, GG, IGS, MDR1, TNS, SN, Raine
CEA-EVG	Collie eye anomaly (by cooperating lab EVG Maribor)
CEA-SG	Collie eye anomaly (partner lab under Slovgen supervision)
CKCSID*	dry eye curly coat syndrome for Cavalier King Charles Spaniel - congenital keratoconjuctivitis sicca and ichthyosiform dermatosis
CMO	Craniomandibular osteopathy - for terriers
CMR1	Canine multifocal retinopathy type 1 in Great Pyrenees, English Mastiffs, Bullmastiffs, Cane Corso, Dogue de Bordeaux, English Bulldog, American Bulldog, Pero se Presa Canario and Australian shepherds
CN*- GCS	Cyclic Neutropenia (CN) - **GREY COLLIE SYNDROME** - grey collie, smooth and rough collie
CNM	(Centronuclear Myopathy)/ HMLR (Hereditary myopathy of Labrador retrievers)
CSNB	Congenital Stationary Night Blindness for Briard
CYS	Cystinuria for Newfoundland dogs and Landseer
Fa VII	Factor VII deficiency - Deficit faktoru VII (Airedale Terrier, Alaskan Klee Kai (Miniature Alaskan Husky), Giant Schnautzer and Scottish Deerhound)
DM- DogTest partner lab	Degenerative myelopathy (by cooperating lab **DogTest lab**)
DM-SG partner lab	Degenerative myelopathy (partner lab under Slovgen supervision)
DM - BSP exon 1	Degenerative myelopathy for Bernese mountain dog
DMS	Dermatomyositis Pan2/MAP3K7CL/DLA-DRB1*002:01 for collies and shelties
DWARF	dwarfism (pituary nanism) German shepherds, Saarloos Wolfdogs and Czechoslovakian Wolfdogs
DWARF-LABR/SD2	SD2 - skeletal dysplasia 2 for Labrador
EFS	(Episodic Falling Syndrome) for Cavalier King Charles Spaniel
FN	Familiar Nephropathy – English Cocker Spaniels
FUCA	Fucosidosis in English Springer Spaniels
GG	Goniodysgenesis and glaucoma for Border Collies , Flat coated retriever, Leonberger, Dandie dinmont terrier, Basset, Magyar Viszla, Golden retriever
GRMD	Golden retriever muscular dystrophy
GR-PRA1*	for Golden retriever
GR-PRA2*	for Golden retriever

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1) Analyses	2) Description
HC/HSF4*	hereditary cataract Australian shepherd, Staffordshire Bull Terrier a Boston Terrier
HEM-P2Y12	Hemorrhage – P2Y12 receptor **(**Greater Swiss Mountain dog)
HUU*	Hyperuricosuria
ICT-A:	Congenital Ichthyosis*for Golden retriever
IGS	Imerslund-Gräsbeck syndrom (Border collie)
IVDD (CDPA / CDDY)	Chondrodysplasia, chondrodystrophy and risk of Intervertebral Disc Disease: Dachshunds, Welsh Corgi, Pekingese, Shih Tzu, Cocker Spaniel, French Bulldog and Beagle
JLPP*	Juvenile Laryngeal Paralysis & Polyneuropathy (JLPP) - Rottweiler, Black Russian Terriers
JME	Juvenile Myoclonic Epilepsy - Rhodesian Ridgeback
IMGD	Inherited Myopathy in Great Danes
JRD	Juvenile renal dysplasia - several breeds
L-2-HGA*	L-2-HGA - L-2-hydroxyglutaric aciduria in Staffordshire bull terriers
MDL	Muscular dystrophy for Landseer
MH*	Malignant Hyperthermia, all dogs
MDR1	multidrug resistance gene – direct detection of nt230(del4)
MLS	Musladin-Leuke Syndrom (Beagle)
NAD	Neuroaxonal dystrophy for Papillons
NAR	Narcolepsy for Labrador Retriever
NBT	Bob Tail/ Short Tail
NCCD	Neonatal cerebellar cortical degeneration - cerebellar abiotrophy - Beagle
NCL5	(neuronal ceroid lipofuscinosis – for Border collie
NCL	(neuronal ceroid lipofuscinosis – for Tibetan Terriers
NCL6*	(neuronal ceroid lipofuscinosis – for Australian Shepherd
N. can	Neospora caninum
NEWS	Neonatal encephalopathy with seizures – Standard Poodle
OCA-2	OCA-2 - Spitz - Oculocutaneous albinism type 2
OCA-4 Bull	OCA - Oculocutaneous albinism Bullmastiff (c.1287delC in gene SLC45A2)
OCA-4 Dob	OCA - Oculocutaneous albinism Doberman (SLC45A2, 4081bp del)
pap-PRA1	progressive retinal atrophy - for Pappilons, Phalens
PK-BEAG	Deficiency of pyruvate kinase for Beagle
PK-LABR	Deficiency of pyruvate kinase for Labrador retriever
PK-MOPS	Deficiency of pyruvate kinase for Pug (Mops)
PK-WHWT	Deficiency of pyruvate kinase for Cairn Terrier and West Highland White Terrier
PLL	Primary lens luxation
PRA1-GR*	Progressive retinal atrophy for Golden retriever = **GR-PRA1***
PRA2-GR*	Progressive retinal atrophy for Golden retriever = **GR-PRA2***
PRA-cord1	Progressive retinal atrophy for Standard wire-haired dachshunds, miniature long-haired dachshunds, English springer spaniels and pit bull terriers

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1) Analyses	2) Description
PRA-cord2	Progressive retinal atrophy for standard wire-haired dachshunds
PRA-prcd	Progressive retinal atrophy for several breeds
PRA-rcd2	Progressive retinal atrophy for Collie
PRA-rcd3	Progressive retinal atrophy for Cardigan Welsh Corgi
PRA-rcd4*	Progressive retinal atrophy for Terriers, Setters etc.
PRA-shet	Progressive retinal atrophy for Sheltie (CNGA1)
Raine Syndrome (DH)	Raine syndrome - Dental Hypomineralization for Border Collie
SD2 - Dwarf LABR	SD2 - skeletal dysplasia 2 - Dwarf for Labrador
SDCA1	SDCA1 -Spongy cerebellar degeneration with cerebellar ataxia (Belgian shepherds – malinois)
SDCA2	SDCA2 -Spongy cerebellar degeneration with cerebellar ataxia (Belgian shepherds – malinois)
SN	Sensory neuropathy for Border Collie
TNS	(Trapped Neutrophil Syndrome – for Border collie)
T. gondii	Toxoplasma gondii
VDEG	Van den Ende-Gupta syndrome in Wire Fox Terrier
vWDI	von Willebrand disease for several breed Basethound, Bernese mountain dog, Coton de Tulear, Dachshund (standard, mini), Doberman pinscher, Drentsche Patrijschond, German Shepherd, Golden retriever, Goldendoodle, Keeshound, Kerry blue terrier, Labradoodle, Manchester terrier, Miniature Schnauzer, Papillion, Poodle, Rotttweiler, Stabyhoun, Welsh Corgi Pembroke
vWDII	von Willebrand disease for German Shorthaired Pointers, German Wirehaired Pointers and Collies
vWDIII	von Willebrand disease for Shetland Sheepdog

GENETICKÝ PROFIL A POTVRDENIE RODIČOVSTVA (PARENTITA)

1) TYP ANALÝZY	2) POPIS
DNA profile	STR-21 loci + amelogenin (ISAG 2006), profil DNA včítane prípadneho potvrdenia rodičov

FARBY A KVALITA SRSTI PSOV

1) TYP ANALÝZY	2) POPIS
Bez-srstosť - Hairlessness	Nahatosť - for Chinese Crested, Peruvian Inca Orchid and Mexican Xoloitzcuintle
FGF5	Fluffy gene, long or short hair
KRT71	Curly coat/ kučeravosť u psov
Locus A	(coat colour – ay >aw> at> a)
Locus B	(coat colour)
Locus BE*	(coat colour) – brown for Australian Shepherd
Locus D	(coat colour)
Locus E	(coat colour)
Locus K	(coat colour)
Locus EM	(coat colour) -melanistic mask
Locus M - Merle	(coat colour), merle/cryptic merle
Locus S (spotting)	(coat colour), white factor
RSPO2/IC	improper coat

OCHORENIA MAČIEK

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1) TYP ANALÝZY	2) POPIS
PKD	Polycystic Kidney Disease (for cats)

Vtáci

1) TYP ANALÝZY	2) POPIS
Sex	Určenie pohlavia
PBFD	Psittacine Beak and Feather Disease
APV	Avian Polyoma Virus
Ch. psittaci	Chlamydophila psittaci