

**ANALYTE:**  
**WNK1**

<b>NAME:</b>	WNK lysine deficient protein kinase 1
<b>SYMBOL:</b>	WNK1
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<b>SYNONYMS:</b>	HSAN2 PPP1R167 protein phosphatase 1, regulatory subunit 167
<b>XREF(S):</b>	<a href="#">Orphanet</a> <a href="#">Ensembl</a> <a href="#">Genatlas</a> <a href="#">HGNC</a> <a href="#">OMIM</a> <a href="#">Reactome</a> <a href="#">SwissProt</a>
<b>CREATED:</b>	13 May 2019 - 01:01
<b>CHANGED:</b>	22 Jun 2023 - 16:14

## RELATED CONTENT

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### Related Genetic Tests

- [Charcot-Marie-Tooth \(other than type 1A\) \(gene panel, IPN panel\)](#)
- [Dermatogenetic panel, severe, rare and hereditary genodermatoses \(gene panel - 394 genes\)](#)
- [Inherited Kidney Diseases \(Gene Panel\)](#)
- [Nephrogenetics / Nephropathy \(gene panel\)](#)
- [Nephropathies, hereditary \(gene panel\)](#)
- [Neuromuscular disorders \(548 genes\)](#)
- [Neuropathy \(gene panel\)](#)
- [Neuropathy \(gene panel\)](#)
- [Peripheral neuropathy \(gene panel\)](#)
- [Tubulopathy \(gene panel\)](#)

### Related Diseases

- [Hereditary sensory and autonomic neuropathy type 2](#)
- [Pseudohypoaldosteronism type 2C](#)

### Related Gene Panels

- [Inherited Peripheral Neuropathies gene panel \(139 genes\) - KUL](#)
- [Dermatogenetic / severe, rare and hereditary genodermatoses \(394 genes\) - ULB](#)
- [Nephropathies, hereditary \(219 genes\) - KUL](#)
- [Nephropathy panel - UGent](#)
- [Neuromuscular disorders \(548 genes\) - ULB](#)
- [Neuropathy \(148 genes\) - IPG](#)

- Neuropathy (genepanel) - UZA
- Neuropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG

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