

**ANALYTE:**  
**SLC5A2**

<b>NAME:</b>	solute carrier family 5 member 2
<b>SYMBOL:</b>	SLC5A2
<b>VERSION OF ORPHANET:</b>	2023-06-22 14:14:43
<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

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- [Nephropathies, hereditary \(gene panel\)](#)
- [Neurodevelopmental disorders \(1300 genes\)](#)
- [Neurodevelopmental disorders gene panel](#)
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- [Neurodevelopmental disorders \(1300 genes\) - ULB](#)
- [Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders \(1162 genes\) - VUB](#)
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- [Tubulopathy/Nephrolithiasis \(106 genes\) - IPG](#)