ANALYTE:
MSX2

NAME:	msh homeobox 2
SYMBOL:	MSX2
VERSION OF ORPHANET:	2023-06-22 14:14:43
SYNONYMS:	CRS2 FPP HOX8 MSH PFM craniosynostosis, type 2
XREF(S):	Orphanet Reactome Ensembl Genatlas HGNC OMIM SwissProt
CREATED:	13 May 2019 - 01:01
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/analyte/1786

RELATED CONTENT

Related Genetic Tests

- Congenital malformation (gene panel - 1721 genes)
- Congenital malformation gene panel
- Craniosynostosis (gene panel)
- Early onset epileptic encephalopathy (gene panel - 845 genes)
- Epilepsy gene panel
- Intellectual disability (virtual gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)
- Skeletal dysplasia (gene panel)

Related Diseases

- Craniosynostosis, Boston type
- Enlarged parietal foramina
- Parietal foramina with clavicular hypoplasia

Related Gene Panels

- Congenital malformation (1721 genes) - ULB
- Congenital malformation gene panel - VUB
- Craniosynostosis (32 genes) - KUL
- Early onset epileptic encephalopathy (845 genes) - ULB
- Epilepsy gene panel - VUB
- Intellectual disability (gene panel)

- Skeletal dysplasia (394 genes) - VUB
- Skeletal dysplasia (genepanel) - UZA
- Skeletal dysplasia - UGent
