

GeDiPNet Cluster 70 -- Top Shared Genes

Curated genes (disease\_gdp) linked to 2 or more of this cluster's member diseases, most widely shared first. Data as of: 2026-10-07 03:15:01. · Rows: 8 · Generated: 2026-10-07 01:46:08

| Gene   | Member Diseases Linked | Cluster Size | Linked Diseases                                                                                                                                                                                                             |
|--------|------------------------|--------------|-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| REN    | 7                      | 13           | familial juvenile hyperuricemic nephropathy type 2; Juvenile hyperuricemic nephropathy; Liddle syndrome; Malignant hypertension; Portal hypertension; renal tubular dysgenesis of genetic origin; Tonne-kalscheuer syndrome |
| SCNN1A | 4                      | 13           | Bronchiectasis; Liddle syndrome; pseudohypoaldosteronism, type IB1, autosomal recessive; Pseudohypoparathyroidism                                                                                                           |
| SCNN1B | 3                      | 13           | Bronchiectasis; Liddle syndrome; Pseudohypoparathyroidism                                                                                                                                                                   |
| SCNN1G | 3                      | 13           | Bronchiectasis; Liddle syndrome; Pseudohypoparathyroidism                                                                                                                                                                   |
| CUL3   | 2                      | 13           | pseudohypoaldosteronism type 2E; Pseudohypoparathyroidism                                                                                                                                                                   |
| EDEM3  | 2                      | 13           | Bronchiectasis; congenital disorder of glycosylation, type 2v                                                                                                                                                               |
| KLHL3  | 2                      | 13           | pseudohypoaldosteronism type 2D; Pseudohypoparathyroidism                                                                                                                                                                   |
| LTBR   | 2                      | 13           | Bronchiectasis; Pseudohypoparathyroidism                                                                                                                                                                                    |