ANALYTE:
TWIST1

NAME:	twist family bHLH transcription factor 1
SYMBOL:	TWIST1
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	BPES2 CRS1 H-twist SCS Saethre-Chotzen syndrome bHLHa38
XREF(S):	Orphanet Reactome HGNC OMIM SwissProt Ensembl Genatlas
CREATED:	13 May 2019 - 01:01
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RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Congenital structural heart defects (gene panel)
- Craniosynostosis (gene panel)
- Intellectual disability & Epilepsy (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Neurodevelopmental disorders (1300 genes)
- Neurodevelopmental disorders gene panel
- Saethre-Chotzen syndrome
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Isolated plagiocephaly
- Non-syndromic bicoronal craniosynostosis
- Non-syndromic sagittal craniosynostosis
- Saethre-Chotzen syndrome

Related Gene Panels

- Congenital malformation (1721 genes) - ULB

- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Craniosynostosis (32 genes) - KUL
- Intellectual disability & Epilepsy - UGent
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Neurodevelopmental disorders (1300 genes) - ULB
- Neurodevelopmental disorders: developmental delay, intellectual disability, autistic disorders (1162 genes) - VUB
- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
