GENETIC TEST:
Craniosynostosis syndrome (hot spot mutation - p.Pro252Arg)

FULL NAME:	Craniosynostosis syndrome (hot spot mutation - p.Pro252Arg)
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):	2 months
CREATED:	20 Aug 2019 - 11:57
CHANGED:	25 Jan 2023 - 11:38
URL:	https://labogidsmedgen.uza.be/analyses/pfeiffer-syndroom

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