

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: 2026-10-10 03:00:13. Filters applied: disease name contains "PMM2-congenital disorder of glycosylation" (5 matching pairs). · Rows: 5 · Generated: 2026-10-10 02:34:02

Disease A	Disease B	Shared Genes	Corroborated Shared Genes	Shared Gene Names	Similarity Score	Overlap Coefficient	P-value	FDR Q-value	Shared Cluster
hyperinsulinemic hypoglycemia with polycystic kidney disease	PMM2-congenital disorder of glycosylation	1	0	PMM2 (1)	0.50000	1.00000	6.494e-5	2.331e-4	
Cerebral atrophy	PMM2-congenital disorder of glycosylation	1	1	PMM2 (2)	0.12500	1.00000	4.546e-4	9.726e-4	
Hyperinsulinemic hypoglycemia	PMM2-congenital disorder of glycosylation	1	1	PMM2 (2)	0.08333	1.00000	7.144e-4	1.335e-3	
Pituitary stalk interruption syndrome	PMM2-congenital disorder of glycosylation	1	1	PMM2 (2)	0.03125	1.00000	2.013e-3	2.901e-3	
Cerebellar hypoplasia/atrophy, epilepsy, and global developmental delay	PMM2-congenital disorder of glycosylation	1	1	PMM2 (2)	0.02703	1.00000	2.338e-3	3.243e-3	