



NPHS2 gene

NPHS2, podocin

Normal Function

The *NPHS2* gene provides instructions for making a protein called podocin. Podocin is primarily found in the kidneys, which are organs that filter waste products from the blood and remove them in urine. Specifically, podocin is found in cells called podocytes, which are located in specialized kidney structures called glomeruli. Podocin is located at the cell surface in the area between two podocytes called the slit diaphragm. The slit diaphragm is known as a filtration barrier because it captures proteins in blood so that they remain in the body while allowing other molecules like sugars and salts to be excreted in urine. Podocin likely helps bring other proteins that are needed for a functional slit diaphragm to the podocyte cell surface. The protein also is involved with podocyte cell signaling, helping the cell adapt to changes that occur during the filtration process.

Health Conditions Related to Genetic Changes

Congenital nephrotic syndrome

At least 170 mutations in the *NPHS2* gene have been found to cause congenital nephrotic syndrome. This condition is a kidney disorder that begins in infancy and typically leads to irreversible kidney failure (end-stage renal disease) by early childhood. Mutations in this gene appear to be the most frequent cause of congenital nephrotic syndrome. Most *NPHS2* gene mutations change single protein building blocks (amino acids) in the podocin protein. These mutations result in a reduction or absence of functional protein, which impairs the formation of normal slit diaphragms. Without a functional slit diaphragm, molecules pass through the kidneys abnormally and are excreted in urine. The filtering ability of the kidneys worsens from birth, eventually leading to end-stage renal disease.

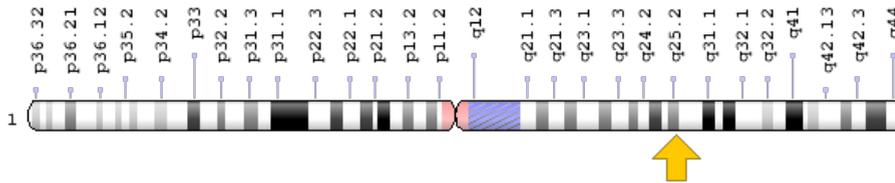
Other disorders

NPHS2 gene mutations can cause other forms of nephrotic syndrome that develop later in life. In one form, called infantile nephrotic syndrome, signs and symptoms of the condition appear between 4 and 12 months of age. The features of this condition are similar to congenital nephrotic syndrome (described above), but they are often less severe. It is likely that *NPHS2* gene mutations that cause infantile nephrotic syndrome have less effect on podocin function than those that cause congenital nephrotic syndrome, accounting for the later onset of the disorder.

Chromosomal Location

Cytogenetic Location: 1q25.2, which is the long (q) arm of chromosome 1 at position 25.2

Molecular Location: base pairs 179,550,539 to 179,575,987 on chromosome 1 (Homo sapiens Updated Annotation Release 109.20190607, GRCh38.p13) (NCBI)



Credit: Genome Decoration Page/NCBI

Other Names for This Gene

- nephrosis 2, idiopathic, steroid-resistant (podocin)
- NPHS2 podocin
- PDCN
- podocin isoform 1
- podocin isoform 2
- SRN1

Additional Information & Resources

Educational Resources

- National Institute of Diabetes and Digestive and Kidney Diseases: The Kidneys and How They Work
<https://www.niddk.nih.gov/health-information/kidney-disease/kidneys-how-they-work>

Scientific Articles on PubMed

- PubMed
<https://www.ncbi.nlm.nih.gov/pubmed?term=%28%28NPHS2%5BTIAB%5D%29+OR+%28podocin%5BTIAB%5D%29%29+AND+%28%28Genes%5BMH%5D%29+OR+%28Genetic+Phenomena%5BMH%5D%29%29+AND+english%5Bla%5D+AND+human%5Bmh%5D+AND+%22last+1440+days%22%5Bdp%5D>

Catalog of Genes and Diseases from OMIM

- **PODOCIN**
<http://omim.org/entry/604766>

Research Resources

- Atlas of Genetics and Cytogenetics in Oncology and Haematology
http://atlasgeneticsoncology.org/Genes/GC_NPHS2.html
- ClinVar
<https://www.ncbi.nlm.nih.gov/clinvar?term=NPHS2%5Bgene%5D>
- HGNC Gene Symbol Report
https://www.genenames.org/data/gene-symbol-report#!/hgnc_id/HGNC:13394
- Monarch Initiative
<https://monarchinitiative.org/gene/NCBIGene:7827>
- NCBI Gene
<https://www.ncbi.nlm.nih.gov/gene/7827>
- UniProt
<https://www.uniprot.org/uniprot/Q9NP85>

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