

GeDiPNet Shared-Gene Disease Pairs

Similarity: Jaccard index on shared curated genes (disease_gdp). Significance: upper-tail hypergeometric test ($P(X \geq k)$), 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: 2026-10-11 03:00:14. Filters applied: disease name contains "Monomelic amyotrophy" (5 matching pairs). · Rows: 5 · Generated: 2026-10-11 20:39:18

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Hydrops fetalis	Monomelic amyotrophy	1	0	RYR3 (1)	0.04545	0.16667	6.219e-3	7.474e-3	
Encephalocele	Monomelic amyotrophy	1	0	CPLANE1 (1)	0.10000	0.25000	1.558e-3	2.376e-3	
Brachial plexus neuritis	Monomelic amyotrophy	1	1	SEPTIN9 (3)	0.14286	1.00000	3.897e-4	8.665e-4	
joubert syndrome 17	Monomelic amyotrophy	1	1	CPLANE1 (2)	0.14286	1.00000	3.897e-4	8.665e-4	
Monomelic amyotrophy	Neuralgic amyotrophy	1	1	SEPTIN9 (4)	0.14286	1.00000	3.897e-4	8.665e-4	