Belgian Genetic Tests database

ANALYTE:
BSND

NAME:	barttin CLCNK type accessory subunit beta
SYMBOL:	BSND
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BART
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- End-stage renal disease, ESRD (gene panel)
- Hearing loss (deafness), (gene panel)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability (virtual gene panel)
- Nephrocalcinosis and nephrolithiasis (gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Tubulopathy (gene panel)

Related Diseases

- Bartter syndrome type 4
- Rare autosomal recessive non-syndromic sensorineural deafness type DFNB

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- End-stage renal disease (106 genes) - IPG
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability (gene panel)

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