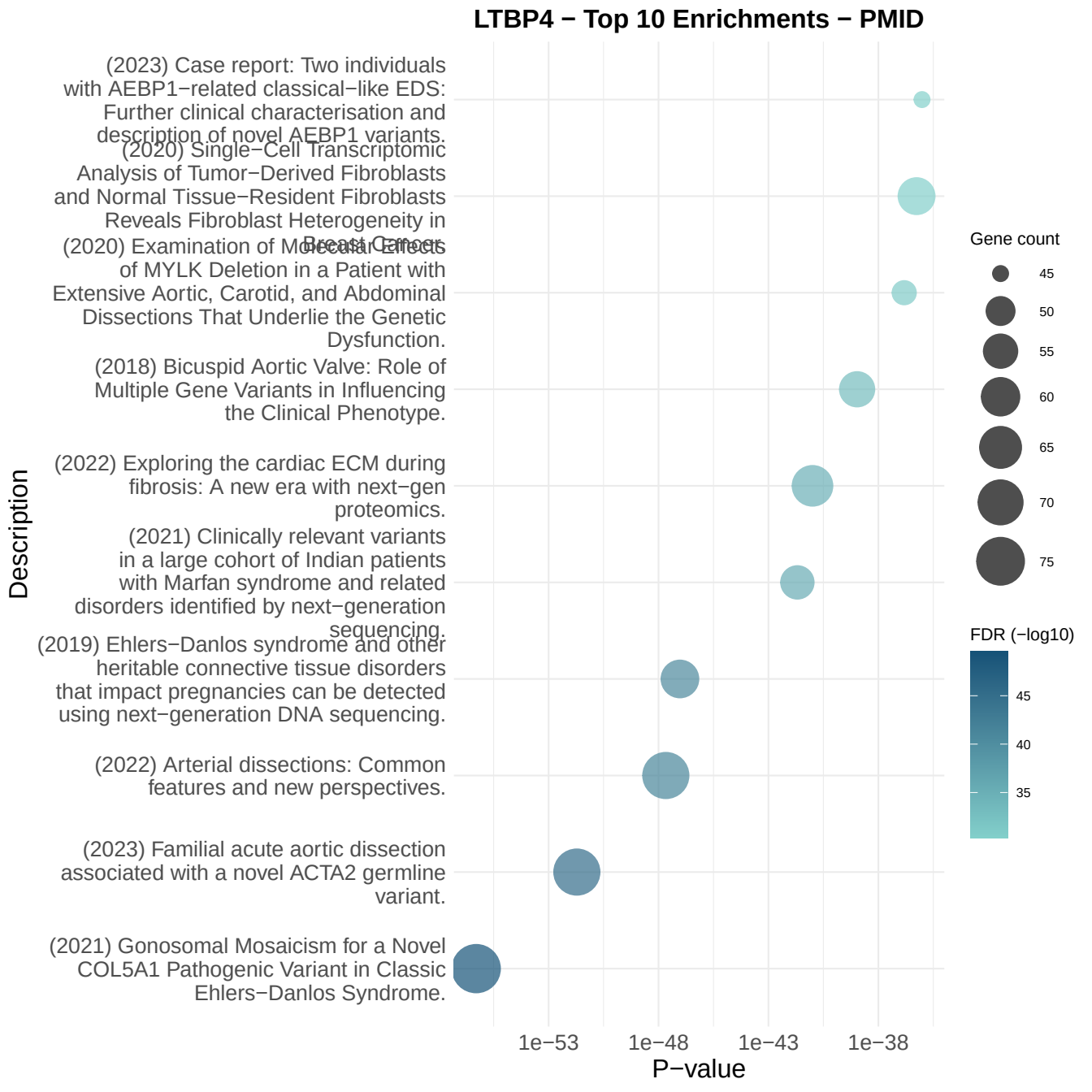
LTBP4 − Top 10 Enrichments − PMID
(2023) Case report: Two individuals with AEBP1−related classical−like EDS: Further clinical characterisation and description of novel AEBP1 variants.
(2020) Single−Cell Transcriptomic Analysis of Tumor−Derived Fibroblasts and Normal Tissue−Resident Fibroblasts Reveals Fibroblast Heterogeneity in Breast Cancer.
(2020) Examination of Molecular Effects of MYLK Deletion in a Patient with Extensive Aortic, Carotid, and Abdominal Dissections That Underlie the Genetic Dysfunction.
(2018) Bicuspid Aortic Valve: Role of Multiple Gene Variants in Influencing the Clinical Phenotype.
(2022) Exploring the cardiac ECM during fibrosis: A new era with next−gen proteomics.
(2021) Clinically relevant variants in a large cohort of Indian patients with Marfan syndrome and related disorders identified by next−generation sequencing.
(2019) Ehlers−Danlos syndrome and other heritable connective tissue disorders that impact pregnancies can be detected using next−generation DNA sequencing.
(2022) Arterial dissections: Common features and new perspectives.
(2023) Familial acute aortic dissection associated with a novel ACTA2 germline variant.
(2021) Gonosomal Mosaicism for a Novel COL5A1 Pathogenic Variant in Classic Ehlers−Danlos Syndrome.
Description
1e−53
1e−48
1e−43
1e−38
P−value
Gene count
45
50
55
60
65
70
75
FDR (−log10)
45
40
35