

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  $\geq 2$  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: 2026-10-10 03:00:13. Filters applied: disease name contains "congenital microcephaly - severe encephalopathy - progressive cerebral atrophy syndrome" (5 matching pairs). · Rows: 5 · Generated: 2026-10-10 08:32:18

| 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 - severe encephalopathy - progressive cerebral atrophy syndrome | Lipidoses                                                                               | 1            | 1                         | ASNS (2)          | 0.05000          | 1.00000             | 1.234e-3 | 1.986e-3    |                |
| congenital microcephaly - severe encephalopathy - progressive cerebral atrophy syndrome | Lipoidosis                                                                              | 1            | 1                         | ASNS (2)          | 0.05000          | 1.00000             | 1.234e-3 | 1.986e-3    |                |
| Congenital microcephaly                                                                 | congenital microcephaly - severe encephalopathy - progressive cerebral atrophy syndrome | 1            | 1                         | ASNS (5)          | 0.03125          | 1.00000             | 2.013e-3 | 2.901e-3    |                |
| congenital microcephaly - severe encephalopathy - progressive cerebral atrophy syndrome | Spinocerebellar ataxia                                                                  | 1            | 1                         | ASNS (2)          | 0.00952          | 1.00000             | 6.754e-3 | 8.025e-3    |                |
| congenital microcephaly - severe encephalopathy - progressive cerebral atrophy syndrome | Liver neoplasms                                                                         | 1            | 1                         | ASNS (2)          | 0.00654          | 1.00000             | 9.871e-3 | 1.120e-2    |                |