

Result certificate #012345

Dog Chondrodysplasia (CDPA) and Chondrodystrophy (CDDY) with risk of intervertebral disk disease (IVDD)

Customer: Jan Novák, Dlouhá 1, 30000 Plzeň, Czech Republic

Sample: Sample: 21-12345 Date received: 01.02.2021 Sample type: blood

Information provided by the customer Name: Lassie DEMO Breed: Plemeno

Tattoo number: 1392013 Microchip: 123 456 789 012 345 Reg. number: REGQ12345 Date of birth: 1.1.2020 Sex: female Date of sampling: 01.02.2021 The identity of the animal has been checked.

Result Chondrodysplasia: N/CDPA Result Chondrodystrophy and IVDD risk: N/CDDY

Explanation

The presence or absence of two insertion mutations in retrogene for fibroblast growth factor FGF4 in chromosomes 12 and 18 was examined. FGF4 retrogene is involved in many biological processes incl. bone development. Dogs that carry both mutations show a more drastic reduction of leg length (e.g. Basset Hounds, Dachshund, Welsh Corgi and Scottish terriers).

Chondrodysplasia (CDPA) is caused by the insertion of FGF4 retrogene into chromosome 18, is responsible for short legs in baset, welsh, dachshund, western, Scottish terrier and more. Inheritance of CDPA mutation is autosomal dominant, only one allele passed from one of the parents is sufficient to show the shortened legs.

Chondrodystrophy (CDDY) is caused by the insertion of FGF4 retrogen into chromosome 12 and explains the leg shortening of other breeds. The mutation is also connected with predisposition to abnormal growth and development of intervertebral discs. In affected dogs, premature calcification before 1 year of age results in loss of flexibility and gradual degeneration of intervertebral discs. These abnormal discs are predisposed to herniation into the spinal canal where the inflammation and haemorrhage can cause severe pain and neurological dysfunction termed Intervertebral Disc Disease of type I (IVDD).

The CDDY is inherited as a semi-dominant trait for height, meaning that dogs with 2 copies of the mutation (CDDY/CDDY) are smaller than dogs with only 1 copy (N/CDDY). As to predisposition to degeneration of intervertebral discs, the inheritance follows an autosomal dominant mode meaning that 1 allele obtained from one of its parents is sufficient to expressing the phenotype.

Genomia s.r.o, Republikánská 6, 31200 Plzeň, Czech Republic www.genomia.cz, laborator@genomia.cz, tel: +420 373 749 999



Result certificate #012345

Chondrodysplasia result codes:

N/N	No copies of CDPA mutation.
N/CDPA	1 copy of CDPA. Mutation causes leg shortening compared to N/N dogs.
CDPA/CDPA	2 copies of CDPA. Muta <mark>tion cause</mark> s leg shortening compared to N/N dogs.

Chondrodystrophy (CDDY and IVDD Risk) result codes:

N/N	No copies of CDDY mut <mark>ation.</mark>	
N/CDDY		at risk for IVDD. Mutation causes leg shortening od to an N/N dog will produce 50% of normal sized sk for IVDD.
CDDY/CDDY		r IVDD. Mutation causes leg shortening compared to suppies with shorter legs at risk for IVDD.

Method: SOP176-IVDD, ASA-PCR

Date of issue: 06.02.2021 Date of testing: 01.02.2021 - 06.02.2021 Approved by: Mgr. Martina Šafrová, Laboratory Manager



Genomia s.r.o, Republikánská 6, 31200 Plzeň, Czech Republic www.genomia.cz, laborator@genomia.cz, tel: +420 373 749 999