Belgian Genetic Tests database

ANALYTE:
PBX1

NAME:	PBX homeobox 1
SYMBOL:	PBX1
VERSION OF ORPHANET:	2023-06-22 14:14:43
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Source URL: http://gentest.healthdata.be/analyte/3868

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Inherited Kidney Diseases (Gene Panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (virtual gene panel)
- Nephrogenetics / Nephropathy (gene panel)
- Nephropathies, hereditary (gene panel)
- Renal or urinary tract malformation (CAKUT) (gene panel)

Related Diseases

- B-lymphoblastic leukemia/lymphoma with recurrent genetic abnormality
- B-lymphoblastic leukemia/lymphoma with t(1;19)(q23;p13.3)
- Precursor B-cell acute lymphoblastic leukemia
- Renal hypoplasia, bilateral

Related Gene Panels

- Cakut (congenital anomalies of the kidney and urinary tract-1) (69 genes) - IPG
- Congenital malformation (1721 genes) - ULB
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Nephropathies, hereditary (219 genes) - KUL
- Nephropathy panel - UGent
- Panel Nephro-ULG-V1
