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: 2026-10-20 03:00:20. Filters applied: disease name contains "Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6" (5 matching pairs). · Rows: 5 · Generated: 2026-10-20 04:47:05

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6	Seckel syndrome	1	1	DNA2 (5)	0.05263	1.00000	1.162e-3	1.729e-3	
Mitochondrial myopathy	Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6	1	1	DNA2 (2)	0.05556	1.00000	1.098e-3	1.651e-3	
Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6	Rothmund-thomson syndrome	1	1	DNA2 (4)	0.07692	1.00000	7.748e-4	1.247e-3	
Microcephalic primordial dwarfism with or without poikiloderma and cataracts	Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6	1	0	DNA2 (1)	0.50000	1.00000	6.457e-5	1.985e-4	
Mitochondrial dna deletion syndrome	Progressive external ophthalmoplegia with mitochondrial dna deletions, autosomal dominant, 6	1	1	DNA2 (3)	0.50000	1.00000	6.457e-5	1.985e-4	