

**ANALYTE:**  
**PRKCSH**

<b>NAME:</b>	protein kinase C substrate 80K-H
<b>SYMBOL:</b>	PRKCSH
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	GIIB VASAP-60 advanced glycation end-product receptor 2 glucosidase 2 subunit beta glucosidase II beta subunit hepatocystin
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Reactome</a> <a href="#">SwissProt</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

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## RELATED CONTENT

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### Related Genetic Tests

- [Ciliopathy / polycystic kidney and liver diseases / ADTKD/ nephronophtisis / Bardet-Biedl syndromes and kidney cancers \(gene panel\)](#)
- [Congenital malformation \(gene panel - 1721 genes\)](#)
- [Hepatology \(gene panel\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders \(213 genes\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)

### Related Diseases

- [Isolated polycystic liver disease](#)

### Related Gene Panels

- [Ciliopathy, polycystic kidney and liver diseases, ADTKD, nephronophtisis, Bardet-Biedl syndromes and kidney cancers \(146 genes\) - IPG](#)
- [Congenital malformation \(1721 genes\) - ULB](#)
- [Hepatology panel - UGent](#)
- [Metabolic disorders \(213 genes\) - VUB](#)
- [Nephropathy panel - UGent](#)
- [Panel Nephro-ULG-V1](#)