ANALYTE:
HOXD13

NAME:	homeobox D13
SYMBOL:	HOXD13
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	synpolydactyly
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
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Source URL: http://gentest.healthdata.be/analyte/868

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Synpolydactyly / Brachydactyly

Related Diseases

- Brachydactyly type E
- Brachydactyly-syndactyly, Zhao type
- NON RARE IN EUROPE: Brachydactyly type D
- Syndactyly type 5
- Synpolydactyly type 1
- VACTERL/VATER association
- Zygodactyly type 3

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Intellectual disability (gene panel)
- Skeletal dysplasia (394 genes) - VUB

- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
