Belgian Genetic Tests database

ANALYTE:
PTCH2

NAME:	patched 2
SYMBOL:	PTCH2
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XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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Source URL: http://gentest.healthdata.be/analyte/379

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Gorlin syndrome (gene panel)
- Medulloblastoma (3 genes)
- Onco-endocrine pathologies (gene panel)

Related Diseases

- Gorlin syndrome
- Tessier number 7 facial cleft

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Gorlin syndrome (3 genes)
- Medulloblastoma (3 genes) - UCL
- Onco-endocine pathologies (50 genes) - UCL