

GeDiPNet Cluster 94 -- 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 03:00:12. · Rows: 31 · Generated: 2026-10-07 05:26:59

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
Accessory skin tag	Darier disease	0.80000	8	1.150e-28	3.531e-27
Accessory skin tag	Benign pemphigus	0.80000	8	1.150e-28	3.531e-27
Benign pemphigus	Darier disease	0.72727	8	1.035e-27	3.075e-26
Accessory skin tag	Rothmund-thomson syndrome	0.61538	8	6.327e-27	1.813e-25
Darier disease	Rothmund-thomson syndrome	0.57143	8	5.693e-26	1.536e-24
Benign pemphigus	Rothmund-thomson syndrome	0.57143	8	5.693e-26	1.536e-24
Accessory skin tag	Cutis laxa	0.38095	8	1.610e-24	4.028e-23
Cutis laxa	Darier disease	0.36364	8	1.448e-23	3.454e-22
Benign pemphigus	Cutis laxa	0.36364	8	1.448e-23	3.454e-22
Cutis laxa	Rothmund-thomson syndrome	0.32000	8	7.948e-22	1.738e-20
Accessory skin tag	autosomal recessive cutis laxa type 2B	0.11111	1	5.195e-4	1.041e-3
Accessory skin tag	autosomal recessive cutis laxa type 2C	0.11111	1	5.195e-4	1.041e-3
Accessory skin tag	autosomal recessive cutis laxa type 2D	0.11111	1	5.195e-4	1.041e-3
autosomal recessive cutis laxa type 2D	Benign pemphigus	0.10000	1	5.845e-4	1.135e-3
Benign familial pemphigus	Benign pemphigus	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2D	Darier disease	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2C	Darier disease	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2C	Benign pemphigus	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2B	Darier disease	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2B	Benign pemphigus	0.10000	1	5.845e-4	1.135e-3
autosomal recessive cutis laxa type 2D	Rothmund-thomson syndrome	0.07692	1	7.793e-4	1.386e-3
autosomal recessive cutis laxa type 2C	Rothmund-thomson syndrome	0.07692	1	7.793e-4	1.386e-3
autosomal recessive cutis laxa type 2B	Rothmund-thomson syndrome	0.07692	1	7.793e-4	1.386e-3
Dna2-related mitochondrial dna deletion syndrome	Rothmund-thomson syndrome	0.07692	1	7.793e-4	1.386e-3
Rothmund-thomson syndrome	Rothmund-Thomson syndrome type 1	0.07692	1	7.793e-4	1.386e-3
Accessory skin tag	Demyelinating hereditary motor and sensory neuropathy	0.10000	1	1.039e-3	1.720e-3
Benign pemphigus	Demyelinating hereditary motor and sensory neuropathy	0.09091	1	1.169e-3	1.876e-3
Darier disease	Demyelinating hereditary motor and sensory neuropathy	0.09091	1	1.169e-3	1.876e-3
autosomal recessive cutis laxa type 2C	Cutis laxa	0.04762	1	1.299e-3	2.044e-3
autosomal recessive cutis laxa type 2B	Cutis laxa	0.04762	1	1.299e-3	2.044e-3

Disease A	Disease B	Similarity Score	Shared Genes	P-value	FDR Q-value
Demyelinating hereditary motor and sensory neuropathy	Rothmund-thomson syndrome	0.07143	1	1.558e-3	2.360e-3