

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 "neuroferritinopathy" (5 matching pairs). · Rows: 5 · Generated: 2026-10-11 21:08:04

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Neurodegeneration with brain iron accumulation	neuroferritinopathy	1	1	FTL (3)	0.06667	1.00000	9.092e-4	1.576e-3	
Basal ganglia disease	neuroferritinopathy	1	1	FTL (2)	0.07143	1.00000	8.443e-4	1.499e-3	
Hyperferritinemia	neuroferritinopathy	1	1	FTL (4)	0.33333	1.00000	1.299e-4	3.931e-4	
Genetic hyperferritinemia without iron overload	neuroferritinopathy	1	1	FTL (3)	0.50000	1.00000	6.494e-5	2.331e-4	
hereditary hyperferritinemia with congenital cataracts	neuroferritinopathy	1	0	FTL (1)	0.50000	1.00000	6.494e-5	2.331e-4	