sciensano **Belgian Genetic Tests database**

GENETIC TEST:
Craniosynostosis, Muenke syndrome (hot spot mutation - p.Pro250 in FGFR3 gene)

FULL NAME:	Craniosynostosis, Muenke syndrome (hot spot mutation - p.Pro250 in FGFR3 gene)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565390-565401
TURNAROUND TIME (MAXIMUM):	6 weeks
CREATED:	29 Jul 2019 - 14:29
CHANGED:	16 May 2022 - 09:38

RELATED CONTENT

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