

## French Bulldog



[Labgenvet](#)  
[Donner, 2018](#)  
[Donner, 2023](#)

| Disease                                                                   | Gene       | Reference       | # tests | M/N    |          |
|---------------------------------------------------------------------------|------------|-----------------|---------|--------|----------|
| <a href="#">Cystinuria Type I-A (SLC3A1 p.S698G)</a>                      | SLC3A1     | Donner, 2023    | 13114   | 36.4%  |          |
| <a href="#">Cystinuria Type I-A (SLC3A1 p.I192V)</a>                      | SLC3A1     | Donner, 2023    | 13114   | 36.0%  |          |
| <a href="#">Degenerative Myelopathy (DM)</a>                              | SOD1       | Labgenvet, 2021 | 5       | 32.0%  |          |
|                                                                           |            | Donner, 2018    | 85      | 28.0%  |          |
|                                                                           |            | Donner, 2023    | 13111   | 31.5%  |          |
| <a href="#">Chondrodystrophy, Intervertebral Disc Disease Risk (CDDY)</a> | FGF4       | Donner, 2023    | 13062   | 21.3%  | DOMINANT |
| <a href="#">Cone-Rod Dystrophy (cord1-PRA/crd4)</a>                       | RPGRIP1    | Labgenvet, 2021 | 4       | 25.0%  |          |
|                                                                           |            | Donner, 2018    | 85      | 23.5%  |          |
|                                                                           |            | Donner, 2023    | 13097   | 14.7%  |          |
| <a href="#">Canine Multifocal Retinopathy 1 (Mastiff; CMR1)</a>           | BEST1      | Donner, 2023    | 13114   | 10.2%  |          |
| <a href="#">Hereditary cataracts (terriers)</a>                           | HSF4       | Donner, 2018    | 62      | 8.8%   |          |
| <a href="#">Dilated Cardiomyopathy risk factor (PDK4-related)</a>         | PDK4       | Donner, 2023    | 13114   | 6.6%   | DOMINANT |
| <a href="#">Cystinuria Type I-B (SLC7A9 p.A217T)</a>                      | SLC7A9     | Donner, 2023    | 13113   | 1.3%   |          |
| <a href="#">Hyperuricosuria (HUU)</a>                                     | SLC2A9     | Donner, 2023    | 13114   | 0.4%   |          |
| <a href="#">Hypocalasias</a>                                              | CAT        | Donner, 2023    | 13113   | 0.03%  |          |
| <a href="#">Von Willebrand's Disease, Type 1 (vWD 1)</a>                  | VWF        | Donner, 2023    | 13114   | 0.03%  |          |
| Mutation reported at very low frequency (Donner, 2023):                   |            |                 |         |        |          |
| Shar-Pei Autoinflammatory Disease (SPAID)                                 | MTBP, HAS2 | Donner, 2023    | 13114   | >0.01% |          |
| Early-Onset Progressive Polyneuropathy (Greyhound)                        | NDRG1      | Donner, 2023    | 13109   | >0.01% |          |
| Craniomandibular Osteopathy (Terrier)                                     | SLC37A2    | Donner, 2023    | 13114   | >0.01% | DOMINANT |
| Neonatal Cerebellar Cortical Degeneration (NCCD)                          | SPTBN2     | Donner, 2023    | 13114   | >0.01% |          |