ANALYTE:
PAX3

NAME:	paired box 3
SYMBOL:	PAX3
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	HUP2
XREF(S):	Orphanet Ensembl Genatlas HGNC OMIM Reactome SwissProt
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)
- Dermatogenetic panel, severe, rare and hereditary genodermatoses (gene panel - 394 genes)
- Hearing loss (deafness), (gene panel)
- Intellectual disability (gene panel)
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skin disorders (gene panel)
- Waardenburg Syndrome types I and III
- Waardenburg syndrome (gene panel)
- cleft lip with/whitout cleft palate (virtual gene panel)

Related Diseases

- Alveolar rhabdomyosarcoma
- Craniofacial-deafness-hand syndrome
- Waardenburg syndrome type 1
- Waardenburg syndrome type 3

Related Gene Panels

- Cleft lip and palate / dysmorphic facial features / craniofacial anomalies (255 genes)) - UCL

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Congenital structural heart defects - UGent
- Dermatogenetic / severe, rare and hereditary genodermatoses (394 genes) - ULB
- Hearing loss (deafness) (genepanel) - UZA
- Hearing loss (deafness) syndromic (59 genes) - UZA
- Intellectual disability (gene panel)
- Intellectual disability/Epilepsy (1091 genes) - ULG
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
- Skin disorders - UGent
- Waardenburg syndrome (6 genes) - UZA
