

GeDiPNet Cluster 82 -- Pairs Within Cluster

Similarity: Jaccard index on shared curated genes (disease_gdp). Significance: upper-tail hypergeometric test, Benjamini-Hochberg FDR-corrected across all tested pairs. See scripts/build_related.php on gedipnet.bicnirrh.res.in. Data as of: 2026-10-08 03:00:13. · Rows: 21 · Generated: 2026-10-08 20:57:03

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
Constipation	Nausea	0.18182	2	1.770e-6	9.988e-6
Rhabdomyoma	Sacroiliitis	0.50000	1	6.494e-5	2.317e-4
Eye pain	Rhabdomyoma	0.50000	1	6.494e-5	2.317e-4
Eye pain	Sacroiliitis	0.50000	1	6.494e-5	2.317e-4
Arginase deficiency	Immune system disorder	0.16667	1	3.247e-4	7.680e-4
Immune system disorder	Sacroiliitis	0.16667	1	3.247e-4	7.680e-4
Immune system disorder	Rhabdomyoma	0.16667	1	3.247e-4	7.680e-4
Eye pain	Immune system disorder	0.16667	1	3.247e-4	7.680e-4
Immune system disorder	pulmonary veno-occlusive disease and/or pulmonary capillary haemangiomatosis	0.16667	1	3.247e-4	7.680e-4
Constipation	Sacroiliitis	0.16667	1	3.247e-4	7.680e-4
Constipation	Rhabdomyoma	0.16667	1	3.247e-4	7.680e-4
Constipation	Eye pain	0.16667	1	3.247e-4	7.680e-4
Anterior cruciate ligament injury	Immune system disorder	0.16667	1	3.247e-4	7.680e-4
Eye pain	Pancreatic diseases	0.14286	1	3.897e-4	8.637e-4
Pancreatic diseases	Rhabdomyoma	0.14286	1	3.897e-4	8.637e-4
Pancreatic diseases	Sacroiliitis	0.14286	1	3.897e-4	8.637e-4
Eye pain	Nausea	0.12500	1	4.546e-4	9.697e-4
Eye pain	Facial nerve disorder	0.11111	1	5.195e-4	1.058e-3
Facial nerve disorder	Rhabdomyoma	0.11111	1	5.195e-4	1.058e-3
Facial nerve disorder	Sacroiliitis	0.11111	1	5.195e-4	1.058e-3
Congenital hereditary facial paralysis with variable hearing loss syndrome	Facial nerve disorder	0.11111	1	5.195e-4	1.058e-3