# ANALYTE:
# SLC3A1

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|---|---|
| **NAME:** | solute carrier family 3 member 1 |
| **SYMBOL:** | SLC3A1 |
| **VERSION OF ORPHANET:** | 2023-06-22 14:14:43 |
| **SYNONYMS:** | ATR1<br>CSNU1<br>D2H<br>NBAT<br>RBAT |
| **XREF(S):** | Orphanet<br>Ensembl<br>Genatlas<br>HGNC<br>OMIM<br>Reactome<br>SwissProt |
| **CREATED:** | 13 May 2019 - 01:01 |
| **CHANGED:** | 22 Jun 2023 - 16:14 |

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# RELATED CONTENT

## Related Genetic Tests

- Inherited Kidney Diseases (Gene Panel)
- Metabolic disorders including disorders of glycosylation, peroxisomal disorders, organic acidurias, glycogenosis disorders, neurotransmitter disorders (213 genes)
- Nephrocalcinosis and nephrolithiasis (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

## Related Diseases

- 2p21 microdeletion syndrome
- Atypical hypotonia-cystinuria syndrome
- Cystinuria type A
- Hypotonia-cystinuria syndrome

## Related Gene Panels

- Metabolic disorders (213 genes) - VUB
- Nephrocalcinosis and nephrolithiasis (37 genes) - IPG
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
- Tubulopathy/Nephrolithiasis (106 genes) - IPG