GENETIC TEST:
Craniosynostosis / Apert syndrome (hot spot mutations - exon 7)

FULL NAME:	Craniosynostosis / Apert syndrome (hot spot mutations - exon 7)
TEST TYPE:	Clinical
TEST SPECIALTY:	Molecular Genetics
TEST PURPOSE:	Post-natal Diagnosis, Prenatal diagnosis
SPECIMEN:	Peripheral (whole) blood on EDTA, Amniotic fluid, Chorionic villi, DNA
METHOD CATEGORY:	Targeted variant analysis
METHOD TECHNIQUE:	Bi-directional Sanger Sequence analysis
RIZIV CODE:	565390-565401
EQA:	• Skeletal dysplasis
TURNAROUND TIME (MAXIMUM):	21 days

CREATED:	20 Aug 2019 - 12:05
CHANGED:	16 Jan 2024 - 13:19
URL:	https://www.chuliege.be/jcms/c_525300/fr/apert-syndrome

Source URL: http://gentest.healthdata.be/genetic_test/818

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