PKD1 gene  
**polycystin 1, transient receptor potential channel interacting**

**Normal Function**

The *PKD1* gene provides instructions for making a protein called polycystin-1. This protein is most active in kidney cells before birth; much less of the protein is made in normal adult kidneys. Although its exact function is not well understood, polycystin-1 appears to interact with a smaller, somewhat similar protein called polycystin-2.

Polycystin-1 spans the cell membrane of kidney cells, so that one end of the protein remains inside the cell and the other end projects from the outer surface of the cell. This positioning of the protein allows it to interact with other proteins, carbohydrates, and fat molecules (lipids) outside the cell and to receive signals that help the cell respond to its environment. When a molecule binds to polycystin-1 on the surface of the cell, the protein interacts with polycystin-2 to trigger a cascade of chemical reactions inside the cell. These chemical reactions instruct the cell to undergo certain changes, such as maturing to take on specialized functions. Polycystin-1 and polycystin-2 likely work together to help regulate cell growth and division (proliferation), cell movement (migration), and interactions with other cells.

Polycystin-1 is also found in cell structures called primary cilia. Primary cilia are tiny, fingerlike projections that line the small tubes where urine is formed (renal tubules). Researchers believe that primary cilia sense the movement of fluid through these tubules, which appears to help maintain the tubules' size and structure. The interaction of polycystin-1 and polycystin-2 in renal tubules promotes the normal development and function of the kidneys.

**Health Conditions Related to Genetic Changes**

**Polycystic kidney disease**

More than 250 mutations in the *PKD1* gene have been identified in people with polycystic kidney disease. These mutations are responsible for about 85 percent of cases of autosomal dominant polycystic kidney disease (ADPKD), which is the most common type of this disorder. Mutations in the *PKD1* gene include deletions or insertions of DNA building blocks (base pairs) and alterations of one or more base pairs. Most *PKD1* mutations are predicted to produce an abnormally small, nonfunctional version of the polycystin-1 protein. Although researchers are uncertain how a lack of polycystin-1 leads to the formation of cysts, it probably disrupts the protein's signaling function within the cell and in primary cilia. As a result, cells lining the renal tubules may grow and divide abnormally, leading to the growth of numerous cysts characteristic of polycystic kidney disease.
Chromosomal Location

Cytogenetic Location: 16p13.3, which is the short (p) arm of chromosome 16 at position 13.3

Molecular Location: base pairs 2,088,708 to 2,135,898 on chromosome 16 (Homo sapiens Updated Annotation Release 109.20191205, GRCh38.p13) (NCBI)

Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- Lov-1
- PBP
- Pc-1
- PC1
- PKD1_HUMAN
- polycystic kidney disease 1 (autosomal dominant)
- polycystin-1
- TRPP1

Additional Information & Resources

Clinical Information from GeneReviews
- Polycystic Kidney Disease, Autosomal Dominant
  https://www.ncbi.nlm.nih.gov/books/NBK1246

Scientific Articles on PubMed
- PubMed
  https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28PKD1%5BTIAB%5D%29+OR+%28polycystic+kidney+disease+1%5BTIAB%5D%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+720+days%22%5Bdp%5D
Catalog of Genes and Diseases from OMIM

- POLYCYSTIN 1
  http://omim.org/entry/601313

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
  http://atlasgeneticsoncology.org/Genes/PKD1ID41725ch16p13.html
- ClinVar
  https://www.ncbi.nlm.nih.gov/clinvar?term=PKD1%5Bgene%5D
- HGNC Gene Symbol Report
- Monarch Initiative
  https://monarchinitiative.org/gene/NCBIGene:5310
- NCBI Gene
- UniProt
  https://www.uniprot.org/uniprot/P98161

Sources for This Summary

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