

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, seizures, and developmental delay" (5 matching pairs). · Rows: 5 · Generated: 2026-10-10 23:44:36

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Congenital microcephaly	microcephaly, seizures, and developmental delay	1	1	PNKP (2)	0.03125	1.00000	2.013e-3	2.901e-3	
Cerebellar atrophy	microcephaly, seizures, and developmental delay	1	1	PNKP (2)	0.03226	1.00000	1.948e-3	2.828e-3	
microcephaly, seizures, and developmental delay	Pyridoxine dependent epilepsy	1	1	PNKP (2)	0.11111	1.00000	5.195e-4	1.061e-3	
Ataxia with oculomotor apraxia	microcephaly, seizures, and developmental delay	1	1	PNKP (7)	0.16667	1.00000	3.247e-4	7.709e-4	
Dna repair-deficiency disorders	microcephaly, seizures, and developmental delay	1	1	PNKP (3)	0.33333	1.00000	1.299e-4	3.931e-4	