Belgian Genetic Tests database

DISEASE:
Crouzon syndrome

NAME:	Crouzon syndrome
DESCRIPTION:	Crouzon disease is characterized by craniosynostosis and facial hypoplasia.
ORPHACODE:	207
SYNONYMS:	Crouzon craniofacial dysostosis
XREF(S):	Orphanet ICD-10 OMIM
ANALYTE(S):	FGFR2 ERF
CREATED:	13 May 2019 - 01:02
CHANGED:	22 Jun 2023 - 16:14

Source URL: http://gentest.healthdata.be/disease/179

RELATED CONTENT

Related Genetic Tests

- Craniosynostosis (gene panel)
- Craniosynostosis / Crouzon syndrome (hot spot mutation - exon 9)
- Craniosynostosis syndromes (Apert, Crouzon)

Related Laboratories

- Centre de Génétique Humaine - CHU Sart-Tilman
- Centrum Medische Genetica - UZ Antwerpen
- Centrum Menselijke Erfelijkheid - KUL

Related Analytes

- ETS2 repressor factor
- fibroblast growth factor receptor 2

Related Gene Panels

- Craniosynostosis (32 genes) - KUL