Belgian Genetic Tests database

DISEASE:
Muenke syndrome

NAME:	Muenke syndrome
DESCRIPTION:	Muenke syndrome is a syndromic craniosynostosis with significant phenotypic variability, usually characterized by coronal synostosis, midfacial retrusion, strabismus, hearing loss and developmental delay.
ORPHACODE:	53271
XREF(S):	Orphanet MeSH OMIM ICD-10
ANALYTE(S):	FGFR3
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Source URL: http://gentest.healthdata.be/disease/2992

RELATED CONTENT

Related Genetic Tests

- Craniosynostosis (gene panel)
- Craniosynostosis / Muenke syndrome (hot spot mutation - p.Pro250Arg)
- Craniosynostosis, Muenke syndrome (hot spot mutation - p.Pro250 in FGFR3 gene)
- Craniosynostosis, Muenke syndrome (hot spot mutation - p.Pro250Arg)
- Craniosynostosis, Muenke syndrome (hot spot mutation - p.Pro250Arg)
- Short Stature (gene panel)
- Short stature/ Growth retardation/ (gene panel)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centre de Génétique-Institut de Pathologie et de Génétique (IPG)
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Medische Genetica - UZ Gent
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- fibroblast growth factor receptor 3

Related Gene Panels

- Craniosynostosis (32 genes) - KUL
- Growth retardation/short stature (genepanel) - UZA
- Short Stature (46 genes) - IPG
