DISEASE:
Congenital nephrotic syndrome, Finnish type

NAME:	Congenital nephrotic syndrome, Finnish type
DESCRIPTION:	A rare congenital nephrotic syndrome characterized by massive protein loss and marked edema manifesting in utero or during the first 3 months of life.
ORPHACODE:	839
SYNONYMS:	Finnish congenital nephrosis
XREF(S):	Orphanet MeSH MedDRA ICD-10 OMIM
ANALYTE(S):	NPHS1
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RELATED CONTENT

Related Genetic Tests

- Nephrotic syndrome, Focal Segmental Glomerulosclerosis (FSGS) , Alport syndrome and podocytopathy (gene panel)

Related Laboratories

- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)

Related Analytes

- NPHS1 adhesion molecule, nephrin

Related Gene Panels

- Nephrotic syndrome, FSGS, Alport syndrome (76 genes) - IPG