GeDiPNet Shared-Gene Disease Pairs

Similarity: Jaccard index on shared curated genes (disease_gdp). Significance: upper-tail hypergeometric test (P(X>=k)), Benjamini-Hochberg FDR-corrected across all tested pairs. Corroborated Shared Genes: count of shared genes backed by >=2 distinct disease_gdp.main_source values combined across both diseases (e.g. GWAS-backed for one, ClinVar-backed for the other still counts); Shared Gene Names lists each gene's own source count in parentheses. Overlap Coefficient: shared genes / MIN(each disease's own total gene count) -- complements the Jaccard-based Similarity Score by showing how much of the SMALLER disease's entire known gene-set is covered by this pair, which Jaccard (dividing by the union) can understate. Shared Cluster: cluster id if both diseases fall in the same label-propagation cluster (scripts/build_disease_clusters.php), blank otherwise. See scripts/build_related.php on gedipnet.bicnirrh.res.in. Data as of: 2026-10-21 12:58:11. Filters applied: disease name contains "Myopathy caused by variation in POMT2" (5 matching pairs). · Rows: 5 · Generated: 2026-10-21 13:15:21

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Myopathy caused by variation in POMT2	Walker-warburg syndrome	1	1	POMT2 (4)	0.05882	1.00000	1.033e-3	1.574e-3	
Muscle eye brain disease	Myopathy caused by variation in POMT2	1	1	POMT2 (4)	0.06250	1.00000	9.685e-4	1.497e-3	
Muscular dystrophy-dystroglycanopathy (congenital with brain and eye anomalies), type a	Myopathy caused by variation in POMT2	1	1	POMT2 (2)	0.14286	1.00000	3.874e-4	7.329e-4	
Muscular dystrophy-dystroglycanopathy (congenital with brain and eye anomalies), type a2	Myopathy caused by variation in POMT2	1	1	POMT2 (2)	0.50000	1.00000	6.457e-5	1.985e-4	
Muscular dystrophy-dystroglycanopathy (congenital with impaired intellectual development), type b2	Myopathy caused by variation in POMT2	1	1	POMT2 (2)	0.50000	1.00000	6.457e-5	1.985e-4	