

GeDiPNet Cluster 74 -- 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 17:17:46. · Rows: 21 · Generated: 2026-10-07 12:44:24

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
Anhydramnios	autosomal dominant polycystic kidney disease	0.18750	3	9.478e-9	7.637e-8
autosomal recessive polycystic kidney disease	Caroli disease	0.33333	2	1.518e-7	1.033e-6
Anhydramnios	autosomal recessive polycystic kidney disease	0.15385	2	1.391e-6	7.989e-6
Anhydramnios	Caroli disease	0.14286	2	2.782e-6	1.510e-5
Congenital hypoplasia of aortic arch	Periportal fibrosis	0.50000	1	6.494e-5	2.317e-4
Congenital hypoplasia of aortic arch	Cystathionine beta-synthase deficiency	0.33333	1	1.299e-4	3.908e-4
Cystathionine beta-synthase deficiency	Periportal fibrosis	0.33333	1	1.299e-4	3.908e-4
autosomal recessive polycystic kidney disease	Congenital hypoplasia of aortic arch	0.25000	1	1.948e-4	5.325e-4
autosomal recessive polycystic kidney disease	Periportal fibrosis	0.25000	1	1.948e-4	5.325e-4
Congenital hypoplasia of aortic arch	Ventricular hypertrophy	0.25000	1	1.948e-4	5.325e-4
Periportal fibrosis	Ventricular hypertrophy	0.25000	1	1.948e-4	5.325e-4
Biliary-renal-neuro-skeletal syndrome	Caroli disease	0.20000	1	2.598e-4	6.484e-4
Caroli disease	Congenital hypoplasia of aortic arch	0.20000	1	2.598e-4	6.484e-4
Caroli disease	Periportal fibrosis	0.20000	1	2.598e-4	6.484e-4
Cystathionine beta-synthase deficiency	Ventricular hypertrophy	0.20000	1	3.896e-4	8.637e-4
autosomal recessive polycystic kidney disease	Cystathionine beta-synthase deficiency	0.20000	1	3.896e-4	8.637e-4
autosomal dominant polycystic kidney disease	renal-hepatic-pancreatic dysplasia 2	0.12500	1	4.546e-4	9.697e-4
radioulnar synostosis with amegakaryocytic thrombocytopenia 1	Urogenital abnormalities	0.10000	1	5.845e-4	1.154e-3
renal tubular acidosis, distal, 2, with progressive sensorineural hearing loss	Urogenital abnormalities	0.10000	1	5.845e-4	1.154e-3
Congenital hypoplasia of aortic arch	Urogenital abnormalities	0.10000	1	5.845e-4	1.154e-3
Periportal fibrosis	Urogenital abnormalities	0.10000	1	5.845e-4	1.154e-3