### MAE

## Veterinary Report by Embark

embarkvet.com

Test Date: June 6th, 2023

## Customer-supplied information

Owner Name: William Gibson

Dog Name: Mae Sex: Female Date of birth: n/a Breed type: N/A

Breed: Labrador Retriever

Breed registration: The Kennel Club (KC)

Microchip: N/A

### Genetic summary

Genetic breed identification:

**Labrador Retriever** 

Breed ancestry:

Labrador Retriever: 100.0%

Predicted adult weight: **61 lbs** Calculated from 17 size genes.

### **Health Report**

#### How to interpret Mae's genetic health results:

If Mae inherited any of the variants that we tested, they will be listed at the top of the Health Report section, along with a description of how to interpret this result. We also include all of the variants that we tested Mae for that we did not detect the risk variant for.

#### A genetic test is not a diagnosis

This genetic test does not diagnose a disease. Please talk to your vet about your dog's genetic results, or if you think that your pet may have a health condition or disease.

#### **Summary**

Mae is not at increased risk for the genetic health conditions that Embark tests.

**Breed-relevant** (18)

**Other** (225)

# **Health Report**

### **BREED-RELEVANT RESULTS**

Research studies indicate that these results are more relevant to dogs like Mae, and may influence her chances of developing certain health conditions.

| Alexander Disease (GFAP)  | Clear |
|---|-------|
| Oanine Elliptocytosis (SPTB Exon 30)  | Clear |
| Centronuclear Myopathy, CNM (PTPLA)   | Clear |
| Congenital Myasthenic Syndrome, CMS (COLQ, Labrador Retriever Variant)                  | Clear |
| Oay Blindness (CNGA3 Exon 7, Labrador Retriever Variant)                                | Clear |
| Exercise-Induced Collapse, EIC (DNM1)   | Clear |
| Golden Retriever Progressive Retinal Atrophy 2, GR-PRA2 (TTC8)                          | Clear |
| Hereditary Nasal Parakeratosis, HNPK (SUV39H2)  | Clear |
| Macular Corneal Dystrophy, MCD (CHST6)  | Clear |
| Narcolepsy (HCRTR2 Intron 6, Labrador Retriever Variant)                                | Clear |
| Progressive Retinal Atrophy, crd4/cord1 (RPGRIP1)                                       | Clear |
| Progressive Retinal Atrophy, prcd (PRCD Exon 1)   | Clear |
| Pyruvate Kinase Deficiency (PKLR Exon 7, Labrador Retriever Variant)                    | Clear |
| Skeletal Dysplasia 2, SD2 (COL11A2, Labrador Retriever Variant)                         | Clear |
| Stargardt Disease (ABCA4 Exon 28, Labrador Retriever Variant)                           | Clear |
| Ullrich-like Congenital Muscular Dystrophy (COL6A3 Exon 10, Labrador Retriever Variant) | Clear |
|   | Clear |
| X-Linked Myotubular Myopathy (MTM1, Labrador Retriever Variant)                         | Clear |