

GeDiPNet Cluster 68 -- 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 03:00:12. · Rows: 30 · Generated: 2026-10-08 09:03:55

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
Ductal carcinoma	Intraductal noninfiltrating carcinoma	0.30769	4	7.044e-13	8.653e-12
Anaplasia	Intraductal noninfiltrating carcinoma	0.33333	2	8.436e-8	5.953e-7
Anaplasia	Ductal carcinoma	0.16667	2	4.640e-7	2.904e-6
Glucose transporter type 1 deficiency syndrome	GLUT1 deficiency syndrome	0.50000	1	6.494e-5	2.317e-4
Epilepsy with myoclonic absence	GLUT1 deficiency syndrome	0.50000	1	6.494e-5	2.317e-4
Epilepsy with myoclonic absence	Glucose transporter type 1 deficiency syndrome	0.50000	1	6.494e-5	2.317e-4
Chromosome 17q23.1-q23.2 deletion syndrome	Glucose transporter type 1 deficiency syndrome	0.50000	1	6.494e-5	2.317e-4
Chromosome 17q23.1-q23.2 deletion syndrome	Epilepsy with myoclonic absence	0.50000	1	6.494e-5	2.317e-4
Chromosome 17q23.1-q23.2 deletion syndrome	GLUT1 deficiency syndrome	0.50000	1	6.494e-5	2.317e-4
Cryohydrocytosis	GLUT1 deficiency syndrome	0.33333	1	1.299e-4	3.908e-4
Childhood-onset glut1 deficiency syndrome 2	GLUT1 deficiency syndrome	0.33333	1	1.299e-4	3.908e-4
Cryohydrocytosis	Epilepsy with myoclonic absence	0.33333	1	1.299e-4	3.908e-4
Cryohydrocytosis	Glucose transporter type 1 deficiency syndrome	0.33333	1	1.299e-4	3.908e-4
Chromosome 17q23.1-q23.2 deletion syndrome	Cryohydrocytosis	0.33333	1	1.299e-4	3.908e-4
Childhood-onset glut1 deficiency syndrome 2	Epilepsy with myoclonic absence	0.33333	1	1.299e-4	3.908e-4
Childhood-onset glut1 deficiency syndrome 2	Glucose transporter type 1 deficiency syndrome	0.33333	1	1.299e-4	3.908e-4
Childhood-onset glut1 deficiency syndrome 2	Chromosome 17q23.1-q23.2 deletion syndrome	0.33333	1	1.299e-4	3.908e-4
Childhood-onset glut1 deficiency syndrome 2	Cryohydrocytosis	0.25000	1	2.598e-4	6.484e-4
Epilepsy with myoclonic absence	Intraductal noninfiltrating carcinoma	0.16667	1	3.247e-4	7.680e-4
Glucose transporter type 1 deficiency syndrome	Intraductal noninfiltrating carcinoma	0.16667	1	3.247e-4	7.680e-4
Chromosome 17q23.1-q23.2 deletion syndrome	Intraductal noninfiltrating carcinoma	0.16667	1	3.247e-4	7.680e-4
Choreoathetosis	Epilepsy with myoclonic absence	0.14286	1	3.897e-4	8.637e-4
Choreoathetosis	Chromosome 17q23.1-q23.2 deletion syndrome	0.14286	1	3.897e-4	8.637e-4
Choreoathetosis	Glucose transporter type 1 deficiency syndrome	0.14286	1	3.897e-4	8.637e-4
Epilepsy with myoclonic absence	Epilepsy with myoclonic atonic seizures	0.12500	1	4.546e-4	9.697e-4
Epilepsy with myoclonic atonic seizures	GLUT1 deficiency syndrome	0.12500	1	4.546e-4	9.697e-4
Epilepsy with myoclonic atonic seizures	Glucose transporter type 1 deficiency syndrome	0.12500	1	4.546e-4	9.697e-4
Epilepsy with myoclonic atonic seizures	Intellectual developmental disorder seizures epilepsy	0.12500	1	4.546e-4	9.697e-4
Chromosome 17q23.1-q23.2 deletion syndrome	Epilepsy with myoclonic atonic seizures	0.12500	1	4.546e-4	9.697e-4
Bladder disease	Epilepsy with myoclonic atonic seizures	0.10000	1	1.363e-3	2.146e-3