

GeDiPNet Cluster 108 -- 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: 2026-10-07 17:17:47. · Rows: 19 · Generated: 2026-10-07 15:12:01

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
11p11.2 deletion syndrome	Potocki-shaffer syndrome	0.75000	3	1.644e-12	1.934e-11
exostoses, multiple, type 2	Seizures, scoliosis, and macrocephaly/microcephaly syndrome	0.50000	1	6.494e-5	2.317e-4
Frontonasal dysplasia with alopecia and genital anomaly	Parietal foramina	0.33333	1	1.299e-4	3.908e-4
craniosynostosis 2	Parietal foramina	0.33333	1	1.299e-4	3.908e-4
11p11.2 deletion syndrome	Frontonasal dysplasia with alopecia and genital anomaly	0.25000	1	1.948e-4	5.325e-4
Potocki-shaffer syndrome	Seizures, scoliosis, and macrocephaly/microcephaly syndrome	0.25000	1	1.948e-4	5.325e-4
Intellectual developmental disorder behavioral dysmorphic	Potocki-shaffer syndrome	0.25000	1	1.948e-4	5.325e-4
Frontonasal dysplasia with alopecia and genital anomaly	Potocki-shaffer syndrome	0.25000	1	1.948e-4	5.325e-4
exostoses, multiple, type 2	Potocki-shaffer syndrome	0.25000	1	1.948e-4	5.325e-4
11p11.2 deletion syndrome	Intellectual developmental disorder behavioral dysmorphic	0.25000	1	1.948e-4	5.325e-4
11p11.2 deletion syndrome	exostoses, multiple, type 2	0.25000	1	1.948e-4	5.325e-4
11p11.2 deletion syndrome	Seizures, scoliosis, and macrocephaly/microcephaly syndrome	0.25000	1	1.948e-4	5.325e-4
11p11.2 deletion syndrome	Parietal foramina	0.20000	1	3.896e-4	8.637e-4
Parietal foramina	Potocki-shaffer syndrome	0.20000	1	3.896e-4	8.637e-4
craniosynostosis 2	Posterior cortical atrophy	0.07143	1	8.443e-4	1.497e-3
combined immunodeficiency due to ZAP70 deficiency	Posterior cortical atrophy	0.07143	1	8.443e-4	1.497e-3
Parietal foramina	Posterior cortical atrophy	0.06667	1	1.688e-3	2.529e-3
Exostoses	exostoses, multiple, type 2	0.03448	1	1.818e-3	2.675e-3
Exostoses	Seizures, scoliosis, and macrocephaly/microcephaly syndrome	0.03448	1	1.818e-3	2.675e-3