

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 "microcephaly 26, primary, autosomal dominant" (5 matching pairs). · Rows: 5 · Generated: 2026-10-10 22:26:23

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Demyelinating leukodystrophy	microcephaly 26, primary, autosomal dominant	1	1	LMNB1 (2)	0.50000	1.00000	6.494e-5	2.331e-4	
microcephaly 26, primary, autosomal dominant	Primary microcephaly	1	1	LMNB1 (4)	0.02778	1.00000	2.273e-3	3.191e-3	
Microcephaly	microcephaly 26, primary, autosomal dominant	1	1	LMNB1 (5)	0.00935	1.00000	6.884e-3	8.163e-3	
Dyslexia	microcephaly 26, primary, autosomal dominant	1	1	LMNB1 (2)	0.00893	1.00000	7.209e-3	8.515e-3	
microcephaly 26, primary, autosomal dominant	Parkinson disease	1	1	LMNB1 (2)	0.00189	1.00000	3.436e-2	3.633e-2	